THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 2093 **Seq:** 1 **Gender:** F
Reported: 28/07/1989 **Weight:**
Hospitalisation: **Age:** 77Y
Onset Date: 21/05/1989 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hallucination			

Medicine Details:

PANADOL (Other drug) **Reason:**
0.0
Batch: **Started:** 11/05/1989 **Stopped:** 30/05/1989

DYMADON CO (Other drug) **Reason:**
0.0
Batch: **Started:** 11/05/1989 **Stopped:** CONTIN

RANITIDINE (Other drug) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 11/05/1989 **Stopped:** CONTIN

SUCRALFATE (Other drug) **Reason:**
4.0 Gram Daily
Batch: **Started:** 11/05/1989 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Case Sequenced to: 61792/2

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 2093 **Seq:** 1
Reported: 28/07/1989
Hospitalisation:
Onset Date: 21/05/1989
Outcome: Not yet recovered

Gender: F
Weight:
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** 15/05/1989 **Stopped:** 21/05/1989

Laboratory Investigations:

Additional Information:

Case Sequenced to: 61792/2

Report Detail

Case Number: 2112 **Seq:** 1 **Gender:** M
Reported: 06/09/1989 **Weight:** 96.00
Hospitalisation: **Age:** 53Y
Onset Date: 25/08/1989 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous			Observation

Medicine Details:

RENITEC (Suspected) **Reason:** Essential benign hypertension
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 25/08/1989 **Stopped:** 25/08/1989

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 25.0 Dose Unspecified Daily Oral
Batch: **Started:** 08/08/1989 **Stopped:** CONTIN

MINIPRESS (Suspected) **Reason:** Essential benign hypertension
Tablet 1.0 Milligram Daily Oral
Batch: **Started:** 22/08/1989 **Stopped:** 24/08/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Case Sequenced to: 62369/2

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 2764 **Seq:** 1 **Gender:** M
Reported: 02/11/1990 **Weight:** 75.00
Hospitalisation: **Age:** 52Y
Onset Date: 31/05/1988 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased Testicular disorder			

Medicine Details:

CHOLESTYRAMINE (Other drug) **Reason:** Xanthomatosis
Tablet 12.0 Gram Daily Oral
Batch: **Started:** 01/12/1987 **Stopped:** 28/08/1990

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 01/12/1987 **Stopped:** 28/08/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				10/7/90 range serum testosterone: 6.0 (10-49)

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. Case Sequenced to: 68158/2

Report Detail

Case Number: 58056 **Seq:** 1 **Gender:** U
Reported: 05/01/1989 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			Dilosyn 4mg bd

Medicine Details:

ENDONE (Suspected)			**Reason:** Pain			
Tablet	5.0	Milligram	As necessary	Oral		
Batch:	**Started:**			**Stopped:**		CONTIN
PROLADONE (Suspected)			**Reason:** Pain			
Suppository	2.0	Dose Unspecified	Daily	Rectal		
Batch:	**Started:**			**Stopped:**		CONTIN
PANADOL (Suspected)			**Reason:** Pain			
Tablet	4.0	Gram	Daily	Oral		
Batch:	**Started:**			**Stopped:**		CONTIN
TRYPTANOL (Suspected)			**Reason:** Specific disorders of sleep			
Tablet	50.0	Milligram	Daily	Oral		
Batch:	**Started:**			**Stopped:**		CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 58403 **Seq:** 1 **Gender:** F
Reported: 26/01/1989 **Weight:** 60.00
Hospitalisation: **Age:** 77Y
Onset Date: 20/01/1989 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Other diseases of bladder
25.0 Milligram Daily
Batch: **Started:** 01/01/1989 **Stopped:** 20/01/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 58793 **Seq:** 1 **Gender:** F
Reported: 11/02/1989 **Weight:** 70.00
Hospitalisation: **Age:** 55Y
Onset Date: 02/02/1989 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

ZANTAC (Suspected) **Reason:** Diaph hern of abd cav w/o obst
300.0 Milligram Daily
Batch: **Started:** 02/02/1989 **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** 01/06/1987 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 59137 **Seq:** 1 **Gender:** U
Reported: 06/03/1989 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dental caries Dry mouth			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 0.0 Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 59427 **Seq:** 1 **Gender:** F
Reported: 17/03/1989 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 31/12/1987 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cardiac failure Cardiomyopathy	Life		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** 01/01/1979 **Stopped:** 15/11/1987

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				1biventricular chamber dilatation

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 60485 **Seq:** 1 **Gender:** M
Reported: 19/05/1989 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 15/05/1989 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Confusional state		Visual and auditory hallucinations	

Medicine Details:

NAVIDREX (Other drug) **Reason:**
Tablet 3.0 Dose Unspecified Weekly Oral
Batch: **Started:** **Stopped:**

MIDAMOR (Other drug) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

VENTOLIN (Other drug) **Reason:**
Inhalation 8.0 Dose Unspecified Daily Inhalation
Batch: **Started:** **Stopped:**

THEO-DUR (Other drug) **Reason:**
200.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 60485 **Seq:** 1 **Gender:** M
Reported: 19/05/1989 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 15/05/1989 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

PREDNISONE (Other drug)			**Reason:**	
	5.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	
TRYPTANOL (Suspected)			**Reason:**	Herpes zoster
	2.0 Dose Unspecified	Daily		
Batch:	**Started:**	10/05/1989	**Stopped:**	15/05/1989

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 60978 **Seq:** 1
Reported: 16/06/1989
Hospitalisation: Hospitalisation prolonged
Onset Date: 01/06/1989
Outcome: Not yet recovered

Gender: M
Weight: 102.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Hepatocellular damage Hypoglycaemia Shock	Life		Cessation of medication, 10% dextrose infusion and saline by central venous catheter

Medicine Details:

TRENTAL 400 (Suspected) **Reason:** Otr&unspec peripher vascul dis
Tablet 1.2 Gram Daily Oral
Batch: **Started:** 01/04/1989 **Stopped:** 01/06/1989

FELDENE (Suspected) **Reason:** Osteoarthritis
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 01/06/1988 **Stopped:** 01/06/1989

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 01/06/1989

DIABINESE (Suspected) **Reason:** Diabetes mellitus
Tablet 2.5 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 01/06/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				Ast 1224 alt 603 alp 1.9 prothrombin 1.9

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 60978 **Seq:** 1
Reported: 16/06/1989
Hospitalisation: Hospitalisation prolonged
Onset Date: 01/06/1989
Outcome: Not yet recovered

Gender: M
Weight: 102.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ZYLOPRIM (Suspected)				**Reason:**	Gout	
Tablet	300.0	Milligram	Daily		Oral	
Batch:	**Started:**			**Stopped:**		L TERM
VENTOLIN (Suspected)				**Reason:**	Otr respiratory systm diseases	
Inhalation	8.0	Dose Unspecified	Daily		Inhalation	
Batch:	**Started:**			**Stopped:**		L TERM

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 61340 **Seq:** 1 **Gender:** F
Reported: 10/07/1989 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: 31/05/1989 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue Neutropenia			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** 01/01/1987 **Stopped:** 31/01/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haematology				1 neutropenia (1.41) 2/5/89. neutrophils 2.74 on 17/6/89.

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Report Detail

Case Number: 61573 **Seq:** 1 **Gender:** F
Reported: 19/07/1989 **Weight:** 58.00
Hospitalisation: **Age:** 76Y
Onset Date: 05/07/1989 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Fluid restriction

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr periph nerve dis not auton
25.0 Milligram Daily
Batch: **Started:** 13/05/1989 **Stopped:** 07/07/1989

TRIMETHOPRIM (Suspected) **Reason:** Othr diseases of urinary tract
300.0 Milligram Daily
Batch: **Started:** 02/05/1989 **Stopped:** 05/07/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 62032 **Seq:** 1 **Gender:** U
Reported: 22/08/1989 **Weight:**
Hospitalisation: **Age:**
Onset Date: 04/07/1989 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hallucination			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

PREDNISOLONE (Suspected) **Reason:** Other rheumatoid arthritis

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 62935 **Seq:** 1 **Gender:** F
Reported: 12/10/1989 **Weight:** 48.00
Hospitalisation: **Age:** 38Y
Onset Date: 10/10/1989 **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Pain
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 25/09/1989 **Stopped:** 08/10/1989

TRYPTANOL (Suspected) **Reason:** Pain
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 10/10/1989 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 63688 **Seq:** 1 **Gender:** M
Reported: 12/12/1989 **Weight:**
Hospitalisation: **Age:** 22Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Oral application 175.0 Milligram Daily Oral

Batch: **Started:** 01/03/1989 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 65623 **Seq:** 1 **Gender:** M
Reported: 23/05/1990 **Weight:** 60.00
Hospitalisation: **Age:** 66Y
Onset Date: 01/05/1990 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspepsia			

Medicine Details:

RENITEC (Other drug) **Reason:** Congestive heart failure
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

LANOXIN PG (Other drug) **Reason:** Congestive heart failure
Tablet 187.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SEREPAX (Other drug) **Reason:** Otr spec symp psychopathol nec
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 01/05/1990 **Stopped:** 06/05/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 65652 **Seq:** 1 **Gender:** M
Reported: 04/05/1990 **Weight:** 69.00
Hospitalisation: **Age:** 77Y
Onset Date: 12/04/1990 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular Syncope			Changed to desipramine hcl

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 10/04/1990 **Stopped:** 17/04/1990

METHADONE HYDROCHLORIDE (Suspected) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 65903 **Seq:** 1 **Gender:** F
Reported: 04/06/1990 **Weight:** 18.00
Hospitalisation: **Age:** 03Y
Onset Date: 09/04/1990 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Personality disorder Pruritus Urticaria			

Medicine Details:

AMITRIP (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 03/04/1990 **Stopped:** 11/04/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 66366 **Seq:** 1 **Gender:** F
Reported: 28/06/1990 **Weight:** 55.00
Hospitalisation: **Age:** 14Y
Onset Date: 22/06/1990 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypokalaemia			Amphotericin increased in successive doses: 30, 40, 50mg. ceased amphotericin 24/6/90. total of 3.5 umol/kg/24hrs. kci infusion (0.5mmol/kg/24hrs & tpn 3 mmol/kg/24hrs).

Medicine Details:

DIAZEPAM (Suspected) **Reason:**
4.0 Milligram As necessary
Batch: **Started:** **Stopped:** 20/06/1990

AMPHOTERICIN (Suspected) **Reason:** Moniliasis
Injection 15.0 Milligram Daily Intravenous
Batch: **Started:** 21/06/1990 **Stopped:** 24/06/1990

CARBAMAZEPINE (Suspected) **Reason:**
800.0 Milligram Daily
Batch: **Started:** **Stopped:**

NILSTAT (Suspected) **Reason:**
30.0 Millilitre Daily
Batch: **Started:** **Stopped:** 25/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrolytes				22/06, 24/06, 25/06 3.9 3.0(1008),3.6 (0205) after kcl infusion 10 mmol/hr 2.7(1620),3.2 (1150) for 5 hrs

Additional Information:

Report Detail

Case Number: 66366 **Seq:** 1
Reported: 28/06/1990
Hospitalisation:
Onset Date: 22/06/1990
Outcome: Not yet recovered

Gender: F
Weight: 55.00
Age: 14Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

CISAPRIDE (Suspected)			**Reason:**	
	30.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	
METHYLDOPA (Suspected)			**Reason:**	
	750.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	25/06/1990
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	25/06/1990
PROPRANOLOL HYDROCHLORIDE (Suspected)			**Reason:**	
	120.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	20/06/1990

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 66366 **Seq:** 1
Reported: 28/06/1990
Hospitalisation:
Onset Date: 22/06/1990
Outcome: Not yet recovered

Gender: F
Weight: 55.00
Age: 14Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

CEFOTAXIME SODIUM (Suspected) **Reason:**
6.0 Gram Daily
Batch: **Started:** 20/06/1990 **Stopped:** 25/06/1990

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 66370 **Seq:** 1 **Gender:** M
Reported: 03/07/1990 **Weight:** 70.00
Hospitalisation: **Age:** 39Y
Onset Date: 18/06/1990 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperhidrosis Malaise Paraesthesia Pruritus			

Medicine Details:

SEREPAX (Other drug) **Reason:** Depression
30.0 Milligram As necessary
Batch: **Started:** 02/05/1988 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 18/06/1990 **Stopped:** 21/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 66567 **Seq:** 1 **Gender:** F
Reported: 13/07/1990 **Weight:**
Hospitalisation: **Age:** 89Y
Onset Date: 09/07/1990 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dermatitis exfoliative Rash			Prednisolone 50mg orally stat.

Medicine Details:

CLINORIL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

LASIX (Suspected) **Reason:** Congestive heart failure
80.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

LANOXIN PG (Suspected) **Reason:** Congestive heart failure
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 66567 **Seq:** 1 **Gender:** F
Reported: 13/07/1990 **Weight:**
Hospitalisation: **Age:** 89Y
Onset Date: 09/07/1990 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

THYROXINE SODIUM (Suspected)			**Reason:**	
	50.0	Microgram	Daily	
Batch:	**Started:**			**Stopped:**
SLOW-K (Suspected)			**Reason:**	
	6.0	Dose Unspecified	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 66581 **Seq:** 1 **Gender:** M
Reported: 11/07/1990 **Weight:** 84.00
Hospitalisation: **Age:** 30Y
Onset Date: 20/06/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia			Barrier nursing, tobramycin iv 140mg tds.

Medicine Details:

NITROUS OXIDE (Suspected) **Reason:** Pain
0.0
Batch: **Started:** 25/05/1990 **Stopped:** 20/06/1990

CALCIPARINE (Suspected) **Reason:** Other prophylactic procedures
Injection 10.0 Thousand Internat Daily Subcutaneous
Batch: **Started:** 23/05/1990 **Stopped:** 20/06/1990

LARGACTIL (Suspected) **Reason:** Pain
Oral application 175.0 Milligram Daily Oral
Batch: **Started:** 01/05/1990 **Stopped:** 20/06/1990

AMITRIP (Suspected) **Reason:**
Oral application 125.0 Milligram Daily Oral
Batch: **Started:** 26/04/1990 **Stopped:** 20/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haematology				1 1/6/90 5/6/90 13/6/90 20/6/90 23/6/90 1/7/90 neutrophils:8.89 5.63 1.92 0.04 1.22 5.11

Additional Information:

Report Detail

Case Number: 66581 **Seq:** 1 **Gender:** M
Reported: 11/07/1990 **Weight:** 84.00
Hospitalisation: **Age:** 30Y
Onset Date: 20/06/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CLAFORAN (Suspected)			**Reason:**	
Injection	4.0 Gram		Daily	Intravenous
Batch:	**Started:** 17/06/1990		**Stopped:**	20/06/1990
GENTAMICIN SULPHATE (Suspected)			**Reason:**	
Injection	300.0 Milligram		Daily	Intravenous
Batch:	**Started:** 17/06/1990		**Stopped:**	20/06/1990

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 66583 **Seq:** 1 **Gender:** F
Reported: 12/07/1990 **Weight:**
Hospitalisation: **Age:** 80Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			

Medicine Details:

RANITIDINE (Suspected) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 06/07/1990

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 03/07/1990

ATENOLOL (Suspected) **Reason:**
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 03/07/1990

HALOPERIDOL (Suspected) **Reason:**
Injection 5.0 Milligram Daily Intramuscular
Batch: **Started:** 28/06/1990 **Stopped:** 03/07/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 66824 **Seq:** 1 **Gender:** F
Reported: 04/08/1990 **Weight:** 54.00
Hospitalisation: **Age:** 66Y
Onset Date: 19/06/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Hepatocellular damage Nausea Pruritus			

Medicine Details:

VALIUM (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

PRIMOLUT-N (Other drug) **Reason:**
0.0
Batch: **Started:** 01/04/1990 **Stopped:**

CORDILOX (Other drug) **Reason:**
0.0
Batch: **Started:** 01/04/1990 **Stopped:**

DONNATAB (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				20/06/90, 26/06/90, 03/07/90, 01/08/90 bili (0-20) 18- 11 12 5 ggt (0-35) 175 105 67 35 ast (0-40) 164 29 15 15 alk (30-120) 83 66 63 62

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 66824 **Seq:** 1 **Gender:** F
Reported: 04/08/1990 **Weight:** 54.00
Hospitalisation: **Age:** 66Y
Onset Date: 19/06/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected)			**Reason:**	Lumbalgia	
	0.0				
Batch:	**Started:**			**Stopped:**	21/06/1990
ORUDIS SR (Suspected)			**Reason:**	Osteoarthritis	
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**	22/05/1990		**Stopped:**	20/06/1990

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 66940 **Seq:** 1 **Gender:** M
Reported: 14/08/1990 **Weight:** 102.00
Hospitalisation: **Age:** 50Y
Onset Date: 20/06/1990 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance			

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 15/06/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 67124 **Seq:** 1 **Gender:** F
Reported: 29/08/1990 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 19/07/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Jaundice			

Medicine Details:

FORANE (Suspected) **Reason:** Other disturbance of sensation
Inhalation 0.0 Inhalation
Batch: **Started:** 16/07/1990 **Stopped:** 16/07/1990

THIOPENTONE SODIUM (Suspected) **Reason:** Other disturbance of sensation
Injection 325.0 Milligram Daily Intravenous
Batch: **Started:** 16/07/1990 **Stopped:** 16/07/1990

PAVULON (Suspected) **Reason:** Other disturbance of sensation
Injection 6.0 Milligram Daily Intravenous
Batch: **Started:** 16/07/1990 **Stopped:** 16/07/1990

MEFOXIN (Suspected) **Reason:**
Injection 4.0 Gram Daily Intravenous
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				06/07, 20/07, 24/07, 27/07, 30/07, 03/08 sgot/ast (12-40) 48 200 68 32 27 20 sgpt/alt (0-30) 13 147 87 64 40 23 ldh (95-365) 155 361 250 257 234 159 alk (35-100) 87 197 288 203 171 131
	Misclassified at [illegible]				Ggt (0-40) 8 160 362 241 164 100 bili (0-19) 5 12 8 9 8 6

Additional Information:

Report Detail

Case Number: 67124 **Seq:** 1 **Gender:** F
Reported: 29/08/1990 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 19/07/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Oral application	25.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 67274 **Seq:** 1 **Gender:** F
Reported: 05/09/1990 **Weight:** 50.00
Hospitalisation: **Age:** 48Y
Onset Date: 17/08/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haematuria			

Medicine Details:

PROGYNOVA (Suspected) **Reason:**
Tablet 1.0 Milligram Daily Oral
Batch: **Started:** 13/08/1990 **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 17/08/1990 **Stopped:** 18/08/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 67336 **Seq:** 1 **Gender:** M
Reported: 31/07/1990 **Weight:** 51.00
Hospitalisation: **Age:** 82Y
Onset Date: 21/07/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous			Ceased serenace,started application of topical celestone-m ointment daily and paraffin cream bd to rash

Medicine Details:

SERENACE (Suspected) **Reason:** Unspecified psychosis
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** 11/07/1990 **Stopped:**

SERENACE (Suspected) **Reason:** Unspecified psychosis
Oral application 15.0 Milligram Daily Oral
Batch: **Started:** 12/07/1990 **Stopped:** 24/07/1990

TRYPTANOL (Suspected) **Reason:** Incontinence of urine
30.0 Milligram Daily
Batch: **Started:** 19/03/1990 **Stopped:** 17/07/1990

MELLERIL (Suspected) **Reason:** Disturbance of sleep
50.0 Milligram Daily
Batch: **Started:** 05/06/1990 **Stopped:** 11/07/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 68124 **Seq:** 1
Reported: 31/10/1990
Hospitalisation:
Onset Date: 24/10/1990
Outcome: Recovered with sequelae

Gender: F
Weight: 72.00
Age: 30Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Chest pain Tachycardia			Sequelae-slight chest pain

Medicine Details:

PANADEINE (Other drug) **Reason:** Migraine
6.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** L TERM

LAROXYL (Suspected) **Reason:** Migraine
150.0 Milligram Total
Batch: **Started:** 23/10/1990 **Stopped:** 27/10/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 68177 **Seq:** 1
Reported: 07/11/1990
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 85.00
Age: 40Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema			Lasix & slow-k. ceased deseril.

Medicine Details:

DESERIL (Suspected) **Reason:** Migraine
Oral application 6.0 Milligram Daily Oral
Batch: **Started:** 01/06/1990 **Stopped:** 28/09/1990

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Migraine
200.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

MAGMIN (Suspected) **Reason:** Migraine
1.5 Gram Daily
Batch: **Started:** 01/06/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 68623 **Seq:** 1 **Gender:** M
Reported: 04/12/1990 **Weight:** 64.00
Hospitalisation: **Age:** AD
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Dysgeusia Nightmare			

Medicine Details:

CAPSIG (Other drug) **Reason:**
Ointment 0.0 Topical
Batch: **Started:** 29/10/1990 **Stopped:**

AMPRACE (Other drug) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 11/01/1990 **Stopped:** CONTIN

PROTAPHANE HM (Other drug) **Reason:**
Injection 10.0 Millilitre Daily Subcutaneous
Batch: **Started:** 01/03/1990 **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** 05/11/1990 **Stopped:** 09/11/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 68696 **Seq:** 1 **Gender:** F
Reported: 04/12/1990 **Weight:** 60.00
Hospitalisation: **Age:** 42Y
Onset Date: 17/05/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			Tryptanol reduced to 25mg

Medicine Details:

TENORMIN (Other drug) **Reason:** Essential benign hypertension
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** 06/04/1990 **Stopped:** 20/11/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry Misclassified at				17/5/90 23/8/90 r.r. bilirubin 9 8 (2-20) alk.phos 112 127 (30-115) ggt 72 181 (up to 45) alt 23 51 (up to 45) Ast 20 50 (up to 41)

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 68909 **Seq:** 1 **Gender:** M
Reported: 28/12/1990 **Weight:**
Hospitalisation: **Age:** 52Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

0.0

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				Documented decrease in testosterone after commencing amitriptyline. recovered to normal on ceasing and recurred on

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 69138 **Seq:** 1 **Gender:** M
Reported: 15/01/1991 **Weight:** 56.00
Hospitalisation: **Age:** 63Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension			

Medicine Details:

CAPOTEN (Suspected) **Reason:** Congestive heart failure
25.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** 01/07/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 69837 **Seq:** 1 **Gender:** F
Reported: 28/02/1991 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 20/12/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hyponatraemia Syncope			Hyperosmotic saline infusion, fluid restriction.

Medicine Details:

DI-GESIC (Other drug) **Reason:**
Tablet 6.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:** CONTIN

LANOXIN PG (Other drug) **Reason:**
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** CONTIN

MODURETIC (Suspected) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 18/12/1990

TRYPTANOL (Suspected) **Reason:**
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 25/12/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrolytes				Patient had serum sodium of 105mmol/l.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70290 **Seq:** 1
Reported: 07/04/1991
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight: 90.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Malaise Nausea Periorbital oedema			

Medicine Details:

ZYLOPRIM (Other drug) **Reason:** Gout
400.0 Milligram Daily
Batch: **Started:** 01/01/1970 **Stopped:**

APTIN (Other drug) **Reason:** Chron isch heart dis no hyper
200.0 Milligram Daily
Batch: **Started:** 01/01/1986 **Stopped:**

MODURETIC (Other drug) **Reason:** Chron isch heart dis no hyper
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1986 **Stopped:**

CLINORIL (Suspected) **Reason:** Osteoarthritis
400.0 Milligram Daily
Batch: **Started:** 01/08/1990 **Stopped:** 28/11/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70290 **Seq:** 1
Reported: 07/04/1991
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight: 90.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
100.0 Milligram Daily
Batch: **Started:** 01/01/1984 **Stopped:**

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70357 **Seq:** 1 **Gender:** F
Reported: 11/04/1991 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 31/03/1991 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bone marrow failure			

Medicine Details:

PREDNISONE (Other drug) **Reason:** Other rheumatoid arthritis
Tablet 8.0 Milligram Daily Oral
Batch: **Started:** 01/01/1977 **Stopped:**

MOGADON (Other drug) **Reason:**
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

METHOTREXATE (Suspected) **Reason:** Other rheumatoid arthritis
Tablet 12.0 Milligram Weekly Oral
Batch: **Started:** 15/11/1990 **Stopped:** 28/03/1991

FELDENE (Suspected) **Reason:** Other rheumatoid arthritis
Capsule 20.0 Milligram Daily Oral
Batch: **Started:** 29/11/1989 **Stopped:** 28/03/1991

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70357 **Seq:** 1 **Gender:** F
Reported: 11/04/1991 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 31/03/1991 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 04/11/1980 **Stopped:** 28/03/1991

Laboratory Investigations:

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 70362 **Seq:** 1 **Gender:** F
Reported: 11/04/1991 **Weight:** 55.00
Hospitalisation: **Age:** 54Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myopathy			

Medicine Details:

CAPTOPRIL (Suspected) **Reason:** Essential benign hypertension
75.0 Milligram Daily
Batch: **Started:** 27/11/1990 **Stopped:** 23/01/1991

VERAPAMIL HYDROCHLORIDE (Suspected) **Reason:** Essential benign hypertension
240.0 Milligram Daily
Batch: **Started:** 27/11/1990 **Stopped:** 23/01/1991

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
125.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Histology				No evidence of an inflammatory process is present. moderately severe type 2b fibre atrophy is noted, which although non-specific, is commonly seen in disuse, cachexia, thyroid abnormalities and steroid myopathy

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70380 **Seq:** 1 **Gender:** M
Reported: 08/04/1991 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 18/03/1991 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Nightmare			

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 07/03/1991 **Stopped:** 18/03/1991

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Generic substitution problem

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70837 **Seq:** 1
Reported: 03/05/1991
Hospitalisation:
Onset Date: 29/04/1991
Outcome: Not yet recovered

Gender: M
Weight: 54.00
Age: 64Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Otr periph nerve dis not auton

50.0 Milligram Daily

Batch: **Started:** 26/04/1991 **Stopped:** 29/04/1991

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 70854 **Seq:** 1 **Gender:** F
Reported: 17/05/1991 **Weight:** 60.00
Hospitalisation: **Age:** 76Y
Onset Date: 10/05/1991 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pollakiuria Urinary incontinence			

Medicine Details:

MINIPRESS (Suspected) **Reason:** Essential benign hypertension
1.0 Milligram Daily
Batch: **Started:** 08/05/1991 **Stopped:** 17/05/1991

TENORMIN (Suspected) **Reason:** Essential benign hypertension
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 15/05/1990 **Stopped:**

ZOCOR (Suspected) **Reason:** Othr&unspec metabolic diseases
10.0 Milligram Daily
Batch: **Started:** 05/02/1991 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 70854 **Seq:** 1
Reported: 17/05/1991
Hospitalisation:
Onset Date: 10/05/1991
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 76Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

VALIUM (Suspected) **Reason:** Anxiety neurosis
2.5 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 71037 **Seq:** 1 **Gender:** F
Reported: 28/05/1991 **Weight:** 60.00
Hospitalisation: **Age:** 78Y
Onset Date: 05/05/1991 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Glaucoma			Needed laser peripheral

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 05/05/1991 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 71666 **Seq:** 1
Reported: 17/07/1991
Hospitalisation: Admitted to hospital
Onset Date: 10/04/1991
Outcome: Not yet recovered

Gender: M
Weight:
Age: 72Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysphagia Oesophagitis ulcerative Duodenal ulcer		Difficulty in swallowing Oesophageal ulcers	

Medicine Details:

ASPIRIN (Suspected) **Reason:** Acute myocard infarc,no hypert
150.0 Milligram Daily
Batch: **Started:** 03/04/1991 **Stopped:**

INDOCID (Suspected) **Reason:** Gout
100.0 Milligram Daily
Batch: **Started:** 08/04/1991 **Stopped:** 27/06/1991

TRYPTANOL (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

SEREPAX (Suspected) **Reason:**
15.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Endoscopy- very aggressive peptic ulcer disease- multiple oesaphageal ulcers and du.

Additional Information:

Report Detail

Case Number: 71972 **Seq:** 1 **Gender:** F
Reported: 29/07/1991 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Abnormal lft's	

Medicine Details:

PANADEINE (Other drug) **Reason:**
Oral application 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** 01/06/1991 **Stopped:** CONTIN

PHENYTOIN SODIUM (Suspected) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 24/05/1991 **Stopped:** 07/06/1991

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 06/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	<55 U/L	19/05/1991	6	
	ALT = SGPT	<55 U/L	05/06/1991	129	
	ALT = SGPT	<55 U/L	11/06/1991	78	
	ALT = SGPT	<55 U/L	06/06/1991	107	
	ALT = SGPT	<55 U/L	26/06/1991	16	
	ALT = SGPT	<55 U/L	17/06/1991	31	
	AST = SGOT	3-65 U/L	19/05/1991	11	

Additional Information:

Report Detail

Case Number: 71972 **Seq:** 1 **Gender:** F
Reported: 29/07/1991 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detai

Medicine Details:

PANADEINE (Other drug)			**Reason:**		
Oral application	2.0	Dose Unspecified	As necessary	Oral	
Batch:	**Started:**	01/06/1991	**Stopped:**		CONTIN
PHENYTOIN SODIUM (Suspected)			**Reason:**		
Oral application	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	24/05/1991	**Stopped:**	07/06/1991	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Oral application	20.0	Milligram	Daily	Oral	
Batch:	**Started:**	L TERM	**Stopped:**	06/06/1990	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	3-65 U/L	05/06/1991	34	
	AST = SGOT	3-65 U/L	11/06/1991	30	
	AST = SGOT	3-65 U/L	17/06/1991	17	
	AST = SGOT	3-65 U/L	06/06/1991	39	
	AST = SGOT	3-65 U/L	26/06/1991	16	
	Bilirubin	<17 UMOL/L	19/05/1991	8	
	Bilirubin	<17 UMOL/L	05/06/1991	4	

Additional Information:

Report Detail

Case Number: 71972 **Seq:** 1 **Gender:** F
Reported: 29/07/1991 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detai

Medicine Details:

PANADEINE (Other drug)			**Reason:**		
Oral application	2.0	Dose Unspecified	As necessary	Oral	
Batch:	**Started:**	01/06/1991	**Stopped:**		CONTIN
PHENYTOIN SODIUM (Suspected)			**Reason:**		
Oral application	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	24/05/1991	**Stopped:**	07/06/1991	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Oral application	20.0	Milligram	Daily	Oral	
Batch:	**Started:**	L TERM	**Stopped:**	06/06/1990	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	<17 UMOL/L	06/06/1991	4	
	Bilirubin	<17 UMOL/L	26/06/1991	4	
	Bilirubin	<17 UMOL/L	17/06/1991	3	
	Bilirubin	<17 UMOL/L	11/06/1991	3	
	GGT = SGGT =	7-43 U/L	06/06/1991	672	
	GGT = SGGT =	7-43 U/L	26/06/1991	198	
	GGT = SGGT =	7-43 U/L	17/06/1991	360	

Additional Information:

Report Detail

Case Number: 71972 **Seq:** 1 **Gender:** F
Reported: 29/07/1991 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

PANADEINE (Other drug)			**Reason:**		
Oral application	2.0	Dose Unspecified	As necessary	Oral	
Batch:	**Started:**	01/06/1991	**Stopped:**		CONTIN
PHENYTOIN SODIUM (Suspected)			**Reason:**		
Oral application	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	24/05/1991	**Stopped:**	07/06/1991	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Oral application	20.0	Milligram	Daily	Oral	
Batch:	**Started:**	L TERM	**Stopped:**	06/06/1990	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT =	7-43 U/L	11/06/1991	531	
	SAP = ALP	30-110 U/L	19/05/1991	62	
	SAP = ALP	30-110 U/L	05/06/1991	248	
	SAP = ALP	30-110 U/L	06/06/1991	238	
	SAP = ALP	30-110 U/L	11/06/1991	154	
	SAP = ALP	30-110 U/L	17/06/1991	109	
	SAP = ALP	30-110 U/L	26/06/1991	80	

Additional Information:

Report Detail

Case Number: 72410 **Seq:** 1 **Gender:** F
Reported: 28/08/1991 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: 04/08/1991 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura Epistaxis Thrombocytopenia		Bruising of lips	Prednisone

Medicine Details:

ERGODRYL (Other drug) **Reason:** Migraine
0.0
Batch: **Started:** **Stopped:**

QUININE (Suspected) **Reason:** Pain in limb
0.0
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 72745 **Seq:** 1 **Gender:** F
Reported: 08/09/1991 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 29/06/1991 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cheilitis		Dry lips.	
Dry mouth		Dry mouth.	
Muscle twitching		Twitchiness	
Agitation			
Insomnia			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 28/06/1991 **Stopped:** 30/06/1991

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 73272 **Seq:** 1 **Gender:** F
Reported: 03/10/1991 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 01/10/1991 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Duodenal ulcer perforation			

Medicine Details:

PANADEINE FORTE (Suspected) **Reason:**
2.0 Dose Unspecified As necessary
Batch: **Started:** L TERM **Stopped:**

TENOXICAM (Suspected) **Reason:** Spondylitis osteoarthritica
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 03/12/1990 **Stopped:** 01/10/1991

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 73489 **Seq:** 1 **Gender:** M
Reported: 15/10/1991 **Weight:**
Hospitalisation: **Age:** 63Y
Onset Date: 31/12/1990 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cardiomyopathy Left ventricular failure	Severe		

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Migraine

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** 01/01/1980 **Stopped:** 31/01/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Report Detail

Case Number: 73810 **Seq:** 1
Reported: 25/10/1991
Hospitalisation:
Onset Date: 03/09/1991
Outcome: Death, maybe drug

Gender: F
Weight: 86.00
Age: 64
DOB: 14/01/1927
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Encephalopathy Hepatic failure			

Medicine Details:

SODIUM VALPROATE (Suspected) **Reason:**
0.0
Batch: **Started:** 28/08/1991 **Stopped:**

VENTOLIN (Suspected) **Reason:**
6.0 Dose Unspecified Daily
Batch: **Started:** 16/08/1991 **Stopped:**

BECOTIDE (Suspected) **Reason:**
4.0 Dose Unspecified Daily
Batch: **Started:** 16/08/1991 **Stopped:**

ASCORBIC ACID (Suspected) **Reason:**
500.0 Milligram Daily
Batch: **Started:** 18/07/1991 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient died on the 10/9/1991. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 73810 **Seq:** 1
Reported: 25/10/1991
Hospitalisation:
Onset Date: 03/09/1991
Outcome: Death, maybe drug

Gender: F
Weight: 86.00
Age: 64
DOB: 14/01/1927
Causality: Causality possible

Reaction Detail

Medicine Details:

TRANSIDERM-NITRO (Suspected)			**Reason:**	Angina pectoris w/o hyperten
	25.0 Milligram		Daily	
Batch:	**Started:** 01/05/1991			**Stopped:**
PROVERA (Suspected)			**Reason:**	Neoplasm unspecified of uterus
	400.0 Milligram		Daily	
Batch:	**Started:**	L TERM		**Stopped:**
TRYPTANOL (Suspected)			**Reason:**	Depression
Tablet	150.0 Milligram		Daily	Oral
Batch:	**Started:** 18/07/1991			**Stopped:**

Laboratory Investigations:

Additional Information:

Patient died on the 10/9/1991. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 74318 **Seq:** 1 **Gender:** F
Reported: 21/11/1991 **Weight:** 60.00
Hospitalisation: **Age:** 35Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia Tinnitus Tremor		Tingling in hands and around mouth	

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
100.0 Milligram Daily
Batch: **Started:** 29/08/1991 **Stopped:** 14/11/1991

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 74829 **Seq:** 1 **Gender:** F
Reported: 20/12/1991 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 17/12/1991 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Nausea Pruritus			Reaction reversed over 4 hours by a stat im injection of promethazine 25mg.

Medicine Details:

ANTENEX (Other drug) **Reason:** Depression
7.5 Milligram Daily
Batch: **Started:** 16/12/1991 **Stopped:**

ENDEP (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 17/12/1991 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallanged on the 18/12/1991 with same reaction occuring.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 75205 **Seq:** 1
Reported: 17/01/1992
Hospitalisation:
Onset Date: 18/12/1991
Outcome: Not yet recovered

Gender: M
Weight: 60.00
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paresis Gait disturbance		Proximal muscle weakness. Shuffling gait.	Physiotherapy

Medicine Details:

CAPSIG (Suspected) **Reason:** Other disturbance of sensation
Ointment 25.0 Milligram As necessary Topical
Batch: **Started:** 20/08/1991 **Stopped:**

ENDONE (Suspected) **Reason:**
Tablet 5.0 Milligram As necessary Oral
Batch: **Started:** 15/08/1991 **Stopped:** 02/01/1992

ALODORM (Suspected) **Reason:** Specific disorders of sleep
Tablet 5.0 Milligram As necessary Oral
Batch: **Started:** L TERM **Stopped:**

EPILIM (Suspected) **Reason:** Other disturbance of sensation
Oral application 600.0 Milligram Daily Oral
Batch: **Started:** 17/12/1991 **Stopped:** 06/01/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 75205 **Seq:** 1
Reported: 17/01/1992
Hospitalisation:
Onset Date: 18/12/1991
Outcome: Not yet recovered

Gender: M
Weight: 60.00
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MEXITIL (Suspected)				**Reason:**	Other disturbance of sensation	
Capsule	150.0	Milligram	Daily		Oral	
Batch:	**Started:**	26/07/1991		**Stopped:**	11/12/1991	
ENDEP (Suspected)				**Reason:**	Other disturbance of sensation	
Tablet	100.0	Milligram	Daily		Oral	
Batch:	**Started:**	19/08/1991		**Stopped:**	13/01/1992	
ZANTAC (Suspected)				**Reason:**	Ulcer of duodenum w/o ment perf	
Oral application	300.0	Milligram	Daily		Oral	
Batch:	**Started:**	01/12/1990		**Stopped:**		
PILOPT (Suspected)				**Reason:**	Unspecified glaucoma	
Eye Drops	4.0	Dose Unspecified	Daily		Ophthalmic	
Batch:	**Started:**	L TERM		**Stopped:**		

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 75205 **Seq:** 1
Reported: 17/01/1992
Hospitalisation:
Onset Date: 18/12/1991
Outcome: Not yet recovered

Gender: M
Weight: 60.00
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

BETOPTIC (Suspected)			**Reason:**	Unspecified glaucoma	
Eye Drops	2.0	Dose Unspecified	Daily		Ophthalmic
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 75639 **Seq:** 1 **Gender:** M
Reported: 19/02/1992 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction		Gross rash on skin from sun.	Dilosyn and calestone v 1/2 cream, paraffin cream

Medicine Details:

MODURETIC (Suspected) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1992 **Stopped:** 02W

NATRILIX (Suspected) **Reason:** Essential benign hypertension
2.5 Milligram Daily
Batch: **Started:** **Stopped:** 19/02/1992

MINIPRESS (Suspected) **Reason:** Essential benign hypertension
6.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Nervousness
75.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 75639 **Seq:** 1 **Gender:** M
Reported: 19/02/1992 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ZANTAC (Suspected)			**Reason:**	
	300.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
WARFARIN SODIUM (Suspected)			**Reason:**	Phlebitis and thrombophlebitisof intracranial ven
	5.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 75848 **Seq:** 1
Reported: 05/03/1992
Hospitalisation: Admitted to hospital
Onset Date: 19/11/1991
Outcome: Recovered

Gender: F
Weight:
Age: 81
DOB: 30/12/1909
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Leukopenia Extrapyramidal disorder Dementia		Low wwc Parkinsonism.	

Medicine Details:

MELLERIL (Suspected) **Reason:**
30.0 Milligram Daily
Batch: **Started:** **Stopped:**

MELLERIL (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** 19/11/1991 **Stopped:** 21/11/1991

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

CLONAZEPAM (Suspected) **Reason:**
500.0 Microgram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	White blood cells		20/11/1991	2.9	
	White blood cells		25/11/1991	4.8	

Additional Information:

See original report for other medication.

Report Detail

Case Number: 75983 **Seq:** 1
Reported: 13/03/1992
Hospitalisation:
Onset Date: 19/02/1992
Outcome: Not yet recovered

Gender: M
Weight: 75.00
Age: 70Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Duodenal ulcer			Losec 1 daily.

Medicine Details:

CORDILOX (Suspected) **Reason:** Essential benign hypertension
Tablet 320.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

DIAZEPAM (Suspected) **Reason:** Otr&nos disord of heart rhythm
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

CLINORIL (Suspected) **Reason:** Osteoarthritis
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 19/02/1992

FELDENE (Suspected) **Reason:** Osteoarthritis
20.0 Milligram Daily
Batch: **Started:** 19/02/1992 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 75983 **Seq:** 1
Reported: 13/03/1992
Hospitalisation:
Onset Date: 19/02/1992
Outcome: Not yet recovered

Gender: M
Weight: 75.00
Age: 70Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Depression
	75.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	
MODURETIC (Suspected)			**Reason:**	Essential benign hypertension
	1.0	Dose Unspecified	Daily	
Batch:	**Started:**		**Stopped:**	04/02/1991
ZANTAC (Suspected)			**Reason:**	
	150.0	Milligram	Daily	
Batch:	**Started:**	01/01/1989	**Stopped:**	26/12/1991

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 76077 **Seq:** 1 **Gender:** M
Reported: 16/03/1992 **Weight:** 63.00
Hospitalisation: **Age:** 70Y
Onset Date: 11/02/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis Jaundice			

Medicine Details:

ROHYPNOL (Other drug) **Reason:** Specific disorders of sleep
2.0 Milligram Daily
Batch: **Started:** 01/01/1988 **Stopped:**

TENORMIN (Other drug) **Reason:** Palpitation
20.0 Milligram Daily
Batch: **Started:** 01/01/1989 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Headache
50.0 Milligram Daily
Batch: **Started:** 17/10/1991 **Stopped:** 14/02/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 76101 **Seq:** 1 **Gender:** F
Reported: 23/03/1992 **Weight:** 43.00
Hospitalisation: **Age:** 85Y
Onset Date: 12/03/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Tremor	 Severe	Unsteady on feet	

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Specific disorders of sleep

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 07/03/1992 **Stopped:** 14/03/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 76470 **Seq:** 1 **Gender:** F
Reported: 03/04/1992 **Weight:** 70.00
Hospitalisation: **Age:** 41Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chloasma			

Medicine Details:

VOLTAREN (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

RIFAMPICIN (Suspected) **Reason:**
Injection 0.0 Intravenous
Batch: **Started:** 01/10/1991 **Stopped:**

FUCIDIN (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

TRENTAL 400 (Suspected) **Reason:** Chronic ulcer of skin
3.0 Dose Unspecified Daily
Batch: **Started:** 02/01/1992 **Stopped:** 18/02/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 76470 **Seq:** 1
Reported: 03/04/1992
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 41Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected)			**Reason:** Pain	
	75.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
FERRO-GRADUMET (Suspected)			**Reason:**	
	2.0	Dose Unspecified	Daily	
Batch:	**Started:**			**Stopped:**
PANADEINE FORTE (Suspected)			**Reason:**	
	12.0	Dose Unspecified	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 76864 **Seq:** 1 **Gender:** M
Reported: 01/05/1992 **Weight:** 60.00
Hospitalisation: **Age:** 74Y
Onset Date: 19/04/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash		On upper body and arms.	

Medicine Details:

ANGININE (Other drug) **Reason:** Chron isch heart dis no hyper
1.0 Dose Unspecified As necessary
Batch: **Started:** L TERM **Stopped:**

DUPHALAC (Other drug) **Reason:** Constipation
30.0 Millilitre Daily
Batch: **Started:** L TERM **Stopped:**

ISORDIL (Other drug) **Reason:** Chron isch heart dis no hyper
15.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

MAREVAN (Other drug) **Reason:** Acute,ill-def cerebrovascular disease w/o hyperte
5.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 76864 **Seq:** 1 **Gender:** M
Reported: 01/05/1992 **Weight:** 60.00
Hospitalisation: **Age:** 74Y
Onset Date: 19/04/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

DYMADON (Other drug)			**Reason:**	
	4.0	Dose Unspecified	Daily	
Batch:	**Started:**	L TERM	**Stopped:**	
ENDEP (Suspected)			**Reason:**	Depression
	10.0	Milligram	Daily	
Batch:	**Started:**	16/04/1992	**Stopped:**	19/04/1992

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 77010 **Seq:** 1 **Gender:** F
Reported: 14/05/1992 **Weight:** 80.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertriglyceridaemia Drug ineffective Hypercholesterolaemia		Amazing jump in cholesterol and	

Medicine Details:

ATROMID-S (Other drug) **Reason:**
1.5 Gram Daily
Batch: **Started:** 22/02/1989 **Stopped:** 30/12/1991

STELAZINE (Suspected) **Reason:** Unspecifid affective psychosis
0.0
Batch: **Started:** 01/01/1985 **Stopped:**

LOPID (Suspected) **Reason:** Othr&unspec metabolic diseases
120.0 Milligram Daily
Batch: **Started:** 30/12/1991 **Stopped:** 14/05/1992

ASPIRIN (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Cholesterol		22/02/1989	8	
	Cholesterol		29/05/1990	9.9	
	Cholesterol		10/09/1990	9.1	
	Cholesterol		06/01/1991	5.5	
	Cholesterol		13/04/1992	15	
	Cholesterol		27/04/1992	19	
	Triglyceride		22/02/1989	4	

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 77010 **Seq:** 1 **Gender:** F
Reported: 14/05/1992 **Weight:** 80.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

CAPOTEN (Suspected) **Reason:** Essential benign hypertension
50.0 Milligram Daily
Batch: **Started:** 04/10/1991 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Unspecifid affective psychosis
75.0 Milligram Daily
Batch: **Started:** 01/01/1985 **Stopped:**

BETALOC (Suspected) **Reason:** Essential benign hypertension
50.0 Milligram Daily
Batch: **Started:** 28/08/1990 **Stopped:**

PROGYNOVA (Suspected) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** 13/11/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Triglyceride		29/05/1990	-	
	Triglyceride		10/09/1990	6.3	
	Triglyceride		06/01/1991	3.4	
	Triglyceride		13/04/1992	23	
	Triglyceride		27/04/1992	31	

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 77634 **Seq:** 1
Reported: 08/06/1992
Hospitalisation: Admitted to hospital
Onset Date: 28/10/1991
Outcome: Recovered

Gender: F
Weight:
Age: 83
DOB: 11/11/1907
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise Myalgia Pyrexia Urticaria			Sorbolene cream with glycerin, atarax 75mg daily,

Medicine Details:

FELODIPINE (Suspected) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Frequency of micturition
10.0 Milligram Daily
Batch: **Started:** 14/10/1991 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 77803 **Seq:** 1 **Gender:** M
Reported: 24/06/1992 **Weight:** 60.00
Hospitalisation: **Age:** 43Y
Onset Date: 11/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Speech disorder Aggression		Rapid speech	

Medicine Details:

ZINVIT (Other drug) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

PANADEINE FORTE (Other drug) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 03D

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 77942 **Seq:** 1 **Gender:** F
Reported: 29/06/1992 **Weight:** 55.00
Hospitalisation: **Age:** 78Y
Onset Date: 05/02/1992 **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Metastases to bone marrow		Metastatic adenocarcinoma to bone marrow, primary site renal	

Medicine Details:

OMEPRAZOLE (Suspected)			**Reason:**	Other diseases of esophagus
	40.0 Milligram	Daily		Oral
Batch:	**Started:** 15/03/1988		**Stopped:**	13/04/1988
OMEPRAZOLE (Suspected)			**Reason:**	Other diseases of esophagus
	20.0 Milligram	Daily		Oral
Batch:	**Started:** 14/04/1988		**Stopped:**	
FOLIC ACID (Suspected)			**Reason:**	
	15.0 Milligram	Daily		Oral
Batch:	**Started:**		**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	
	50.0 Milligram	Daily		Oral
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had cervical cancer in 1970s, colonic cancer in 1980s.

Report Detail

Case Number: 77942 **Seq:** 1
Reported: 29/06/1992
Hospitalisation:
Onset Date: 05/02/1992
Outcome: Death, maybe drug

Gender: F
Weight: 55.00
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

OXAZEPAM (Suspected)			**Reason:**	
	30.0	Milligram	Daily	Oral
Batch:	**Started:**		**Stopped:**	
PHENERGAN (Suspected)			**Reason:**	
	20.0	Milligram	Daily	Oral
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Additional Information:

Patient had cervical cancer in 1970s, colonic cancer in 1980s.

Report Detail

Case Number: 78506 **Seq:** 1 **Gender:** M
Reported: 24/07/1992 **Weight:** 82.00
Hospitalisation: **Age:** 45Y
Onset Date: 31/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Auditory, visual, olfactory hallucinat.	
Consciousness fluctuating		Clouding of consciousness	
Confusional state		Disorientation in time	
Delirium			
Delusion			
Paranoia			

Medicine Details:

BACLOFEN (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 78561 **Seq:** 1 **Gender:** F
Reported: 04/08/1992 **Weight:** 55.00
Hospitalisation: **Age:** 33Y
Onset Date: 20/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hallucination		Disoriented to time and place Visual and tactile hallucinations	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 50.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 78697 **Seq:** 1 **Gender:** F
Reported: 18/08/1992 **Weight:** 57.00
Hospitalisation: **Age:** 71Y
Onset Date: 21/02/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia			

Medicine Details:

VISKEN (Other drug) **Reason:** Essential benign hypertension
Tablet 7.5 Milligram Daily Oral
Batch: **Started:** **Stopped:**

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
Tablet 3.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

GLUCOPHAGE (Other drug) **Reason:** Diabetes mellitus
Tablet 1.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

DIAMICRON (Other drug) **Reason:** Diabetes mellitus
Tablet 240.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 78697 **Seq:** 1 **Gender:** F
Reported: 18/08/1992 **Weight:** 57.00
Hospitalisation: **Age:** 71Y
Onset Date: 21/02/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 21/02/1992 **Stopped:** 25/02/1992

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 78937 **Seq:** 1 **Gender:** U
Reported: 31/07/1992 **Weight:** 55.00
Hospitalisation: **Age:** 80
Onset Date: 17/03/1992 **DOB:** 27/02/1912
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit

Tablet 25.0 Milligram 1 time Oral

Batch: **Started:** 17/03/1992 **Stopped:**

OXAZEPAM (Suspected) **Reason:**

Oral application 60.0 Milligram Total Oral

Batch: **Started:** 16/03/1992 **Stopped:** 17/03/1992

PARACETAMOL (Suspected) **Reason:**

3.0 Dose Unspecified Total

Batch: **Started:** 16/03/1992 **Stopped:** 17/03/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drugs.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 79091 **Seq:** 1 **Gender:** F
Reported: 02/09/1992 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 24/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention			Amitriptyline dose gradually reduced to 150mg at night

Medicine Details:

TEMAZEPAM (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

VALIUM (Other drug) **Reason:**
Tablet 15.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

COLOXYL (Other drug) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

CODRAL FORTE (Other drug) **Reason:**
Tablet 8.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 79091 **Seq:** 1 **Gender:** F
Reported: 02/09/1992 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 24/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 79500 **Seq:** 1 **Gender:** M
Reported: 17/09/1992 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous		Rash affecting legs, arms, back.	

Medicine Details:

SOTACOR (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** 01/03/1992 **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 12.5 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 80275 **Seq:** 1
Reported: 07/10/1992
Hospitalisation: Admitted to hospital
Onset Date: 13/09/1992
Outcome: Not yet recovered

Gender: M
Weight: 74.00
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diabetes mellitus			Actrapid insulin - sliding scale

Medicine Details:

SEPTRIN FORTE (Suspected) **Reason:** Other prophylactic procedures
Tablet 3.0 Dose Unspecified Weekly Oral
Batch: **Started:** 10/07/1991 **Stopped:** CONTIN

MEGOSTAT (Suspected) **Reason:**
Tablet 480.0 Milligram Daily Oral
Batch: **Started:** 25/08/1992 **Stopped:** 13/09/1992

DIFLUCAN (Suspected) **Reason:** Other prophylactic procedures
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** 11/07/1991 L TERM **Stopped:** CONTIN

RETROVIR CAPSULES (Suspected) **Reason:**
Capsule 600.0 Milligram Daily Oral
Batch: **Started:** 10/07/1991 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 80275 **Seq:** 1
Reported: 07/10/1992
Hospitalisation: Admitted to hospital
Onset Date: 13/09/1992
Outcome: Not yet recovered

Gender: M
Weight: 74.00
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	50.0	Milligram	Daily	Oral	
Batch:	**Started:**	16/07/1991		**Stopped:**	CONTIN
VENTOLIN (Suspected)			**Reason:**	Asthma	
Inhalation	4.0	Millilitre	Daily	Inhalation	
Batch:	**Started:**	04/06/1992		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 80699 **Seq:** 1 **Gender:** F
Reported: 29/10/1992 **Weight:** 64.00
Hospitalisation: **Age:** 81Y
Onset Date: 16/10/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Acidosis			

Medicine Details:

ACETAZOLAMIDE (Suspected) **Reason:**
Tablet 500.0 Milligram Daily Oral
Batch: **Started:** 30/01/1992 **Stopped:** 22/10/1992

ENDEP (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

TAMOXIFEN CITRATE (Suspected) **Reason:**
30.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ALLOPURINOL (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrolytes				16/10/92 low s. bicarbonate, 28/10/92 - 20

Additional Information:

Report Detail

Case Number: 80699 **Seq:** 1
Reported: 29/10/1992
Hospitalisation:
Onset Date: 16/10/1992
Outcome: Not yet recovered

Gender: F
Weight: 64.00
Age: 81Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ASCORBIC ACID (Suspected) **Reason:**
500.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 80705 **Seq:** 1 **Gender:** F
Reported: 29/10/1992 **Weight:** 96.00
Hospitalisation: **Age:** 60Y
Onset Date: 26/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Nausea Somnolence Vomiting	Severe		

Medicine Details:

VENTOLIN (Other drug) **Reason:**
0.0
Batch: **Started:** 01/01/1992 **Stopped:**

BECOTIDE (Other drug) **Reason:**
0.0
Batch: **Started:** 01/01/1990 **Stopped:**

OGEN (Other drug) **Reason:** Menopausal symptoms
625.0 Microgram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

PROVERA (Other drug) **Reason:** Menopausal symptoms
10.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 80705 **Seq:** 1 **Gender:** F
Reported: 29/10/1992 **Weight:** 96.00
Hospitalisation: **Age:** 60Y
Onset Date: 26/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Pain
10.0 Milligram Daily
Batch: **Started:** 26/10/1992 **Stopped:** 26/10/1992

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 81070 **Seq:** 1 **Gender:** F
Reported: 23/11/1992 **Weight:** 80.00
Hospitalisation: **Age:** 65Y
Onset Date: 06/07/1992 **DOB:**
Outcome: Recovered with sequelae **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis			

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 16/11/1987 **Stopped:** 06/07/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 81211 **Seq:** 1 **Gender:** M
Reported: 26/11/1992 **Weight:** 75.00
Hospitalisation: **Age:** 34Y
Onset Date: 03/09/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Malaise		Polyarthralgia	Piroxicam, prednisolone.

Medicine Details:

LITHIUM CARBONATE (Suspected) **Reason:**
0.0
Batch: **Started:** 01/08/1992 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** 01/11/1988 **Stopped:** 07/09/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 81687 **Seq:** 1
Reported: 24/11/1992
Hospitalisation:
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 69
DOB: 04/10/1923
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic failure Hepatitis			Icu

Medicine Details:

BROMOCRIPTINE MESYLATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 3M

SINEMET (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

LITHIUM CARBONATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 81687 **Seq:** 1
Reported: 24/11/1992
Hospitalisation:
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 69
DOB: 04/10/1923
Causality: Causality possible

Reaction Detail

Medicine Details:

PANADOL (Suspected)		**Reason:**
	0.0	
Batch:	**Started:**	**Stopped:**
CODEINE (Suspected)		**Reason:**
	0.0	
Batch:	**Started:**	**Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 81815 **Seq:** 1 **Gender:** F
Reported: 24/12/1992 **Weight:**
Hospitalisation: **Age:** 56
Onset Date: 26/11/1992 **DOB:** 25/09/1936
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular			Teldane, hydrocortisone.

Medicine Details:

PREDNISONE (Suspected) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

CYCLOPHOSPHAMIDE (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** 27/10/1992 **Stopped:** 14/12/1992

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** **Stopped:** 08/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83054 **Seq:** 1
Reported: 12/03/1993
Hospitalisation: Admitted to hospital
Onset Date: 02/03/1993
Outcome: Not yet recovered

Gender: M
Weight:
Age: 64Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Dyspnoea Hepatitis			

Medicine Details:

ATROVENT (Other drug) **Reason:**
Inhalation 0.0 Inhalation
Batch: **Started:** **Stopped:**

CAPOTEN (Other drug) **Reason:** Cardiomyopathy
75.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

LASIX (Other drug) **Reason:** Cardiomyopathy
120.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

SLOW-K (Other drug) **Reason:** Cardiomyopathy
6.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83054 **Seq:** 1
Reported: 12/03/1993
Hospitalisation: Admitted to hospital
Onset Date: 02/03/1993
Outcome: Not yet recovered

Gender: M
Weight:
Age: 64Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

OROXINE (Other drug)				**Reason:**	Cardiomyopathy
	200.0	Microgram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
LANOXIN PG (Other drug)				**Reason:**	Cardiomyopathy
	2.0	Dose Unspecified	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
VENTOLIN (Other drug)				**Reason:**	
Inhalation	6.0	Dose Unspecified	Daily		Inhalation
Batch:	**Started:**			**Stopped:**	
METOLAZONE (Other drug)				**Reason:**	Cardiomyopathy
	2.5	Milligram	Alternate days		
Batch:	**Started:**		L TERM	**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 83054 **Seq:** 1
Reported: 12/03/1993
Hospitalisation: Admitted to hospital
Onset Date: 02/03/1993
Outcome: Not yet recovered

Gender: M
Weight:
Age: 64Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

CORDARONE X 200 (Suspected)			**Reason:**	Cardiomyopathy	
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**	20/02/1993		**Stopped:**	05/03/1993
TRYPTANOL (Suspected)			**Reason:**	Cardiomyopathy	
	50.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	05/03/1993
QUININE SULPHATE (Suspected)			**Reason:**	Cardiomyopathy	
	300.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	05/03/1993
NORMISON (Suspected)			**Reason:**	Cardiomyopathy	
	20.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	05/03/1993

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 83240 **Seq:** 1 **Gender:** F
Reported: 12/03/1993 **Weight:** 80.00
Hospitalisation: **Age:** 66Y
Onset Date: 29/05/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Confusional state Urinary retention			

Medicine Details:

DI-GESIC (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

DOLOBID (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

DIAZEPAM (Other drug) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 30/05/1992

SLOW-K (Other drug) **Reason:**
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 83240 **Seq:** 1 **Gender:** F
Reported: 12/03/1993 **Weight:** 80.00
Hospitalisation: **Age:** 66Y
Onset Date: 29/05/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CHLOTRIDE (Other drug)				**Reason:**		
Tablet	500.0	Milligram		Alternate days		Oral
Batch:	**Started:**				**Stopped:**	
BENZTROPINE MESYLATE (Suspected)				**Reason:**	Paralysis agitans	
Tablet	3.0	Dose Unspecified		Daily		Oral
Batch:	**Started:**	22/05/1992			**Stopped:**	29/05/1992
MELLERIL (Suspected)				**Reason:**	Depression	
Tablet	150.0	Milligram		Daily		Oral
Batch:	**Started:**	01/01/1991			**Stopped:**	30/05/1992
TRYPTANOL (Suspected)				**Reason:**	Depression	
Tablet	75.0	Milligram		Daily		Oral
Batch:	**Started:**	01/01/1991			**Stopped:**	30/03/1992

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 83325 **Seq:** 1 **Gender:** M
Reported: 19/03/1993 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 19/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cold sweat Dyspnoea Hyperhidrosis			Adrenaline 300mcg sc, hydrocortisone iv.

Medicine Details:

IOPAMIRO (Suspected)			**Reason:**	
Injection	20.0	Millilitre	1 time	Intravenous
Batch:	**Started:** 19/03/1993			**Stopped:**
ATROPINE (Suspected)			**Reason:**	
Injection	600.0	Microgram	1 time	Intravenous
Batch:	**Started:** 19/03/1993			**Stopped:**
MIDAZOLAM (Suspected)			**Reason:**	
Injection	1.0	Milligram	1 time	Intravenous
Batch:	**Started:** 19/03/1993			**Stopped:**
CEPHALOTHIN SODIUM (Suspected)			**Reason:**	
Injection	2.0	Gram	Daily	Intravenous
Batch:	**Started:** 19/03/1993			**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83325 **Seq:** 1 **Gender:** M
Reported: 19/03/1993 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 19/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

METRONIDAZOLE (Suspected)			**Reason:**	
Injection	500.0	Milligram	Daily	Intravenous
Batch:	**Started:** 19/03/1993			**Stopped:**
PETHIDINE HYDROCHLORIDE (Suspected)			**Reason:**	
Injection	150.0	Milligram	Daily	Intramuscular
Batch:	**Started:** 18/03/1993			**Stopped:**
VALIUM (Suspected)			**Reason:**	
Tablet	10.0	Milligram	Daily	Oral
Batch:	**Started:** 19/03/1993			**Stopped:**
TEMAZEPAM (Suspected)			**Reason:**	
Oral application	20.0	Milligram	Daily	Oral
Batch:	**Started:** 18/03/1993			**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 83325 **Seq:** 1 **Gender:** M
Reported: 19/03/1993 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 19/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Oral application 75.0 Milligram Daily Oral

Batch: **Started:** 18/03/1993 **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 84010 **Seq:** 1
Reported: 30/03/1993
Hospitalisation: Admitted to hospital
Onset Date: 30/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bronchospasm Hypotension			Ventolin neb.

Medicine Details:

FLUVAX (Suspected) **Reason:** Prophylactic inoculation and vaccination
1.0 Dose Unspecified 1 time
Batch: **Started:** **Stopped:**

MODURETIC (Suspected) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** CONTIN

TRANSIDERM-NITRO (Suspected) **Reason:** Pain in chest
Cream 25.0 Milligram Daily Transdermal
Batch: **Started:** 30/03/1993 **Stopped:** 30/03/1993

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 29/03/1993 **Stopped:** 30/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 84010 **Seq:** 1
Reported: 30/03/1993
Hospitalisation: Admitted to hospital
Onset Date: 30/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

METFORMIN HYDROCHLORIDE (Suspected)				**Reason:**	Diabetes mellitus		
Tablet	500.0	Milligram	Daily		Oral		
Batch:		**Started:**	L TERM		**Stopped:**	CONTIN	

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 84104 **Seq:** 1 **Gender:** F
Reported: 15/04/1993 **Weight:**
Hospitalisation: **Age:** 43Y
Onset Date: 31/12/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia		Urge to move legs continously.	

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Tablet 125.0 Milligram Daily Oral

Batch: **Started:** 01/10/1992 **Stopped:** 28/04/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Tryptanol dose was reduced with improvement. reaction later recurred so drug ceased. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 85435 **Seq:** 1 **Gender:** F
Reported: 22/06/1993 **Weight:** 50.00
Hospitalisation: **Age:** 78Y
Onset Date: 31/03/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

FRUSEMIDE (Suspected) **Reason:**
Oral application 40.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 25/03/1993

INDOCID (Suspected) **Reason:** Gout
Capsule 50.0 Milligram Daily Oral
Batch: **Started:** 13/11/1992 **Stopped:** 12/03/1993

ADALAT (Suspected) **Reason:** Essential benign hypertension
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 01/01/1991 **Stopped:** 25/03/1993

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Oral application 75.0 Milligram Daily Oral
Batch: **Started:** 01/09/1992 **Stopped:** 25/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	8-40	15/05/1991	8	
	ALT = SGPT	8-40	27/01/1993	27	
	ALT = SGPT	8-40	15/03/1993	85	
	ALT = SGPT	8-40	25/03/1993	81	
	ALT = SGPT	8-40	06/04/1993	31	
	ALT = SGPT	8-40	31/03/1993	38	
	Albumin	36-47	15/05/1991	36	

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 85435 **Seq:** 1
Reported: 22/06/1993
Hospitalisation:
Onset Date: 31/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 50.00
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

INDOCID (Suspected) **Reason:** Other disturbance of sensation

Suppository 200.0 Milligram Daily Rectal

Batch: **Started:** 12/03/1993 **Stopped:** 24/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Albumin	36-47	15/03/1993	38	
	Albumin	36-47	25/03/1993	42	
	Albumin	36-47	06/04/1993	38	
	Albumin	36-47	31/03/1993	41	
	Albumin	36-47	27/01/1993	40	
	Bilirubin	<20	15/05/1991	12	
	Bilirubin	<20	27/01/1993	<20	

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 85435 **Seq:** 1
Reported: 22/06/1993
Hospitalisation:
Onset Date: 31/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 50.00
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

FRUSEMIDE (Suspected)			**Reason:**		
Oral application	40.0	Milligram	Daily		Oral
Batch:	**Started:**		L TERM	**Stopped:**	25/03/1993
INDOCID (Suspected)			**Reason:**	Gout	
Capsule	50.0	Milligram	Daily		Oral
Batch:	**Started:**	13/11/1992		**Stopped:**	12/03/1993
ADALAT (Suspected)			**Reason:**	Essential benign hypertension	
Tablet	40.0	Milligram	Daily		Oral
Batch:	**Started:**	01/01/1991		**Stopped:**	25/03/1993
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Depression	
Oral application	75.0	Milligram	Daily		Oral
Batch:	**Started:**	01/09/1992		**Stopped:**	25/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	<20	15/03/1993	<20	
	Bilirubin	<20	25/03/1993	<20	
	Bilirubin	<20	31/03/1993	<20	
	Bilirubin	<20	06/04/1993	<20	
	GGT = SGGT =	8-40	15/05/1991	29	
	GGT = SGGT =	8-40	27/01/1993	65	
	GGT = SGGT =	8-40	25/03/1993	483	

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 85435 **Seq:** 1
Reported: 22/06/1993
Hospitalisation:
Onset Date: 31/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 50.00
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

INDOCID (Suspected) **Reason:** Other disturbance of sensation

Suppository 200.0 Milligram Daily Rectal

Batch: **Started:** 12/03/1993 **Stopped:** 24/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT =	8-40	06/04/1993	222	
	GGT = SGGT =	8-40	31/03/1993	390	
	GGT = SGGT =	8-40	15/03/1993	592	
	SAP = ALP	35-120	15/05/1991	65	
	SAP = ALP	35-120	27/01/1993	103	
	SAP = ALP	35-120	15/03/1993	369	
	SAP = ALP	35-120	25/03/1993	242	

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 85435 **Seq:** 1 **Gender:** F
Reported: 22/06/1993 **Weight:** 50.00
Hospitalisation: **Age:** 78Y
Onset Date: 31/03/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FRUSEMIDE (Suspected)				**Reason:**		
Oral application	40.0	Milligram		Daily		Oral
Batch:	**Started:**		L TERM		**Stopped:**	25/03/1993
INDOCID (Suspected)				**Reason:**	Gout	
Capsule	50.0	Milligram		Daily		Oral
Batch:	**Started:**	13/11/1992			**Stopped:**	12/03/1993
ADALAT (Suspected)				**Reason:**	Essential benign hypertension	
Tablet	40.0	Milligram		Daily		Oral
Batch:	**Started:**	01/01/1991			**Stopped:**	25/03/1993
AMITRIPTYLINE HYDROCHLORIDE (Suspected)				**Reason:**	Depression	
Oral application	75.0	Milligram		Daily		Oral
Batch:	**Started:**	01/09/1992			**Stopped:**	25/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP	35-120	31/03/1993	135	
	SAP = ALP	35-120	06/04/1993	100	

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 85435 **Seq:** 1
Reported: 22/06/1993
Hospitalisation:
Onset Date: 31/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 50.00
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

INDOCID (Suspected) **Reason:** Other disturbance of sensation

Suppository 200.0 Milligram Daily Rectal

Batch: **Started:** 12/03/1993 **Stopped:** 24/03/1993

Laboratory Investigations:

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 85552 **Seq:** 1 **Gender:** F
Reported: 25/06/1993 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 31/12/1990 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Visual disturbance			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

0.0

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergies= elastoplast. - the date of onset is not accurate but indicates that onset occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 85830 **Seq:** 1 **Gender:** M
Reported: 02/07/1993 **Weight:** 92.00
Hospitalisation: **Age:** 62Y
Onset Date: 27/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lichenoid keratosis Photosensitivity reaction			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 10/05/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 85854 **Seq:** 1 **Gender:** U
Reported: 05/07/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 30/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope			

Medicine Details:

IMIGRAN (Suspected) **Reason:** Migraine
Tablet 1.0 Dose Unspecified 1 time Oral
Batch: **Started:** 29/06/1993 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 17/06/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 85928 **Seq:** 1 **Gender:** U
Reported: 07/07/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 21/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Nightmare			

Medicine Details:

TEGRETOL (Other drug) **Reason:**
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

ALLOPURINOL (Other drug) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

ASPALGIN (Other drug) **Reason:**
Tablet 8.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

PETHIDINE HYDROCHLORIDE (Suspected) **Reason:**
Injection 50.0 Milligram 2 times Intramuscular
Batch: **Started:** 20/05/1993 **Stopped:** 21/05/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 85928 **Seq:** 1 **Gender:** U
Reported: 07/07/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 21/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CHLORPROMAZINE HYDROCHLORIDE (Suspected)			**Reason:**		
Injection	50.0	Milligram	Total		Intramuscular
Batch:	**Started:**	20/05/1993		**Stopped:**	21/05/1993
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
	100.0	Milligram	Total		
Batch:	**Started:**	20/05/1993		**Stopped:**	21/05/1993

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 86657 **Seq:** 1 **Gender:** M
Reported: 04/08/1993 **Weight:** 82.00
Hospitalisation: **Age:** 33Y
Onset Date: 28/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Psychotic disorder			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 15/04/1993 **Stopped:** 28/07/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 87417 **Seq:** 1 **Gender:** F
Reported: 31/08/1993 **Weight:** 80.00
Hospitalisation: **Age:** 37Y
Onset Date: 17/08/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

VOLTAREN (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	1 - 24 U/L	17/08/1993	133	
	AST = SGOT	1 - 24 U/L	23/08/1993	92	
	Bilirubin	1 - 24 UMOL/L	17/08/1993	4	
	GGT = SGGT =	1 - 30 U/L	17/08/1993	23	
	GGT = SGGT =	1 - 30 U/L	23/08/1993	13	
	SAP = ALP	30 115 U/L	17/08/1993	90	
	SAP = ALP	30 115 U/L	23/08/1993	77	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87630 **Seq:** 1
Reported: 13/09/1993
Hospitalisation:
Onset Date: 28/07/1993
Outcome: Death, maybe drug

Gender: M
Weight: 56.00
Age: 39Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia Anaemia Confusional state Headache Sepsis Thrombocytopenia Vomiting	Severe	Temp 41 degrees.	5u rbc's, 6u haemacel and 5u platelets.

Medicine Details:

BACTRIM (Suspected) **Reason:** Otr&nos infec&parasit diseases
Oral application 6.0 Dose Unspecified Weekly Oral
Batch: **Started:** 21/07/1993 **Stopped:** 02/08/1993

ZALCITABINE (Suspected) **Reason:** Hiv infection
Tablet 4.0 Dose Unspecified Daily Oral
Batch: **Started:** 20/07/1993 **Stopped:** 28/07/1993

FLUCONAZOLE (Suspected) **Reason:** Moniliasis
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 23/03/1993 **Stopped:** 04/08/1993

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 20/07/1993 **Stopped:** 06/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haemoglobin		28/07/1993	7.9	
	Haemoglobin		02/08/1993	7.8	
	Platelets		02/08/1993	42	
	Platelets		04/08/1993	30	
	Platelets		06/08/1993	15	
	Platelets		07/08/1993	22	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87652 **Seq:** 1
Reported: 13/09/1993
Hospitalisation: Admitted to hospital
Onset Date: 12/05/1993
Outcome: Death, maybe drug

Gender: F
Weight: 64.00
Age: 61Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mouth ulceration		Mucosal ulceration.	
Rash		Rash over arms, legs, trunk and face.	
Coagulopathy			
Eosinophilia			
Hepatic failure			
Jaundice			
Malaise			
Renal failure acute			Prednisolone, urs0deoxycholic acid and liver transplant.

Medicine Details:

AUGMENTIN FORTE (Suspected) **Reason:** Acute bronchitis&bronchiolitis
Oral application 5.0 Dose Unspecified Total Oral
Batch: **Started:** 11/05/1993 **Stopped:** 12/05/1993

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 01/01/1983 **Stopped:**

METOPROLOL TARTRATE (Suspected) **Reason:** Essential benign hypertension
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	10-50 U/L	02/07/1993	66	
	AST = SGOT	10-50 U/L	09/07/1993	52	
	AST = SGOT	10-50 U/L	29/07/1993	121	
	AST = SGOT	10-50 U/L	03/08/1993	148	
	Albumin	35-45 G/L	02/07/1993	17	
	Albumin	35-45 G/L	09/07/1993	14	
	Albumin	35-45 G/L	29/07/1993	27	

Additional Information:

Patient died on 07/10/93. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87652 **Seq:** 1
Reported: 13/09/1993
Hospitalisation: Admitted to hospital
Onset Date: 12/05/1993
Outcome: Death, maybe drug

Gender: F
Weight: 64.00
Age: 61Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AUGMENTIN FORTE (Suspected)				**Reason:**	Acute bronchitis&bronchiolitis
Oral application	5.0	Dose Unspecified	Total		Oral
Batch:	**Started:**	11/05/1993		**Stopped:**	12/05/1993
TRYPTANOL (Suspected)				**Reason:**	Depression
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**	01/01/1983		**Stopped:**	
METOPROLOL TARTRATE (Suspected)				**Reason:**	Essential benign hypertension
Oral application	100.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Albumin	35-45 G/L	03/08/1993	27	
	Bilirubin	<17 UMOL/L	02/07/1993	530	
	Bilirubin	<17 UMOL/L	09/07/1993	333	
	Bilirubin	<17 UMOL/L	03/08/1993	565	
	Bilirubin	<17 UMOL/L	29/07/1993	519	

Additional Information:

Patient died on 07/10/93. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87652 **Seq:** 1
Reported: 13/09/1993
Hospitalisation: Admitted to hospital
Onset Date: 12/05/1993
Outcome: Death, maybe drug

Gender: F
Weight: 64.00
Age: 61Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AUGMENTIN FORTE (Suspected)				**Reason:**	Acute bronchitis&bronchiolitis
Oral application	5.0	Dose Unspecified	Total		Oral
Batch:	**Started:**	11/05/1993		**Stopped:**	12/05/1993
TRYPTANOL (Suspected)				**Reason:**	Depression
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**	01/01/1983		**Stopped:**	
METOPROLOL TARTRATE (Suspected)				**Reason:**	Essential benign hypertension
Oral application	100.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Liver biopsy				2/6/93 - the appearances are those of a cholestatic jaundice and would be compatible with a drug induced optiology. 26/7/93 - marked cholestasis and severe bile duct injury with mild hepatitis
	SAP = ALP	30-120 U/L	02/07/1993	337	
	SAP = ALP	30-120 U/L	09/07/1993	274	
	SAP = ALP	30-120 U/L	29/07/1993	694	
	SAP = ALP	30-120 U/L	03/08/1993	956	

Additional Information:

Patient died on 07/10/93. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 87684 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:**
Hospitalisation: **Age:** 46
Onset Date: 23/04/1993 **DOB:** 16/10/1946
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Convulsion			

Medicine Details:

DOLOXENE CO (Other drug) **Reason:**
Capsule 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:** CONTIN

OXYCODONE HYDROCHLORIDE (Other drug) **Reason:**
10.0 Milligram As necessary
Batch: **Started:** **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level				Tricyclic level: 600ug/ml, which was significantly elevated.

Additional Information:

Report Detail

Case Number: 87743 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:** 45.00
Hospitalisation: **Age:** 68Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agranulocytosis			
Anorexia			
Diarrhoea			
Drug ineffective			
Dyspnoea			
Fatigue			
Hyperhidrosis			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
0.0
Batch: **Started:** 01/01/1973 **Stopped:** 28/06/1993

MYLANTA (Suspected) **Reason:**
0.0
Batch: **Started:** 01/08/1993 **Stopped:** 16/08/1993

DISPRIN (Suspected) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:** 16/08/1993

QUININE (Suspected) **Reason:** Pain in limb
Oral application 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** 01/01/1991 **Stopped:** 16/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bone marrow				20/8/93 - bone marrow - early myeloid series.
	Neutrophils	2.00 - 7.50	16/08/1993	-	
	Neutrophils	2.00 - 7.50	22/08/1993	0.15	
	Neutrophils	2.00 - 7.50	24/08/1993	-	
	Neutrophils	2.00 - 7.50	27/08/1993	0.69	
	Neutrophils	2.00 - 7.50	26/08/1993	0.16	
	Neutrophils	2.00 - 7.50	25/08/1993	0.03	

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 87743 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:** 45.00
Hospitalisation: **Age:** 68Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TOLVON (Suspected)			**Reason:**	Depression	
Tablet	30.0	Milligram	Daily		Oral
Batch:	**Started:**	01/06/1993		**Stopped:**	17/08/1993
CORTISONE (Suspected)			**Reason:**		
Ointment	1.0	Dose Unspecified	Daily		Topical
Batch:	**Started:**	01/01/1983		**Stopped:**	16/08/1993
SULPHUR (Suspected)			**Reason:**		
Ointment	1.0	Dose Unspecified	Daily		Topical
Batch:	**Started:**	01/01/1983		**Stopped:**	16/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Neutrophils	2.00 - 7.50	23/08/1993	0.18	
	White blood cells	4.0 - 11.0	16/08/1993	1.4	
	White blood cells	4.0 - 11.0	22/08/1993	1.5	
	White blood cells	4.0 - 11.0	23/08/1993	1.3	
	White blood cells	4.0 - 11.0	24/08/1993	2.2	
	White blood cells	4.0 - 11.0	25/08/1993	1.9	
	White blood cells	4.0 - 11.0	26/08/1993	2.1	

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87743 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:** 45.00
Hospitalisation: **Age:** 68Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:** 01/01/1973		**Stopped:**	28/06/1993
MYLANTA (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:** 01/08/1993		**Stopped:**	16/08/1993
DISPRIN (Suspected)			**Reason:**	
	0.0	As necessary		
Batch:	**Started:**		**Stopped:**	16/08/1993
QUININE (Suspected)			**Reason:**	Pain in limb
Oral application	1.0 Dose Unspecified	As necessary		Oral
Batch:	**Started:** 01/01/1991		**Stopped:**	16/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	White blood cells	4.0 - 11.0	27/08/1993	2.1	

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87743 **Seq:** 1
Reported: 13/09/1993
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 45.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TOLVON (Suspected)				**Reason:**	Depression	
Tablet	30.0	Milligram	Daily		Oral	
Batch:	**Started:**	01/06/1993		**Stopped:**	17/08/1993	
CORTISONE (Suspected)				**Reason:**		
Ointment	1.0	Dose Unspecified	Daily		Topical	
Batch:	**Started:**	01/01/1983		**Stopped:**	16/08/1993	
SULPHUR (Suspected)				**Reason:**		
Ointment	1.0	Dose Unspecified	Daily		Topical	
Batch:	**Started:**	01/01/1983		**Stopped:**	16/08/1993	

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 87990 **Seq:** 1 **Gender:** F
Reported: 29/09/1993 **Weight:**
Hospitalisation: **Age:** 73Y
Onset Date: 03/09/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

TEMAZEPAM (Other drug) **Reason:**
Capsule 10.0 Milligram Daily Oral
Batch: **Started:** 09/08/1993 **Stopped:**

AMILORIDE HYDROCHLORIDE (Other drug) **Reason:** Congestive heart failure
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 03/08/1993 **Stopped:**

FRUSEMIDE (Other drug) **Reason:** Congestive heart failure
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 03/09/1993 **Stopped:**

NAPROXEN (Suspected) **Reason:** Rheum arthrit&allied condition
Tablet 500.0 Milligram Daily Oral
Batch: **Started:** 04/08/1993 **Stopped:** 03/09/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	6 - 42 U/L	03/09/1993	158	
	AST = SGOT	6 - 42 U/L	06/09/1993	16	
	Bilirubin	3 - 20 UMOL/L	03/09/1993	27	
	Bilirubin	3 - 20 UMOL/L	06/09/1993	5	
	Creatine	30 - 180 U/L	03/09/1993	72	
	Creatine	30 - 180 U/L	06/09/1993	37	
	GGT = SGGT =	5 - 40 U/L	03/09/1993	172	

Additional Information:

Subsequently this patient has been found to be hepatitis c positive.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 87990 **Seq:** 1 **Gender:** F
Reported: 29/09/1993 **Weight:**
Hospitalisation: **Age:** 73Y
Onset Date: 03/09/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 100.0 Milligram Daily Oral

Batch: **Started:** 03/08/1993 **Stopped:** 03/09/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP	35 - 135 U/L	03/09/1993	162	
	SAP = ALP	35 - 135 U/L	06/09/1993	104	

Additional Information:

Subsequently this patient has been found to be hepatitis c positive.

Report Detail

Case Number: 88964 **Seq:** 1
Reported: 01/11/1993
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 50Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura Thrombocytopenia			 Prednisolone 100mg daily

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

THYROXINE SODIUM (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

TRIFLUOPERAZINE HYDROCHLORIDE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

TRANYLCYPROMINE SULPHATE (Other drug) **Reason:**
80.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				Borderline - low serum folate 10/93.
	Bone marrow				Bone marrow changes consistent with myeloid dysplasia
	Platelets		20/10/1993	11000	
	Platelets		15/11/1993	315000	

Additional Information:

Report Detail

Case Number: 88964 **Seq:** 1
Reported: 01/11/1993
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 50Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:** Depression		
Tablet	150.0	Milligram	Daily		Oral
Batch:	**Started:** 06/08/1993			**Stopped:**	20/10/1993
CARBAMAZEPINE (Suspected)			**Reason:**		
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	20/10/1993

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 88970 **Seq:** 1 **Gender:** F
Reported: 01/11/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Depression			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Debility&undue fatigue
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 88971 **Seq:** 1 **Gender:** U
Reported: 01/11/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Dysgeusia		Bitter taste.	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 88992 **Seq:** 1
Reported: 02/11/1993
Hospitalisation:
Onset Date: 21/10/1993
Outcome: Not yet recovered

Gender: M
Weight: 55.00
Age: 40Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coombs positive haemolytic anae			Oral prednisone.

Medicine Details:

DAPSONE (Suspected) **Reason:** Hiv infection
2.0 Dose Unspecified Weekly
Batch: **Started:** **Stopped:** 06M

TEGRETOL (Suspected) **Reason:** Herpes zoster
400.0 Milligram Daily
Batch: **Started:** **Stopped:** 03W

TRYPTANOL (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

ACICLOVIR (Suspected) **Reason:** Herpes zoster
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 06M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haemoglobin				21/10/93 hb 5.1, treated with oral prednisone hb 8.0.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 89179 **Seq:** 1 **Gender:** F
Reported: 10/11/1993 **Weight:**
Hospitalisation: **Age:** 57Y
Onset Date: 02/09/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			

Medicine Details:

MACRODANTIN (Other drug)			**Reason:** Othr diseases of urinary tract	
Capsule	100.0 Milligram	Daily	Oral	
Batch:	**Started:** 20/08/1993		**Stopped:**	CONTIN
TRYPTANOL (Suspected)			**Reason:** Headache	
Tablet	25.0 Milligram	Daily	Oral	
Batch:	**Started:** 28/08/1993		**Stopped:** 03/09/1993	
INDERAL (Suspected)			**Reason:** Headache	
Tablet	80.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/09/1993		**Stopped:** 04/09/1993	
PREDNISOLONE (Suspected)			**Reason:**	
Oral application	125.0 Milligram	Daily	Oral	
Batch:	**Started:** 30/08/1993		**Stopped:** 31/08/1993	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 89179 **Seq:** 1 **Gender:** F
Reported: 10/11/1993 **Weight:**
Hospitalisation: **Age:** 57Y
Onset Date: 02/09/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

NORMISON (Suspected)				**Reason:**	Specific disorders of sleep	
Capsule	10.0	Milligram	Daily		Oral	
Batch:	**Started:**	25/08/1993			**Stopped:**	02/09/1993
MAXOLON (Suspected)				**Reason:**	Nausea and vomiting	
Oral application	30.0	Milligram	Daily		Oral	
Batch:	**Started:**	26/08/1993			**Stopped:**	02/09/1993
INDOCID (Suspected)				**Reason:**	Pain	
Suppository	2.0	Dose Unspecified	Daily		Rectal	
Batch:	**Started:**	01/09/1993			**Stopped:**	04/09/1993

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 89616 **Seq:** 1 **Gender:** F
Reported: 02/12/1993 **Weight:** 53.00
Hospitalisation: **Age:** 67Y
Onset Date: 27/08/1993 **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Colitis ulcerative Pancytopenia			Folinic acid, ceptazidime

Medicine Details:

METHOTREXATE (Suspected) **Reason:** Other rheumatoid arthritis
Oral application 12.5 Milligram Weekly Oral
Batch: **Started:** **Stopped:** 27/08/1993

PREDNISOLONE (Suspected) **Reason:** Other rheumatoid arthritis
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 27/08/1993

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 30/08/1993

AUGMENTIN FORTE (Suspected) **Reason:** Othr diseases of urinary tract
Oral Liquid 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 20/08/1993 **Stopped:** 27/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haemoglobin	110-160 G/L	26/07/1993	12.3	
	Haemoglobin	110-160 G/L	28/08/1993	97L	
	Haemoglobin	110-160 G/L	29/08/1993	63L	
	Haemoglobin	110-160 G/L	31/08/1993	152	
	Haemoglobin	110-160 G/L	02/09/1993	110	
	Haemoglobin	110-160 G/L	05/09/1993	105L	
	Platelets	150-450 X10/9/L	26/07/1993	543	

Additional Information:

Report Detail

Case Number: 89616 **Seq:** 1
Reported: 02/12/1993
Hospitalisation:
Onset Date: 27/08/1993
Outcome: Death, maybe drug

Gender: F
Weight: 53.00
Age: 67Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

SEPTRIN FORTE (Suspected) **Reason:** Othr diseases of urinary tract

Tablet 2.0 Dose Unspecified Daily Oral

Batch: **Started:** 17/08/1993 **Stopped:** 20/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Platelets	150-450 X10/9/L	28/08/1993	23L	
	Platelets	150-450 X10/9/L	31/08/1993	33L	
	Platelets	150-450 X10/9/L	05/09/1993	65L	
	Platelets	150-450 X10/9/L	02/09/1993	43L	
	Platelets	150-450 X10/9/L	29/08/1993	29L	
	White blood cells	4.00-11.00	26/07/1993	8.6	
	White blood cells	4.00-11.00	28/08/1993	0.40L	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 89616 **Seq:** 1
Reported: 02/12/1993
Hospitalisation:
Onset Date: 27/08/1993
Outcome: Death, maybe drug

Gender: F
Weight: 53.00
Age: 67Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

METHOTREXATE (Suspected)				**Reason:**	Other rheumatoid arthritis	
Oral application	12.5	Milligram		Weekly		Oral
Batch:	**Started:**				**Stopped:**	27/08/1993
PREDNISOLONE (Suspected)				**Reason:**	Other rheumatoid arthritis	
Oral application	10.0	Milligram		Daily		Oral
Batch:	**Started:**				**Stopped:**	27/08/1993
AMITRIPTYLINE HYDROCHLORIDE (Suspected)				**Reason:**	Depression	
Oral application	100.0	Milligram		Daily		Oral
Batch:	**Started:**				**Stopped:**	30/08/1993
AUGMENTIN FORTE (Suspected)				**Reason:**	Othr diseases of urinary tract	
Oral Liquid	3.0	Dose Unspecified		Daily		Oral
Batch:	**Started:**	20/08/1993			**Stopped:**	27/08/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	White blood cells	4.00-11.00	29/08/1993	0.40L	
	White blood cells	4.00-11.00	31/08/1993	4.72	
	White blood cells	4.00-11.00	02/09/1993	8.33	
	White blood cells	4.00-11.00	05/09/1993	10.38	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 89616 **Seq:** 1
Reported: 02/12/1993
Hospitalisation:
Onset Date: 27/08/1993
Outcome: Death, maybe drug

Gender: F
Weight: 53.00
Age: 67Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

SEPTRIN FORTE (Suspected) **Reason:** Othr diseases of urinary tract

Tablet 2.0 Dose Unspecified Daily Oral

Batch: **Started:** 17/08/1993 **Stopped:** 20/08/1993

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 90103 **Seq:** 1 **Gender:** F
Reported: 24/12/1993 **Weight:**
Hospitalisation: **Age:** 56Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 28/12/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Report Detail

Case Number: 90656 **Seq:** 1 **Gender:** F
Reported: 25/01/1994 **Weight:** 45.00
Hospitalisation: **Age:** 93Y
Onset Date: 07/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Inappropriate antidiuretic hormon			Fluid restriction, demeclocycline 300mg.

Medicine Details:

MORPHINE NOS (Other drug) **Reason:** Pain
Injection 2.5 Milligram As necessary Intravenous
Batch: **Started:** 06/01/1994 **Stopped:**

MAXOLON (Other drug) **Reason:** Nausea and vomiting
Injection 10.0 Milligram As necessary Intravenous
Batch: **Started:** 06/01/1994 **Stopped:**

CARTIA (Other drug) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SEPTRIN (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 07/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum osmolality		07/01/1994	237	
	Sodium		07/01/1994	117	
	Sodium		08/01/1994	119	
	Sodium		09/01/1994	115	
	Sodium		10/01/1994	117	
	Sodium		12/01/1994	130	
	Sodium		13/01/1994	131	

Additional Information:

Report Detail

Case Number: 90656 **Seq:** 1 **Gender:** F
Reported: 25/01/1994 **Weight:** 45.00
Hospitalisation: **Age:** 93Y
Onset Date: 07/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

SEREPAX (Suspected)			**Reason:**		
Tablet	15.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	07/01/1994
TRYPTANOL (Suspected)			**Reason:**	Depression	
Tablet	25.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	07/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Urine osmolality	MMOL/KG.	07/01/1994	519	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 91223 **Seq:** 1
Reported: 23/02/1994
Hospitalisation:
Onset Date: 19/02/1994
Outcome: Not yet recovered

Gender: M
Weight: 75.00
Age: A7
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction	Severe	Sunburn on legs and face.	

Medicine Details:

STELAZINE (Suspected) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 01/11/1992 **Stopped:** CONTIN

ENDEP (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 01/11/1992 **Stopped:** CONTIN

FELDENE (Suspected) **Reason:** Unspecified arthritis
20.0 Milligram Daily
Batch: **Started:** 01/11/1992 **Stopped:**

KALURIL (Suspected) **Reason:**
5.0 Milligram Daily
Batch: **Started:** 01/11/1992 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 91223 **Seq:** 1
Reported: 23/02/1994
Hospitalisation:
Onset Date: 19/02/1994
Outcome: Not yet recovered

Gender: M
Weight: 75.00
Age: A7
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ECOTRIN (Suspected)		**Reason:**	Unspecified arthritis	
	1.0 Dose Unspecified Daily			
Batch:	**Started:** 01/11/1992		**Stopped:**	CONTIN
LASIX (Suspected)		**Reason:**		
	1.0 Dose Unspecified Daily			
Batch:	**Started:** 01/11/1992		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 91465 **Seq:** 1 **Gender:** M
Reported: 09/03/1994 **Weight:** 90.00
Hospitalisation: **Age:** 29
Onset Date: 08/03/1994 **DOB:** 09/07/1964
Outcome: Death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myocarditis Sudden death			

Medicine Details:

BENZTROPINE MESYLATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CLOZARIL (Suspected) **Reason:**
Oral application 275.0 Milligram Daily Oral
Batch: **Started:** 21/02/1994 **Stopped:** 08/03/1994

LITHIUM CARBONATE (Suspected) **Reason:** Unspecifid affective psychosis
1.2 Gram Daily
Batch: **Started:** 17/02/1994 **Stopped:** 08/03/1994

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** 28/02/1994 **Stopped:** 08/03/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Histology				The morphologic features within the heart are typical of hypersensitivity myocarditis.

Additional Information:

Report Detail

Case Number: 91754 **Seq:** 1
Reported: 23/03/1994
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight:
Age: 63Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension Dysarthria		Bp was 280/140	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

PRAZOSIN HYDROCHLORIDE (Interaction) **Reason:** Essential benign hypertension
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

METOPROLOL TARTRATE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

CHLORPROMAZINE HYDROCHLORIDE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 91755 **Seq:** 1 **Gender:** F
Reported: 23/03/1994 **Weight:** 62.00
Hospitalisation: **Age:** 54Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Drug withdrawal syndrome Muscle spasms Tachycardia	Severe		Amitriptyline increased to 50mg daily

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp vertebrogen pain synd

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 09/03/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 91756 **Seq:** 1 **Gender:** F
Reported: 23/03/1994 **Weight:**
Hospitalisation: **Age:** 26Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast enlargement Breast pain			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 92241 **Seq:** 1 **Gender:** M
Reported: 15/04/1994 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 92419 **Seq:** 1 **Gender:** F
Reported: 26/04/1994 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 22/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuroleptic malignant syndrome			

Medicine Details:

DIAZEPAM (Other drug) **Reason:** Unspecified schizophrenia
Tablet 5.0 Milligram As necessary Oral
Batch: **Started:** 07/07/1993 **Stopped:**

PARACETAMOL (Other drug) **Reason:** Pyrexia of unknown origin
Oral application 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 07/07/1993 **Stopped:**

SEPTRIN FORTE (Other drug) **Reason:** Othr diseases of urinary tract
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 07/07/1993 **Stopped:**

CHLORPROMAZINE HYDROCHLORIDE (Suspected) **Reason:** Unspecified schizophrenia
Oral application 100.0 Milligram As necessary Oral
Batch: **Started:** 07/07/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry	< 0.06 MG/L	22/07/1993	107.00	Myoglobin
	Biochemistry	< 0.06 MG/L	22/07/1993	34.80	Myoglobin
	Biochemistry	< 0.38 MG/L	22/07/1993	3180	Urine myoglobin
	Creatine kinase MB	< 10U/L	22/07/1993	517	
	Creatine phosphokinase	25 - 200 U/L	22/07/1993	23690	
	Creatine phosphokinase	25 - 200 U/L	22/07/1993	139178	
	Creatine phosphokinase	25 - 200 U/L	23/07/1993	98778	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 92419 **Seq:** 1 **Gender:** F
Reported: 26/04/1994 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 22/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

MODECATE (Suspected)				**Reason:**	Unspecified schizophrenia
Injection	50.0	Milligram	Fortnightly		Intramuscular
Batch:	**Started:**			**Stopped:**	14/07/1993
TRIFLUOPERAZINE HYDROCHLORIDE (Suspected)				**Reason:**	Unspecified schizophrenia
Oral application	30.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)				**Reason:**	
Tablet	150.0	Milligram	Daily		Oral
Batch:	**Started:**	21/07/1993		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine [illegible]	25 - 200 U/L	24/07/1993	42708	
	Creatine [illegible]	25 - 200 U/L	25/07/1993	16831	
	Creatine [illegible]	25 - 200 U/L	27/07/1993	3731	

Additional Information:

Report Detail

Case Number: 93240 **Seq:** 1 **Gender:** F
Reported: 30/05/1994 **Weight:**
Hospitalisation: **Age:** 70Y
Onset Date: 24/05/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erythema multiforme			

Medicine Details:

ISOPTIN SR (Suspected) **Reason:** Essential benign hypertension
120.0 Milligram Daily
Batch: **Started:** 10/05/1994 **Stopped:** 25/05/1994

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
10.0 Milligram Daily
Batch: **Started:** 01/03/1993 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 93243 **Seq:** 1 **Gender:** F
Reported: 30/05/1994 **Weight:**
Hospitalisation: **Age:** 44Y
Onset Date: 13/04/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Pruritus		Head to toe rash	

Medicine Details:

MICROGYNON (Other drug) **Reason:** Contraception
0.0
Batch: **Started:** L TERM **Stopped:** CONTIN

CALCIUM NOS (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:** CONTIN

VENTOLIN (Other drug) **Reason:** Asthma
Inhalation 0.0 As necessary Inhalation
Batch: **Started:** **Stopped:**

ALDACTONE (Other drug) **Reason:** Menopausal symptoms
100.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous reactions to doxycycline - nausea, tetracycline - photosensitivity. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 93243 **Seq:** 1 **Gender:** F
Reported: 30/05/1994 **Weight:**
Hospitalisation: **Age:** 44Y
Onset Date: 13/04/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TEGRETOL (Suspected)		**Reason:** Pain	
	200.0 Milligram	Daily	
Batch:	**Started:** 03/04/1994		**Stopped:** 13/04/1994
TRYPTANOL (Suspected)		**Reason:** Pain	
	10.0 Milligram	Daily	
Batch:	**Started:** 03/04/1994		**Stopped:** 13/04/1994

Laboratory Investigations:

Additional Information:

Previous reactions to doxycycline - nausea, tetracycline - photosensitivity. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 93763 **Seq:** 1 **Gender:** F
Reported: 20/06/1994 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: 03/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Agitation Confusional state Delusion Nightmare		Auditory hallucinations	

Medicine Details:

PREDNISONE (Other drug) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

RANITIDINE (Other drug) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

THYROXINE SODIUM (Other drug) **Reason:** Myxedema
Tablet 150.0 Microgram Daily Oral
Batch: **Started:** **Stopped:**

TAMOXIFEN CITRATE (Other drug) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93763 **Seq:** 1 **Gender:** F
Reported: 20/06/1994 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: 03/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ENDONE (Suspected)			**Reason:**	Otr&unsp forms neuralg&neurit	
Tablet	30.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	04/06/1994
ENDEP (Suspected)			**Reason:**	Otr&unsp forms neuralg&neurit	
Tablet	100.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	04/06/1994
CARBAMAZEPINE (Suspected)			**Reason:**	Otr&unsp forms neuralg&neurit	
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	04/06/1994

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 93939 **Seq:** 1 **Gender:** F
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 06/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cholecystitis Jaundice Nausea Pruritus			

Medicine Details:

CALTRATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 04/06/1994

HEPARIN CALCIUM (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 04/06/1994

FLUCLOXACILLIN SODIUM (Suspected) **Reason:** Postoperative wound infection
Injection 4.0 Gram Daily Intravenous
Batch: **Started:** 14/05/1994 **Stopped:** 23/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	5-35 U/L	06/06/1994	108	
	AST = SGOT	5-35 U/L	08/06/1994	138	
	AST = SGOT	5-35 U/L	09/06/1994	133	
	AST = SGOT	5-35 U/L	10/06/1994	147	
	AST = SGOT	5-35 U/L	22/06/1994	43	
	AST = SGOT	5-35 U/L	14/06/1994	71	
	AST = SGOT	5-35 U/L	12/06/1994	157	

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Report Detail

Case Number: 93939 **Seq:** 1 **Gender:** F
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 06/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FLUCLOXACILLIN SODIUM (Suspected)			**Reason:**		
Injection	4.0	Gram	Daily	Intravenous	
Batch:	**Started:**	03/06/1994		**Stopped:**	06/06/1994
METFORMIN HYDROCHLORIDE (Suspected)			**Reason:**	Diabetes mellitus	
	1.0	Gram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
MINIDIAB (Suspected)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
PREMARIN (Suspected)			**Reason:**		
	300.0	Microgram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	5-35 U/L	11/06/1994	136	
	Bilirubin	0-17 UMOL/L	06/06/1994	114	
	Bilirubin	0-17 UMOL/L	08/06/1994	127	
	Bilirubin	0-17 UMOL/L	09/06/1994	162	
	Bilirubin	0-17 UMOL/L	10/06/1994	152	
	Bilirubin	0-17 UMOL/L	11/06/1994	154	
	Bilirubin	0-17 UMOL/L	12/06/1994	168	

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 93939 **Seq:** 1
Reported: 24/06/1994
Hospitalisation:
Onset Date: 06/06/1994
Outcome: Not yet recovered

Gender: F
Weight:
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detai

Medicine Details:

PROVERA (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	0-17 UMOL/L	14/06/1994	133	
	Bilirubin	0-17 UMOL/L	22/06/1994	64	
	GGT = SGGT =	5-30 U/L	06/06/1994	1371	
	GGT = SGGT =	5-30 U/L	10/06/1994	1325	
	GGT = SGGT =	5-30 U/L	11/06/1994	1522	
	GGT = SGGT =	5-30 U/L	12/06/1994	1725	
	GGT = SGGT =	5-30 U/L	22/06/1994	2508	

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Report Detail

Case Number: 93939 **Seq:** 1 **Gender:** F
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 06/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CALTRATE (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**		**Stopped:**	CONTIN
TRYPTANOL (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**		**Stopped:**	04/06/1994
HEPARIN CALCIUM (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**		**Stopped:**	04/06/1994
FLUCLOXACILLIN SODIUM (Suspected)			**Reason:**	Postoperative wound infection
Injection	4.0 Gram	Daily		Intravenous
Batch:	**Started:**	14/05/1994	**Stopped:**	23/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT =	5-30 U/L	14/06/1994	1811	
	GGT = SGGT =	5-30 U/L	09/06/1994	1309	
	GGT = SGGT =	5-30 U/L	08/06/1994	1339	
	SAP = ALP	40-140 U/L	06/06/1994	1710	
	SAP = ALP	40-140 U/L	08/06/1994	2193	
	SAP = ALP	40-140 U/L	09/06/1994	1996	
	SAP = ALP	40-140 U/L	10/06/1994	2094	

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Report Detail

Case Number: 93939 **Seq:** 1 **Gender:** F
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 06/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FLUCLOXACILLIN SODIUM (Suspected)			**Reason:**		
Injection	4.0	Gram	Daily	Intravenous	
Batch:	**Started:**	03/06/1994		**Stopped:**	06/06/1994
METFORMIN HYDROCHLORIDE (Suspected)			**Reason:**	Diabetes mellitus	
	1.0	Gram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
MINIDIAB (Suspected)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
PREMARIN (Suspected)			**Reason:**		
	300.0	Microgram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP	40-140 U/L	11/06/1994	2349	
	SAP = ALP	40-140 U/L	12/06/1994	2529	
	SAP = ALP	40-140 U/L	14/06/1994	2054	
	SAP = ALP	40-140 U/L	22/06/1994	1596	

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Report Detail

Case Number: 93939 **Seq:** 1
Reported: 24/06/1994
Hospitalisation:
Onset Date: 06/06/1994
Outcome: Not yet recovered

Gender: F
Weight:
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

PROVERA (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Additional Information:

Developed acute cholecystitis with gallstones - required surgery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 93977 **Seq:** 1 **Gender:** M
Reported: 27/06/1994 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Death **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Sudden death			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 94033 **Seq:** 1 **Gender:** F
Reported: 28/06/1994 **Weight:** 48.00
Hospitalisation: **Age:** 38Y
Onset Date: 28/05/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Flushing Hyperhidrosis			

Medicine Details:

ALEPAM (Other drug)			**Reason:**	
Tablet	120.0 Milligram		Daily	Oral
Batch:	**Started:** 01/02/1994		**Stopped:**	
MOGADON (Other drug)			**Reason:**	
Tablet	5.0 Milligram		Daily	Oral
Batch:	**Started:** 01/02/1994		**Stopped:**	
ENDEP (Suspected)			**Reason:**	Depression
Tablet	50.0 Milligram		Daily	Oral
Batch:	**Started:** 25/05/1994		**Stopped:**	31/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Generic substitution problem the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 94189 **Seq:** 1 **Gender:** F
Reported: 04/07/1994 **Weight:** 77.00
Hospitalisation: **Age:** 47Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion			

Medicine Details:

SALBUTAMOL (Other drug) **Reason:**
Inhalation 8.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/04/1994 **Stopped:**

PROPANTHELINE BROMIDE (Other drug) **Reason:** Othr diseases of urinary tract
Tablet 45.0 Milligram Daily Oral
Batch: **Started:** 01/02/1994 **Stopped:**

CLOZAPINE (Suspected) **Reason:** Unspecified schizophrenia
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** 27/05/1994 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 14/06/1994 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 94289 **Seq:** 1 **Gender:** M
Reported: 11/07/1994 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 08/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Coma Confusional state Delusion Hallucination Somnolence			2lt oxygen nasally, haloperidol im 5mg. amitriptyline & baclofen dose reduced

Medicine Details:

THIAMINE (Other drug) **Reason:**
Capsule 100.0 Milligram Daily Oral
Batch: **Started:** 01/06/1994 **Stopped:**

CARBAMAZEPINE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Oral application 200.0 Milligram As necessary Oral
Batch: **Started:** 06/06/1994 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 31/05/1994 **Stopped:** 09/06/1994

LIORESAL (Suspected) **Reason:**
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** 06/06/1994 **Stopped:** 09/06/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Alcohol withdrawal

Report Detail

Case Number: 94292 **Seq:** 1 **Gender:** F
Reported: 08/07/1994 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Increased ast	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Other acne of sebaceous glands

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT		15/02/1993	48	
	AST = SGOT		15/03/1994	55	

Additional Information:

Report Detail

Case Number: 94437 **Seq:** 1 **Gender:** M
Reported: 18/07/1994 **Weight:**
Hospitalisation: **Age:** 84Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal Hyperbilirubinaemia			

Medicine Details:

BACTRIM (Suspected) **Reason:** Othr diseases of urinary tract
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** 06/07/1994

RIVOTRIL (Suspected) **Reason:**
1.0 Gram Daily
Batch: **Started:** **Stopped:** 09/07/1994

TRYPTANOL (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** **Stopped:** 09/07/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 94528 **Seq:** 1 **Gender:** M
Reported: 26/07/1994 **Weight:** 80.00
Hospitalisation: **Age:** 45Y
Onset Date: 13/07/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pneumonia Pyrexia Tachycardia		Atypical pneumonia	

Medicine Details:

CLOZAPINE (Suspected) **Reason:** Unspecified schizophrenia

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 28/06/1994 **Stopped:** 15/07/1994

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 01/07/1994 **Stopped:** 15/07/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 94667 **Seq:** 1 **Gender:** F
Reported: 28/07/1994 **Weight:** 56.00
Hospitalisation: **Age:** 66
Onset Date: 31/03/1993 **DOB:** 28/07/1926
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuropathy peripheral			

Medicine Details:

IMIGRAN (Suspected) **Reason:** Migraine
Tablet 400.0 Milligram Monthly Oral
Batch: **Started:** 01/01/1992 **Stopped:**

STELAZINE (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

FOLIC ACID (Suspected) **Reason:**
5.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

EUHYPNOS (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrographics				Migraine and myeloproliferative disorder. slowly pregressive peripheral neuropathy motor and sensory. nerve conduction studies reveal moderate slowing. may relate to myeloproliferative disorder

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 94667 **Seq:** 1
Reported: 28/07/1994
Hospitalisation:
Onset Date: 31/03/1993
Outcome: Not yet recovered

Gender: F
Weight: 56.00
Age: 66
DOB: 28/07/1926
Causality: Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected)		**Reason:**	
	50.0 Milligram	Daily	
Batch:	**Started:** 01/01/1992		**Stopped:**
SANDOMIGRAN (Suspected)		**Reason:**	
	5.0 Milligram	Daily	
Batch:	**Started:** 01/01/1991		**Stopped:**

Laboratory Investigations:

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 94838 **Seq:** 1
Reported: 05/08/1994
Hospitalisation:
Onset Date: 21/07/1994
Outcome: Not yet recovered

Gender: M
Weight: 62.00
Age: 34
DOB: 22/03/1960
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Cerebrovascular disorder Hemiplegia			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Migraine

100.0 Milligram Daily

Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 95169 **Seq:** 1 **Gender:** F
Reported: 17/08/1994 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: 31/07/1993 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertonia			

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:**

Tablet	600.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**

Oral application	175.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Report Detail

Case Number: 95376 **Seq:** 1 **Gender:** F
Reported: 29/08/1994 **Weight:** 70.00
Hospitalisation: **Age:** 80Y
Onset Date: 25/08/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Rash inbetween fingers	
Oedema peripheral		Swelling of fingers	
Fixed eruption			

Medicine Details:

ALPRIM (Suspected) **Reason:** Othr diseases of urinary tract
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/08/1994 **Stopped:** 25/08/1994

CALTRATE (Suspected) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

CAPOTEN (Suspected) **Reason:**
Oral application 75.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 95376 **Seq:** 1
Reported: 29/08/1994
Hospitalisation:
Onset Date: 25/08/1994
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 80Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

URAL (Suspected) **Reason:** Othr diseases of urinary tract

Oral application 3.0 Dose Unspecified Daily Oral

Batch: **Started:** 22/08/1994 **Stopped:** 25/08/1994

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 95635 **Seq:** 1 **Gender:** F
Reported: 02/09/1994 **Weight:**
Hospitalisation: **Age:** 80Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast enlargement Breast pain		Breast swelling Breast tenderness	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 95749 **Seq:** 1 **Gender:** M
Reported: 08/09/1994 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: 25/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Haematemesis			

Medicine Details:

NAPROXEN (Suspected) **Reason:** Other disturbance of sensation
Oral application 1.0 Gram Daily Oral
Batch: **Started:** 22/02/1994 **Stopped:** 02/03/1994

CALCIPARINE (Suspected) **Reason:** Phleb&thrombophleb lowr extrem
Injection 10.0 Thousand Internat Daily Subcutaneous
Batch: **Started:** 18/02/1994 **Stopped:** 01/03/1994

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** 24/02/1994 **Stopped:** 01/03/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 97000 **Seq:** 1 **Gender:** F
Reported: 09/11/1994 **Weight:** 120.00
Hospitalisation: **Age:** 21Y
Onset Date: 30/10/1994 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 28/10/1994 **Stopped:** 04/11/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recommenced amitripline but reaction occured, then disappeared once drug stopped.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 97138 **Seq:** 1 **Gender:** F
Reported: 17/11/1994 **Weight:**
Hospitalisation: **Age:** AD
Onset Date: 17/09/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anticholinergic syndrome Constipation Dry mouth Dysphagia Visual disturbance			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 6W

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 97439 **Seq:** 1
Reported: 29/11/1994
Hospitalisation:
Onset Date: 26/10/1994
Outcome: Death, maybe drug

Gender: M
Weight: 78.00
Age: 50Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Suicide attempt			

Medicine Details:

PROZAC (Suspected) **Reason:**
0.0
Batch: **Started:** 26/09/1994 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** 04/10/1994 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient died 8/10/94.

Report Detail

Case Number: 97702 **Seq:** 1 **Gender:** M
Reported: 12/12/1994 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 31/07/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast enlargement			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Report Detail

Case Number: 97828 **Seq:** 1 **Gender:** M
Reported: 13/12/1994 **Weight:**
Hospitalisation: **Age:** 72Y
Onset Date: 11/04/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Eosinophilia Hepatic function abnormal Nausea Vomiting			

Medicine Details:

IMDUR (Other drug) **Reason:** Chron isch heart dis no hyper
Oral application 60.0 Milligram Daily Oral
Batch: **Started:** 29/03/1994 **Stopped:**

ASPIRIN (Other drug) **Reason:** Chron isch heart dis no hyper
Oral application 150.0 Milligram Daily Oral
Batch: **Started:** 01/02/1994 **Stopped:**

ZANTAC (Other drug) **Reason:** Chron isch heart dis no hyper
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/02/1994 **Stopped:**

DILANTIN (Suspected) **Reason:** Pain in limb
0.0
Batch: **Started:** 06/04/1994 **Stopped:** 11/04/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT	(10-50 U/L)	16/04/1994	232	
	Bilirubin	(<17-UMOL/L)	16/04/1994	21	
	SAP = ALP	(30-120 U/L)	16/04/1994	1056	
	White blood cells	(4-11)	17/04/1994	11.70	

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 97828 **Seq:** 1 **Gender:** M
Reported: 13/12/1994 **Weight:**
Hospitalisation: **Age:** 72Y
Onset Date: 11/04/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMLODIPINE (Suspected)			**Reason:**	Chron isch heart dis no hyper	
Tablet	10.0	Milligram	Daily		Oral
Batch:	**Started:**	29/03/1994		**Stopped:**	17/04/1994
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Pain in limb	
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**	11/04/1994		**Stopped:**	16/04/1994

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 98655 **Seq:** 1 **Gender:** M
Reported: 01/02/1995 **Weight:** 84.00
Hospitalisation: **Age:** 38Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Dysgeusia Paraesthesia			

Medicine Details:

FAMOTIDINE (Other drug) **Reason:** Otr disorders stomach function

Tablet 40.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Specific disorders of sleep

Tablet 50.0 Milligram Daily Oral

Batch: P2426 **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on amitriptyline with same reaction occurring.

Report Detail

Case Number: 98700 **Seq:** 1 **Gender:** F
Reported: 03/02/1995 **Weight:** 80.00
Hospitalisation: **Age:** 72Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

TIMOPTOL (Other drug) **Reason:**
Eye Drops 0.0 Ophthalmic
Batch: **Started:** **Stopped:**

SOLPRIN (Other drug) **Reason:** Other disturbance of sensation
300.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

ENDURON (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Disturbance of sleep
25.0 Milligram Daily
Batch: **Started:** 14/11/1994 **Stopped:** 21/11/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 98808 **Seq:** 1 **Gender:** F
Reported: 08/02/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Dyspepsia			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 5.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 98835 **Seq:** 1 **Gender:** M
Reported: 08/02/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Psychotic disorder			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Anxiety neurosis
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 99547 **Seq:** 1 **Gender:** F
Reported: 13/03/1995 **Weight:** 53.00
Hospitalisation: **Age:** 75Y
Onset Date: 07/02/1995 **DOB:**
Outcome: Recovered with sequelae **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue Rash Thrombocytopenia			Started prednisone, zantac.

Medicine Details:

NATRILIX (Suspected) **Reason:** Essential benign hypertension

Tablet 1.0 Dose Unspecified Daily Oral

Batch: **Started:** 01/07/1994 **Stopped:** 07/02/1995

TRYPTANOL (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 07/02/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bone marrow				9/2/95 bone marrow look like itp.
	Platelets	(150-400)	08/02/1995	12	
	Platelets	(150-400)	10/02/1995	10	
	Platelets	(150-400)	11/02/1995	47	

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 99570 **Seq:** 1
Reported: 13/03/1995
Hospitalisation: Admitted to hospital
Onset Date: 13/12/1994
Outcome: Not yet recovered

Gender: M
Weight: 76.00
Age: 36
DOB: 20/10/1958
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pancreatitis Pyrexia	Severe		Appropriate diet/analgesia/clonazepam/amitrytiline/haloperidol.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CLONAZEPAM (Other drug) **Reason:** Anxiety neurosis
2.0 Milligram As necessary
Batch: **Started:** **Stopped:**

HALOPERIDOL (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

CLOZARIL (Suspected) **Reason:** Unspecified schizophrenia
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** 24/08/1994 **Stopped:** 04/12/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				4 x ct scans of abdomen during admission showed slowly unlining pancreatic pseudo cyst/ pancreatitis.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 99570 **Seq:** 1
Reported: 13/03/1995
Hospitalisation: Admitted to hospital
Onset Date: 13/12/1994
Outcome: Not yet recovered

Gender: M
Weight: 76.00
Age: 36
DOB: 20/10/1958
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** 22/11/1994 **Stopped:** 13/12/1994

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 100057 **Seq:** 1 **Gender:** F
Reported: 04/04/1995 **Weight:** 0.00
Hospitalisation: **Age:** 44Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anaemia Leukopenia Red blood cell sedimentation rate			

Medicine Details:

STELAZINE (Suspected) **Reason:** Unspecified psychosis
10.0 Milligram Daily Oral
Batch: **Started:** 02/03/1978 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/12/2002	Haemoglobin	(115-165)	12/07/2006	112	
17/12/2002	White blood cells	(4.0-11.0)10^9/L	12/07/2006	3.7	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 100066 **Seq:** 1 **Gender:** M
Reported: 05/04/1995 **Weight:** 100.00
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperhidrosis			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr mus rheum,fibrosit&myalgia

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 100412 **Seq:** 1 **Gender:** F
Reported: 28/04/1995 **Weight:** 60.00
Hospitalisation: **Age:** 77Y
Onset Date: 04/02/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Face oedema Tongue oedema			

Medicine Details:

MYLANTA (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ZANTAC (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

ZOCOR (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

MINIPRESS (Other drug) **Reason:**
6.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Cessation of medication by patient.

Report Detail

Case Number: 100412 **Seq:** 1 **Gender:** F
Reported: 28/04/1995 **Weight:** 60.00
Hospitalisation: **Age:** 77Y
Onset Date: 04/02/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ZYLOPRIM (Other drug)			**Reason:**	
	300.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	
TENORMIN (Other drug)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	
NATRILIX (Other drug)			**Reason:**	
	2.5	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Pain
	0.0			
Batch:	**Started:**	25/01/1995	**Stopped:**	30/01/1995

Laboratory Investigations:

Additional Information:
Cessation of medication by patient.

Report Detail

Case Number: 100534 **Seq:** 1
Reported: 01/05/1995
Hospitalisation:
Onset Date: 20/04/1995
Outcome: Not yet recovered

Gender: M
Weight: 74.00
Age: 30Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Agitation Fatigue Hypertonia Insomnia		Jerking of extremeties	

Medicine Details:

ZOLOFT (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** 18/04/1995 **Stopped:** 26/04/1995

ENDEP (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 13/03/1995 **Stopped:** 18/04/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 100551 **Seq:** 1 **Gender:** F
Reported: 03/05/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: 28/03/1995 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Headache
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** 26/03/1995 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 100979 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:**
Hospitalisation: **Age:** 87Y
Onset Date: 02/05/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Visual and auditory hallucinations	

Medicine Details:

KEFLEX (Other drug) **Reason:** Otr&nos infec&parasit diseases
750.0 Milligram Daily
Batch: **Started:** 31/03/1995 **Stopped:** 06/05/1995

ENDEP (Suspected) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 31/03/1995 **Stopped:** 04/05/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 101009 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia Fatigue Thirst			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: P2425 **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 101442 **Seq:** 1
Reported: 09/06/1995
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 74Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haemorrhage			Recieved 4 units of blood

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 12/05/1995 **Stopped:** 19/05/1995

WARFARIN SODIUM (Interaction) **Reason:**
Tablet 2.5 Milligram Daily Oral
Batch: **Started:** 01/01/1992 **Stopped:** CONTIN

DIGOXIN (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

GASTROBROM (Other drug) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	International		19/05/1995	11.0	
	International		12/05/1995	2.0	

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 101442 **Seq:** 1
Reported: 09/06/1995
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 74Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TITRALAC (Other drug)			**Reason:**		
Tablet	0.0			Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
CLONAZEPAM (Other drug)			**Reason:**		
	0.0				
Batch:	**Started:**			**Stopped:**	
CALCITRIOL (Other drug)			**Reason:**		
	0.5	Microgram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 101656 **Seq:** 1
Reported: 21/06/1995
Hospitalisation:
Onset Date: 17/05/1995
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 84Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aplastic anaemia			

Medicine Details:

TEGRETOL (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
200.0 Milligram Daily
Batch: **Started:** 03/01/1995 **Stopped:** 17/05/1995

TRYPTANOL (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
50.0 Milligram Daily
Batch: **Started:** 01/04/1995 **Stopped:** 20/05/1995

TENORMIN (Suspected) **Reason:** Palpitation
50.0 Milligram Daily
Batch: **Started:** 01/01/1989 **Stopped:**

NORVASC (Suspected) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date of death 10/6/95 the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 101656 **Seq:** 1
Reported: 21/06/1995
Hospitalisation:
Onset Date: 17/05/1995
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 84Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

SOLPRIN (Suspected) **Reason:**
0.5 Dose Unspecified Daily
Batch: **Started:** 01/01/1989 **Stopped:** 17/05/1995

Laboratory Investigations:

Additional Information:

Date of death 10/6/95 the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 102091 **Seq:** 1 **Gender:** M
Reported: 10/07/1995 **Weight:**
Hospitalisation: **Age:** 8Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102543 **Seq:** 1 **Gender:** M
Reported: 01/08/1995 **Weight:**
Hospitalisation: **Age:** 83Y
Onset Date: **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102560 **Seq:** 1 **Gender:** F
Reported: 01/08/1995 **Weight:** 84.00
Hospitalisation: **Age:** 42Y
Onset Date: 08/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation			
Dry mouth			
Dysuria			
Palpitations			
Somnolence			
Tremor			
Visual disturbance			

Medicine Details:

DIANE-35 ED (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

PARACETAMOL (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

SPIRONOLACTONE (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

OMEPRAZOLE (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recommenced aminotriptyline hcl on 17/07/1995 at 12.5mg daily.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 102560 **Seq:** 1
Reported: 01/08/1995
Hospitalisation:
Onset Date: 08/07/1995
Outcome: Recovered

Gender: F
Weight: 84.00
Age: 42Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Tablet	50.0	Milligram	Daily	Oral

Batch: **Started:** 30/07/1994 **Stopped:** 14/07/1995

Laboratory Investigations:

Additional Information:

Patient recommenced aminotriptyline hcl on 17/07/1995 at 12.5mg daily.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 102564 **Seq:** 1 **Gender:** F
Reported: 01/08/1995 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Palpitations			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Lumbalgia

Tablet 0.0 Oral

Batch: **Started:** 01/01/1991 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

When patient recommenced amitriptyline reaction reoccured. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 102569 **Seq:** 1 **Gender:** F
Reported: 01/08/1995 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Unspecified arthritis

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102571 **Seq:** 1 **Gender:** F
Reported: 01/08/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Drug withdrawal syndrome Hyperhidrosis			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102854 **Seq:** 1
Reported: 14/08/1995
Hospitalisation:
Onset Date: 31/10/1994
Outcome: Not yet recovered

Gender: F
Weight: 69.00
Age: 69Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea			

Medicine Details:

RENITEC (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1988 **Stopped:**

MODURETIC (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1988 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Headache
450.0 Milligram Daily
Batch: **Started:** 01/07/1994 **Stopped:** 28/11/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on tryptanol in august 1995 and same reaction occurred. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 103300 **Seq:** 1 **Gender:** U
Reported: 05/09/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Systemic lupus erythematosus			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 103303 **Seq:** 1 **Gender:** F
Reported: 05/09/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/03/1995 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Report Detail

Case Number: 103680 **Seq:** 1 **Gender:** M
Reported: 25/09/1995 **Weight:** 81.00
Hospitalisation: **Age:** 24Y
Onset Date: 09/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erectile dysfunction			

Medicine Details:

TEGRETOL (Other drug) **Reason:** Pain

Tablet	800.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Pain

Tablet	75.0	Milligram	Daily	Oral

Batch: **Started:** 29/08/1995 **Stopped:** 14/09/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 103867 **Seq:** 1 **Gender:** F
Reported: 05/09/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Fatigue			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Migraine

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 01/09/1995 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 103880 **Seq:** 1 **Gender:** M
Reported: 05/10/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Constipation			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 104818 **Seq:** 1 **Gender:** F
Reported: 23/11/1995 **Weight:** 77.00
Hospitalisation: **Age:** 74Y
Onset Date: 05/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular			

Medicine Details:

AUGMENTIN FORTE (Suspected) **Reason:** Agranulocytosis
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 06/11/1995

PARACETAMOL (Suspected) **Reason:** Pain
Oral application 4.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

MAXOLON (Suspected) **Reason:** Nausea and vomiting
Oral application 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 104818 **Seq:** 1 **Gender:** F
Reported: 23/11/1995 **Weight:** 77.00
Hospitalisation: **Age:** 74Y
Onset Date: 05/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

COLOXYL WITH SENNA (Suspected) **Reason:** Constipation
Oral application 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 105055 **Seq:** 1 **Gender:** F
Reported: 04/12/1995 **Weight:**
Hospitalisation: **Age:** 82Y
Onset Date: 16/11/1995 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Hepatitis			

Medicine Details:

FRUSEMIDE (Other drug) **Reason:** Left ventricular failure
Oral application 80.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

THIORIDAZINE HYDROCHLORIDE (Other drug) **Reason:** Reactive confusion
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

MONOPRIL (Suspected) **Reason:** Essential benign hypertension
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 17/11/1995

DUCENE (Suspected) **Reason:** Left ventricular failure
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 16/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		16/11/1995	819	
	Bilirubin		16/11/1995	23	
	GGT = SGGT = [illegible]		16/11/1995	512	
	LDH		16/11/1995	658	
	SAP = ALP		16/11/1995	286	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105055 **Seq:** 1 **Gender:** F
Reported: 04/12/1995 **Weight:**
Hospitalisation: **Age:** 82Y
Onset Date: 16/11/1995 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 17/11/1995

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105288 **Seq:** 1 **Gender:** F
Reported: 12/12/1995 **Weight:** 50.00
Hospitalisation: **Age:** 84Y
Onset Date: 30/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hyponatraemia			

Medicine Details:

FRAGMIN (Other drug) **Reason:** Prophylaxis
Injection 5.0 Thousand Internat Daily Subcutaneous
Batch: **Started:** 27/11/1995 **Stopped:** 08/12/1995

FRUSEMIDE (Other drug) **Reason:** Congestive heart failure
Oral application 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

SLOW-K (Other drug) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** CONTIN

MORPHINE NOS (Suspected) **Reason:** Pain
Injection 0.0 Intravenous
Batch: **Started:** 28/11/1995 **Stopped:** 30/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium		30/11/1995	125	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105288 **Seq:** 1
Reported: 12/12/1995
Hospitalisation:
Onset Date: 30/11/1995
Outcome: Recovered

Gender: F
Weight: 50.00
Age: 84Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105517 **Seq:** 1 **Gender:** M
Reported: 23/12/1995 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** 01/01/1993 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105519 **Seq:** 1 **Gender:** F
Reported: 23/12/1995 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 01/01/1990 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105686 **Seq:** 1 **Gender:** M
Reported: 03/01/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Aggression			
Agitation			
Anxiety			
Dizziness			
Dry mouth			
Tremor			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 105784 **Seq:** 1 **Gender:** M
Reported: 10/01/1996 **Weight:**
Hospitalisation: **Age:** A6
Onset Date: 03/01/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Raised, red lumpy rash across forehead	

Medicine Details:

LOSEC (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:** Incontinence of urine
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 02/01/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105818 **Seq:** 1 **Gender:** M
Reported: 12/01/1996 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 26/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urine analysis abnormal Hepatic function abnormal Malaise		Dark urine	

Medicine Details:

CLINORIL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 27/12/1995

TOFRANIL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 27/12/1995

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 27/12/1995

AURORIX (Suspected) **Reason:**
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 15/11/1995 **Stopped:** 27/12/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 105945 **Seq:** 1 **Gender:** M
Reported: 18/01/1996 **Weight:** 82.00
Hospitalisation: **Age:** 54Y
Onset Date: 21/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Hyperhidrosis Nightmare Sleep disorder			

Medicine Details:

TRYPTANOL (Other drug) **Reason:**
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** 01/10/1995 **Stopped:** 21/12/1995

ANATENSOL (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

PRINIVIL (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** 01/06/1995 **Stopped:**

VALIUM (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient substituted endep 25 mg tab for tryptanol 25mg tab as unavailable and reaction occurred 15-20 min after taking endep. resumed on tryptanol and symptoms ceased. history of allergy to sulphur. generic substitution problem the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 105945 **Seq:** 1 **Gender:** M
Reported: 18/01/1996 **Weight:** 82.00
Hospitalisation: **Age:** 54Y
Onset Date: 21/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:**

Tablet	25.0	Milligram	1 time	Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Patient substituted endep 25 mg tab for tryptanol 25mg tab as unavailable and reaction occurred 15-20 min after taking endep. resumed on tryptanol and symptoms ceased. history of allergy to sulphur. generic substitution problem the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 106068 **Seq:** 1 **Gender:** F
Reported: 29/01/1996 **Weight:** 60.00
Hospitalisation: **Age:** 65Y
Onset Date: 28/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dehydration Somnolence			

Medicine Details:

ENDEP (Suspected) **Reason:**
75.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 106197 **Seq:** 1 **Gender:** F
Reported: 01/02/1996 **Weight:**
Hospitalisation: **Age:** 66
Onset Date: **DOB:** 15/08/1929
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Eczema			

Medicine Details:

ENDEP (Suspected) **Reason:**

0.0

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had a previous history of a similar rash in 1983 shortly after taking one 2mg tablet of dexchlorpheniramine maleate. lactose, maize/corn starch and magnesium stearate are common to endep and dexchlorpheniramine maleate.

Report Detail

Case Number: 106220 **Seq:** 1 **Gender:** F
Reported: 02/02/1996 **Weight:** 52.00
Hospitalisation: **Age:** 79Y
Onset Date: 15/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatocellular damage			

Medicine Details:

TEGRETOL (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
100.0 Milligram Daily
Batch: **Started:** 01/05/1995 **Stopped:** 28/01/1996

SURGAM (Suspected) **Reason:** Osteoarthritis
200.0 Milligram Daily
Batch: **Started:** 01/11/1995 **Stopped:** 28/01/1996

ENDEP (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
25.0 Milligram Daily
Batch: **Started:** 01/05/1995 **Stopped:** 28/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	0 - 50 U/L	22/01/1996	2380	
	ALT = SGPT	0 - 50 U/L	01/03/1996	14	
	AST = SGOT	11 - 33 U/L	22/01/1996	2460	
	AST = SGOT	11 - 33 U/L	01/03/1996	19	
	Bilirubin	0 - 20 UMOL/L	22/01/1996	20	
	Bilirubin	0 - 20 UMOL/L	01/03/1996	10	
	GGT = SGGT =	0 - 50 U/L	22/01/1996	681	

Additional Information:

Patient drinks 1glass of wine and 1 can of beer a day. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Report Detail

Case Number: 106220 **Seq:** 1 **Gender:** F
Reported: 02/02/1996 **Weight:** 52.00
Hospitalisation: **Age:** 79Y
Onset Date: 15/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TEGRETOL (Suspected)		**Reason:**	Otr&unsp forms neuralg&neurit
	100.0 Milligram	Daily	
Batch:	**Started:** 01/05/1995		**Stopped:** 28/01/1996
SURGAM (Suspected)		**Reason:**	Osteoarthritis
	200.0 Milligram	Daily	
Batch:	**Started:** 01/11/1995		**Stopped:** 28/01/1996
ENDEP (Suspected)		**Reason:**	Otr&unsp forms neuralg&neurit
	25.0 Milligram	Daily	
Batch:	**Started:** 01/05/1995		**Stopped:** 28/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT = GGTP	0 - 50 U/L	01/03/1996	57	
	SAP = ALP	30 - 120 U/L	22/01/1996	215	
	SAP = ALP	30 - 120 U/L	01/03/1996	95	

Additional Information:

Patient drinks 1glass of wine and 1 can of beer a day. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Report Detail

Case Number: 106502 **Seq:** 1 **Gender:** M
Reported: 16/02/1996 **Weight:** 78.00
Hospitalisation: **Age:** 52Y
Onset Date: 16/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Dizziness Mania			

Medicine Details:

VOLTAREN (Other drug) **Reason:**
Tablet 50.0 Milligram As necessary Oral
Batch: **Started:** **Stopped:** CONTIN

PANADEINE FORTE (Other drug) **Reason:**
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 15/01/1996 **Stopped:** 24/01/1996

PHENERGAN (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 15/01/1996 **Stopped:** 20/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 106815 **Seq:** 1 **Gender:** M
Reported: 01/03/1996 **Weight:**
Hospitalisation: **Age:** 72Y
Onset Date: 19/02/1996 **DOB:**
Outcome: Death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sudden death Tremor			

Medicine Details:

MORPHINE NOS (Suspected) **Reason:** Pain
Injection, intravenous infusion 90.0 Milligram Daily Subcutaneous
Batch: **Started:** 07/02/1996 **Stopped:** 20/02/1996

MAXOLON (Suspected) **Reason:** Nausea and vomiting
Injection, intravenous infusion 60.0 Milligram Daily Subcutaneous
Batch: **Started:** 07/02/1996 **Stopped:** 15/02/1996

STEMETIL (Suspected) **Reason:** Nausea and vomiting
Suppository 50.0 Milligram Daily Rectal
Batch: **Started:** 13/02/1996 **Stopped:** 17/02/1996

ENDEP (Suspected) **Reason:** Specific disorders of sleep
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 13/02/1996 **Stopped:** 17/02/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Palliative care patient: disseminated scc lung, enlarged liver, jaundiced.

Report Detail

Case Number: 107505 **Seq:** 1 **Gender:** F
Reported: 29/03/1996 **Weight:** 58.00
Hospitalisation: **Age:** 63Y
Onset Date: 27/03/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Angioedema			

Medicine Details:

TENORMIN (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 02/11/1994 **Stopped:**

NORVASC (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 09/11/1994 **Stopped:**

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** 11/02/1994 **Stopped:**

RENITEC (Suspected) **Reason:** Essential benign hypertension
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 22/12/1993 **Stopped:** 27/03/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 107505 **Seq:** 1
Reported: 29/03/1996
Hospitalisation:
Onset Date: 27/03/1996
Outcome: Not yet recovered

Gender: F
Weight: 58.00
Age: 63Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected)		**Reason:**	
	75.0 Milligram	Daily	
Batch:	**Started:** 19/03/1996		**Stopped:**
RULIDE (Suspected)		**Reason:**	
	150.0 Milligram	Daily	
Batch:	**Started:** 19/03/1996		**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 107698 **Seq:** 1 **Gender:** F
Reported: 02/04/1996 **Weight:** 80.00
Hospitalisation: **Age:** 34Y
Onset Date: 22/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing Pruritus			 Teldane

Medicine Details:

PETHIDINE HYDROCHLORIDE (Other drug) **Reason:** Migraine
Injection 100.0 Milligram As necessary Intramuscular
Batch: **Started:** L TERM **Stopped:**

MAXOLON (Other drug) **Reason:** Migraine
Injection 1.0 Dose Unspecified As necessary Intramuscular
Batch: **Started:** L TERM **Stopped:**

STEMETIL (Other drug) **Reason:** Migraine
Injection 12.0 Milligram As necessary Intramuscular
Batch: **Started:** 01/01/1994 **Stopped:**

BETALOC (Other drug) **Reason:** Essential benign hypertension
200.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previously had severe vomiting with morphine. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 107698 **Seq:** 1 **Gender:** F
Reported: 02/04/1996 **Weight:** 80.00
Hospitalisation: **Age:** 34Y
Onset Date: 22/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

SANDOMIGRAN (Suspected)				**Reason:**	Migraine	
Injection	4.0	Dose Unspecified	Daily			Intravenous
Batch:	**Started:**	01/01/1994			**Stopped:**	28/02/1996
ENDEP (Suspected)				**Reason:**	Migraine	
	50.0	Milligram	Daily			
Batch:	**Started:**	01/11/1995			**Stopped:**	28/02/1996

Laboratory Investigations:

Additional Information:

Previously had severe vomiting with morphine. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 107992 **Seq:** 1 **Gender:** F
Reported: 22/04/1996 **Weight:**
Hospitalisation: **Age:** 69
Onset Date: **DOB:** 28/06/1926
Outcome: Unknown **Causality:** Causality probable

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Dry mouth Nightmare			Changed to tryptanol.

Medicine Details:

ENDEP (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 2W

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Generic substitution problem

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 108212 **Seq:** 1
Reported: 06/05/1996
Hospitalisation:
Onset Date: 01/04/1996
Outcome: Not yet recovered

Gender: M
Weight: 90.00
Age: 27Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing Hyperhidrosis			

Medicine Details:

ENDEP (Other drug) **Reason:** Depression
100.0 Milligram Daily
Batch: **Started:** 01/04/1996 **Stopped:**

ENDEP (Suspected) **Reason:** Migraine
10.0 Milligram Daily
Batch: **Started:** 11/03/1996 **Stopped:** 01/04/1996

RISPERDAL (Suspected) **Reason:**
3.0 Dose Unspecified Daily
Batch: **Started:** 12/10/1995 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 108837 **Seq:** 1 **Gender:** F
Reported: 29/05/1996 **Weight:**
Hospitalisation: **Age:** 15Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Fibroadenoma of breast		Breast cyst	

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Headache
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 108846 **Seq:** 1 **Gender:** F
Reported: 29/05/1996 **Weight:**
Hospitalisation: **Age:** 57Y
Onset Date: 02/05/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Diarrhea
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 01/05/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109033 **Seq:** 1 **Gender:** F
Reported: 07/06/1996 **Weight:**
Hospitalisation: **Age:** 80Y
Onset Date: 31/05/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Dry mouth Flushing			

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

LOSEC (Suspected) **Reason:**
Capsule 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 109105 **Seq:** 1 **Gender:** F
Reported: 07/06/1996 **Weight:** 60.00
Hospitalisation: **Age:** 66Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective			

Medicine Details:

TRYPTINE (Suspected) **Reason:** Specific disorders of sleep
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Generic substitution problem

Report Detail

Case Number: 109231 **Seq:** 1
Reported: 18/06/1996
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: M
Weight:
Age:
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myocardial infarction Pulmonary embolism			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109232 **Seq:** 1 **Gender:** M
Reported: 18/06/1996 **Weight:**
Hospitalisation: Admitted to hospital **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myocardial infarction Pulmonary embolism			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109263 **Seq:** 1 **Gender:** M
Reported: 19/06/1996 **Weight:** 78.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrasystoles Convulsion		Ventricular ectopics	

Medicine Details:

AROPAX (Suspected) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 01/05/1995 **Stopped:** 28/05/1996

ENDEP (Suspected) **Reason:** Specific disorders of sleep
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 01/06/1995 **Stopped:** 28/04/1996

PROPRANOLOL HYDROCHLORIDE (Suspected) **Reason:** Essential benign hypertension
Tablet 80.0 Milligram Daily Oral
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

RIVOTRIL (Suspected) **Reason:** Anxiety neurosis
Oral application 500.0 Microgram Daily Oral
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrographics				Holter monitor - veb(vt x once 5 beats only with no symptoms)

Additional Information:

Ceased aropax, changed to zoloft - myoclonus cured. veb persists but improved. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 109273 **Seq:** 1 **Gender:** M
Reported: 18/06/1996 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 24/04/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention			

Medicine Details:

DEXAMETHASONE (Other drug) **Reason:**
Oral application 4.0 Milligram Daily Oral
Batch: **Started:** 23/04/1996 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 16/04/1996 **Stopped:** 26/04/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109501 **Seq:** 1 **Gender:** M
Reported: 26/06/1996 **Weight:**
Hospitalisation: **Age:** 5Y
Onset Date: 16/06/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective		Hyperactivity	

Medicine Details:

METHYLPHENIDATE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:** Behavior disorders of childhod
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 16/06/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 109511 **Seq:** 1 **Gender:** U
Reported: 27/06/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Incontinence of urine
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 109515 **Seq:** 1 **Gender:** U
Reported: 27/06/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Breast enlargement			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Incontinence of urine
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109516 **Seq:** 1 **Gender:** U
Reported: 27/06/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Hypotension			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Incontinence of urine
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 109518 **Seq:** 1 **Gender:** U
Reported: 27/06/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Hypertension			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Incontinence of urine
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 110039 **Seq:** 1 **Gender:** M
Reported: 18/07/1996 **Weight:**
Hospitalisation: **Age:** 9Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/07/1995 **Stopped:** 28/07/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 110181 **Seq:** 1 **Gender:** M
Reported: 24/07/1996 **Weight:**
Hospitalisation: **Age:** 9Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Aggression		Patient was violent, angry and tearful	

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 110414 **Seq:** 1
Reported: 01/08/1996
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight: 76.00
Age: 52Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

1.0 Dose Unspecified Daily

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of chronic dizziness

Report Detail

Case Number: 110598 **Seq:** 1 **Gender:** F
Reported: 08/08/1996 **Weight:** 115.00
Hospitalisation: **Age:** 52Y
Onset Date: 11/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paranoia		Suspiousness of being talked about	

Medicine Details:

PANADOL (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
2.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

COVERSYL (Other drug) **Reason:** Essential benign hypertension
4.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
15.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

LASIX (Other drug) **Reason:** Essential benign hypertension
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Brief postpartum psychosis 1964.

Report Detail

Case Number: 110598 **Seq:** 1 **Gender:** F
Reported: 08/08/1996 **Weight:** 115.00
Hospitalisation: **Age:** 52Y
Onset Date: 11/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

PEPCIDINE (Other drug)			**Reason:**	Other diseases of esophagus	
	40.0	Milligram	Daily		
Batch:	**Started:**	L TERM	**Stopped:**	CONTIN	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Headache	
	25.0	Milligram	Daily		
Batch:	**Started:**	05/06/1996	**Stopped:**	11/06/1996	

Laboratory Investigations:

Additional Information:

Brief postpartum psychosis 1964.

Report Detail

Case Number: 110906 **Seq:** 1 **Gender:** M
Reported: 16/08/1996 **Weight:** 70.00
Hospitalisation: **Age:** 24Y
Onset Date: 01/08/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Stevens-Johnson syndrome			Topical corticosteroid cream initially(unresponsive), short course of oral prednisolone, promethazine

Medicine Details:

Keflin Neutral (Suspected) **Reason:** Otr&nos infec&parasit diseases
Injection 12.0 Gram Daily Intravenous
Batch: **Started:** 08/07/1996 **Stopped:** 01/08/1996

FLOPEN (Suspected) **Reason:** Otr&nos infec&parasit diseases
Injection 4.0 Gram Daily Intravenous
Batch: **Started:** 13/06/1996 **Stopped:** 08/07/1996

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** 01/05/1996 **Stopped:** CONTIN

INDOMETHACIN (Suspected) **Reason:**
Oral application 150.0 Milligram Daily Oral
Batch: **Started:** 30/05/1996 **Stopped:** 03/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 110906 **Seq:** 1
Reported: 16/08/1996
Hospitalisation:
Onset Date: 01/08/1996
Outcome: Not yet recovered

Gender: M
Weight: 70.00
Age: 24Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ASPIRIN (Suspected) **Reason:** Prophylaxis

Oral application 150.0 Milligram Daily Oral

Batch: **Started:** 11/05/1996 **Stopped:** CONTIN

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 111349 **Seq:** 1
Reported: 03/09/1996
Hospitalisation:
Onset Date: 30/07/1996
Outcome: Not yet recovered

Gender: M
Weight: 65.00
Age: 16Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Aggression		Visual hallucination	

Medicine Details:

LOSEC (Other drug) **Reason:** Other diseases of esophagus
Capsule 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/05/1996 **Stopped:** CONTIN

DEXAMPHETAMINE SULPHATE (Other drug) **Reason:** Behavior disorders of childhod
25.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Specific disorders of sleep
20.0 Milligram Daily
Batch: **Started:** 30/07/1996 **Stopped:** 10/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 112600 **Seq:** 1
Reported: 28/10/1996
Hospitalisation:
Onset Date: 25/09/1996
Outcome: Death, maybe drug

Gender: F
Weight: 69.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pneumonia Renal failure acute			

Medicine Details:

ULTRAVIST 300 (Suspected) **Reason:** Investigation of circulatory system
50.0 Millilitre 1 time
Batch: **Started:** 25/09/1996 **Stopped:**

CIPROXIN (Suspected) **Reason:**
500.0 Milligram Daily
Batch: **Started:** 18/09/1996 **Stopped:**

SALAZOPYRIN (Suspected) **Reason:**
1.0 Gram Daily
Batch: **Started:** L TERM **Stopped:** 20/09/1996

SALAZOPYRIN (Suspected) **Reason:**
1.5 Gram Daily
Batch: **Started:** 20/09/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatinine		16/09/1996	0.08	
	Creatinine		29/09/1996	0.55	
	White blood cells		16/09/1996	-	
	White blood cells		29/09/1996	21.3	

Additional Information:

Patient died 4/10/96

Report Detail

Case Number: 112600 **Seq:** 1
Reported: 28/10/1996
Hospitalisation:
Onset Date: 25/09/1996
Outcome: Death, maybe drug

Gender: F
Weight: 69.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MS CONTIN (Suspected)			**Reason:**	
	20.0 Milligram		Daily	
Batch:	**Started:** 17/09/1996			**Stopped:**
ZINVIT C-250 (Suspected)			**Reason:**	
	2.0 Dose Unspecified		Daily	
Batch:	**Started:** 18/09/1996			**Stopped:**
TRYPTANOL (Suspected)			**Reason:**	
	75.0 Milligram		Daily	
Batch:	**Started:**	L TERM		**Stopped:**
RENITEC (Suspected)			**Reason:**	
	20.0 Milligram		Daily	
Batch:	**Started:**	L TERM		**Stopped:**

Laboratory Investigations:

Additional Information:

Patient died 4/10/96

Report Detail

Case Number: 112600 **Seq:** 1
Reported: 28/10/1996
Hospitalisation:
Onset Date: 25/09/1996
Outcome: Death, maybe drug

Gender: F
Weight: 69.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

NAPROSYN SR (Suspected) **Reason:**
1.0 Gram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Additional Information:

Patient died 4/10/96

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 113221 **Seq:** 1 **Gender:** F
Reported: 18/11/1996 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			Therapy changed back to endep 50mg.

Medicine Details:

ENDEP (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 113345 **Seq:** 1 **Gender:** M
Reported: 25/11/1996 **Weight:** 74.00
Hospitalisation: **Age:** 39Y
Onset Date: 01/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction			

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
2.0 Milligram As necessary
Batch: **Started:** 01/01/1994 **Stopped:**

PANADEINE FORTE (Other drug) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** 01/01/1994 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** 01/06/1996 **Stopped:** 06/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 113364 **Seq:** 1 **Gender:** F
Reported: 25/11/1996 **Weight:** 80.00
Hospitalisation: **Age:** 36Y
Onset Date: 17/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bronchospasm Dizziness			

Medicine Details:

ENDEP (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 17/11/1996 **Stopped:** 17/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 114016 **Seq:** 1 **Gender:** F
Reported: 19/12/1996 **Weight:** 72.00
Hospitalisation: **Age:** 49Y
Onset Date: 21/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cough		Nocturnal cough	Cessation of amitriptyline

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 20/11/1996 **Stopped:** 10/12/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Cough commenced 24 hours post initiation of amitriptyline: symptoms commenced 15 minutes after administration of amitriptyline.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 114329 **Seq:** 1 **Gender:** F
Reported: 06/01/1997 **Weight:**
Hospitalisation: **Age:** 79
Onset Date: 29/08/1996 **DOB:** 27/03/1917
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pain Somnolence			Morphine 6mg.

Medicine Details:

CARBAMAZEPINE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 114342 **Seq:** 1 **Gender:** F
Reported: 06/01/1997 **Weight:**
Hospitalisation: **Age:** 79
Onset Date: 21/08/1996 **DOB:** 27/03/1917
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Constipation		Not eating	Coloxyl with senna, sorbilax

Medicine Details:

MORPHINE NOS (Suspected) **Reason:**
6.0 Milligram As necessary
Batch: **Started:** **Stopped:**

MS CONTIN (Suspected) **Reason:** Pain
60.0 Milligram Daily
Batch: **Started:** **Stopped:**

PULMICORT (Suspected) **Reason:**
800.0 Microgram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 03/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Mist morphine 6mg prn for breakthrough pain. patient has hisory of morphine and panadeine forte causing severe constipation.

Report Detail

Case Number: 114342 **Seq:** 1 **Gender:** F
Reported: 06/01/1997 **Weight:**
Hospitalisation: **Age:** 79
Onset Date: 21/08/1996 **DOB:** 27/03/1917
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FAMOTIDINE (Suspected)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	30/09/1996
PIROXICAM (Suspected)			**Reason:**	
	10.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	30/08/1996

Laboratory Investigations:

Additional Information:

Mist morphine 6mg prn for breakthrough pain. patient has hisory of morphine and panadeine forte causing severe constipation.

Report Detail

Case Number: 114833 **Seq:** 1
Reported: 29/01/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 40Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome			

Medicine Details:

SERTRALINE HYDROCHLORIDE (Interaction) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

PERICYAZINE (Interaction) **Reason:**
7.5 Milligram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
75.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 115461 **Seq:** 1 **Gender:** F
Reported: 19/02/1997 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 31/12/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Saliva altered Dental caries Dry mouth		Gross change in saliva	

Medicine Details:

DESERIL (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

PREMARIN (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

DUCENE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year.

Report Detail

Case Number: 115685 **Seq:** 1 **Gender:** F
Reported: 27/02/1997 **Weight:**
Hospitalisation: **Age:** 29Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suppressed lactation			

Medicine Details:

ZOLOFT (Suspected) **Reason:**
Tablet 12.5 Milligram Daily Oral
Batch: **Started:** **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 115883 **Seq:** 1 **Gender:** M
Reported: 12/03/1997 **Weight:** 73.00
Hospitalisation: **Age:** 59Y
Onset Date: 24/02/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia			

Medicine Details:

PANADOL (Suspected) **Reason:** Pain
Oral application 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

TICLID (Suspected) **Reason:**
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** 21/01/1997 **Stopped:** 25/02/1997

LIPEX (Suspected) **Reason:** Othr&unspec metabolic diseases
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Neutrophils	2.00 - 7.50 X	05/02/1997	4.74	
	Neutrophils	2.00 - 7.50 X	24/02/1997	1.04	
	Neutrophils	2.00 - 7.50 X	01/04/1997	3.83	
	Neutrophils	2.00 - 7.50 X	05/03/1997	3.17	
	White blood cells		01/04/1997	5.4	
	leucocytes	4.0 - 11.0 X	05/02/1997	6.6	
	leucocytes	4.0 - 11.0 X	24/02/1997	2.3	

Additional Information:

Report Detail

Case Number: 115883 **Seq:** 1
Reported: 12/03/1997
Hospitalisation:
Onset Date: 24/02/1997
Outcome: Not yet recovered

Gender: M
Weight: 73.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

DISPRIN (Suspected) **Reason:**

Oral application	150.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	leucocytes	4.0 - 11.0 X	05/03/1997	4.7	

Additional Information:

Report Detail

Case Number: 115918 **Seq:** 1 **Gender:** U
Reported: 12/03/1997 **Weight:**
Hospitalisation: **Age:** 76
Onset Date: 07/01/1997 **DOB:** 10/05/1920
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention			

Medicine Details:

DI-GESIC (Suspected) **Reason:** Pain
2.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

MODECATE (Suspected) **Reason:** Unspecified schizophrenia
Injection 50.0 Milligram Monthly Intravenous
Batch: **Started:** **Stopped:**

ARTANE (Suspected) **Reason:** Paralysis agitans
15.0 Milligram Daily
Batch: **Started:** **Stopped:** 20/01/1997

SURGAM (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
900.0 Milligram Daily
Batch: **Started:** **Stopped:** 20/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 115918 **Seq:** 1 **Gender:** U
Reported: 12/03/1997 **Weight:**
Hospitalisation: **Age:** 76
Onset Date: 07/01/1997 **DOB:** 10/05/1920
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

SEREPAX (Suspected)			**Reason:**	Otr spec symp psychopathol nec
	90.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
TRYPTANOL (Suspected)			**Reason:**	Otr spec symp psychopathol nec
	75.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 116149 **Seq:** 1
Reported: 19/03/1997
Hospitalisation:
Onset Date: 28/02/1997
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous Pruritus	Severe	Rash on face, neck, shoulders and chest.	

Medicine Details:

ENDEP (Suspected) **Reason:** Specific disorders of sleep
20.0 Milligram Daily
Batch: **Started:** 24/02/1997 **Stopped:** 03/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergy to penicillin.

Report Detail

Case Number: 116389 **Seq:** 1 **Gender:** F
Reported: 01/04/1997 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

PANADOL (Suspected) **Reason:** Osteoarthritis
1.0 Gram As necessary
Batch: **Started:** **Stopped:**

TEGRETOL (Suspected) **Reason:**
400.0 Milligram Daily
Batch: **Started:** **Stopped:** 27/02/1997

VERAPAMIL HYDROCHLORIDE (Suspected) **Reason:** Essential benign hypertension
240.0 Milligram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
10.0 Milligram Daily
Batch: **Started:** **Stopped:** 24/02/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT = SAP = ALP				Peak ggt - 1172 Peak sap - 523

Additional Information:

Past history of hepatitis a.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 116389 **Seq:** 1
Reported: 01/04/1997
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight:
Age: 86Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

COLOXYL WITH SENNA (Suspected)		**Reason:**	Constipation
	4.0 Dose Unspecified	Daily	
Batch:	**Started:**		**Stopped:**
TRIMETHOPRIM (Suspected)		**Reason:**	Othr diseases of urinary tract
	300.0 Milligram	Daily	
Batch:	**Started:** 10/03/1997		**Stopped:** 14/03/1997

Laboratory Investigations:

Additional Information:

Past history of hepatitis a.

Report Detail

Case Number: 116660 **Seq:** 1 **Gender:** F
Reported: 10/04/1997 **Weight:**
Hospitalisation: **Age:** 88
Onset Date: 10/02/1997 **DOB:** 20/10/1908
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium		Anticholinergic delirium	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

PRO-BANTHINE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 116963 **Seq:** 1 **Gender:** M
Reported: 22/04/1997 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 12/04/1997 **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Pulmonary embolism Sudden death		Off balance and staggering	

Medicine Details:

LOVAN (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** 01/01/1997 **Stopped:**

TRYPTANOL (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** 05/04/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient died of pulmonary embolus. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 116967 **Seq:** 1 **Gender:** M
Reported: 21/04/1997 **Weight:** 45.00
Hospitalisation: **Age:** 8
Onset Date: 16/03/1997 **DOB:** 08/08/1988
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Prolonged seizure	

Medicine Details:

LAMOTRIGINE (Other drug) **Reason:** Other&unspecified epilepsy
250.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** 06/03/1997 **Stopped:** 16/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 117259 **Seq:** 1 **Gender:** F
Reported: 05/05/1997 **Weight:**
Hospitalisation: **Age:** 80
Onset Date: **DOB:** 23/09/1916
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension Syncope			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
75.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 118032 **Seq:** 1 **Gender:** M
Reported: 04/06/1997 **Weight:**
Hospitalisation: **Age:** 72Y
Onset Date: 25/05/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Transaminitis	

Medicine Details:

AMIODARONE HYDROCHLORIDE (Suspected) **Reason:** Otr venous embolism&thrombosis
Tablet 1.2 Gram Daily Oral
Batch: **Started:** 23/05/1997 **Stopped:** 29/05/1997

FRUSEMIDE (Suspected) **Reason:**
Tablet 160.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

CAPTOPRIL (Suspected) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 29/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				25/05/97 ast/alt approx 2500

Additional Information:

On amiodarone 200mg po bd for 2 weeks only. mild transaminitis. when dose increased to 400mg tds ---> massive increase in ast/alt

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 118785 **Seq:** 1 **Gender:** F
Reported: 03/07/1997 **Weight:** 90.00
Hospitalisation: **Age:** 23Y
Onset Date: 12/06/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous		Rash on neck, trunk, and arms.	

Medicine Details:

FIORINAL (Other drug) **Reason:** Headache
2.0 Dose Unspecified As necessary
Batch: **Started:** 02/04/1996 **Stopped:**

TRYPTINE (Suspected) **Reason:** Headache
25.0 Milligram Daily
Batch: **Started:** 02/06/1997 **Stopped:** 13/06/1997

TEGRETOL (Suspected) **Reason:**
Injection 400.0 Milligram Daily Intravenous
Batch: **Started:** 02/06/1997 **Stopped:** 13/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 119078 **Seq:** 1 **Gender:** M
Reported: 14/07/1997 **Weight:**
Hospitalisation: **Age:** 22Y
Onset Date: 07/07/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blood amylase increased Salivary gland enlargement			

Medicine Details:

CLOZAPINE (Suspected) **Reason:** Unspecified schizophrenia

0.0

Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Unspecified schizophrenia

Oral application 0.0 Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Amylase		07/07/1997	285	

Additional Information:

Report Detail

Case Number: 119320 **Seq:** 1 **Gender:** F
Reported: 22/07/1997 **Weight:**
Hospitalisation: **Age:** 92Y
Onset Date: 11/06/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Sudden elevation of liver enzymes.	Amitriptyline ceased

Medicine Details:

PANADEINE FORTE (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** 30/05/1997 **Stopped:** 13/06/1997

COLOXYL WITH SENNA (Suspected) **Reason:**
2.0 Dose Unspecified As necessary
Batch: **Started:** 30/05/1997 **Stopped:** 12/06/1997

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Headache
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** 02/06/1997 **Stopped:** 13/06/1997

CIPROFLOXACIN (Suspected) **Reason:** Other diseases of lung
Oral application 1.0 Gram Daily Oral
Batch: **Started:** 06/06/1997 **Stopped:** 10/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 119320 **Seq:** 1 **Gender:** F
Reported: 22/07/1997 **Weight:**
Hospitalisation: **Age:** 92Y
Onset Date: 11/06/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRIMETHOPRIM (Suspected)			**Reason:**	Other diseases of lung	
Oral application	300.0	Milligram	Daily		Oral
Batch:	**Started:**	10/06/1997		**Stopped:**	13/06/1997
TEMAZEPAM (Suspected)			**Reason:**		
Oral application	20.0	Milligram	Daily		Oral
Batch:	**Started:**	30/05/1997		**Stopped:**	17/06/1997

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 119453 **Seq:** 1 **Gender:** M
Reported: 28/07/1997 **Weight:** 70.00
Hospitalisation: **Age:** 59Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vertigo	Severe	Acute vertigo	

Medicine Details:

ALPRAZOLAM (Other drug) **Reason:**
Tablet 2.0 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Disturbance of sleep
Tablet 25.0 Milligram 1 time Oral
Batch: **Started:** 22/07/1997 **Stopped:** 23/07/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had no previous exposure of tricyclic antidepressants. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 119814 **Seq:** 1
Reported: 12/08/1997
Hospitalisation:
Onset Date: 31/12/1996
Outcome: Not yet recovered

Gender: F
Weight: 66.00
Age: 72Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lichenoid keratosis			Oral and topical steroids

Medicine Details:

DEPTRAN (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ZOCOR (Suspected) **Reason:** Othr&unspec metabolic diseases
10.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:** 31/01/1997

TRYPTANOL (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/11/1987 **Stopped:** 31/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				19/03/1997 microscopy: the epidermis shows moderate to marked hyperkeratosis patchy thinning and vacuola change of the basal layer. there is a slight saw- tooth configuration of the epidermis. a light, patchy lymphocytic infiltrate is present in the papilliary dermis & about upper dermal vessels. no eosinophils are seen.

Additional Information:

Rash responded quickly once zocor stopped. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 119814 **Seq:** 1
Reported: 12/08/1997
Hospitalisation:
Onset Date: 31/12/1996
Outcome: Not yet recovered

Gender: F
Weight: 66.00
Age: 72Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

DEPTRAN (Other drug)			**Reason:**	
	0.0			
Batch:	**Started:**			**Stopped:**
ZOCOR (Suspected)			**Reason:**	Othr&unspec metabolic diseases
	10.0 Milligram	Daily		
Batch:	**Started:** 01/01/1992			**Stopped:** 31/01/1997
TRYPTANOL (Suspected)			**Reason:**	
Tablet	25.0 Milligram	Daily		Oral
Batch:	**Started:** 01/11/1987			**Stopped:** 31/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Conclusion: right upper arm: the findings present are consistent with atrophic lechen planus. the pattern is less suggestive of a lichenoid drug eruption. in the absence of a deep dermal infiltrate, the pattern is less characteristic of lupus erythematosus

Additional Information:

Rash responded quickly once zocor stopped. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Report Detail

Case Number: 120019 **Seq:** 1 **Gender:** F
Reported: 18/08/1997 **Weight:**
Hospitalisation: **Age:** 78Y
Onset Date: 23/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Dehydration Dry mouth			

Medicine Details:

VALIUM (Other drug) **Reason:**
5.0 Milligram As necessary
Batch: **Started:** **Stopped:**

TEMAZE (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

MACRODANTIN (Other drug) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

MORPHINE MIXTURE, GOVERNMENT STORES (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
0.0 As necessary
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 120019 **Seq:** 1 **Gender:** F
Reported: 18/08/1997 **Weight:**
Hospitalisation: **Age:** 78Y
Onset Date: 23/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

MS CONTIN (Suspected)			**Reason:**	
	60.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
TRYPTANOL (Suspected)			**Reason:**	
	75.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 120069 **Seq:** 1 **Gender:** F
Reported: 20/08/1997 **Weight:** 84.00
Hospitalisation: **Age:** 53Y
Onset Date: 31/12/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oesophageal spasm		Esophageal spasm	
Aphasia		Incoherent	
Amnesia			
Coordination abnormal			
Hypertension			
Sleep disorder			
Tachycardia			

Medicine Details:

METOPROLOL TARTRATE (Other drug) **Reason:**
450.0 Milligram Daily
Batch: **Started:** 01/01/1977 **Stopped:**

CLONIDINE HYDROCHLORIDE (Other drug) **Reason:**
225.0 Microgram Daily
Batch: **Started:** 01/01/1975 **Stopped:**

TRYPTANOL (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 01/01/1977 **Stopped:** 31/01/1979

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 120081 **Seq:** 1 **Gender:** M
Reported: 20/08/1997 **Weight:**
Hospitalisation: **Age:** 7Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension Serotonin syndrome			

Medicine Details:

TRYPTANOL (Interaction) **Reason:**

Tablet 100.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

PROZAC (Interaction) **Reason:**

20.0 Milligram Daily

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reported in a published article (journal of paediatrics and child health, 1996 32(3), 265-266).

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 121290 **Seq:** 1 **Gender:** M
Reported: 02/10/1997 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia		Tardive dyskinesia lo limbs, buttocks,	

Medicine Details:

RITALIN (Other drug) **Reason:** Depression
0.0
Batch: **Started:** L TERM **Stopped:**

NAPROSYN (Other drug) **Reason:** Depression
0.0
Batch: **Started:** L TERM **Stopped:**

DIAZEPAM (Other drug) **Reason:** Anxiety neurosis
0.0 As necessary
Batch: **Started:** **Stopped:** 31/01/1994

TRYPTANOL (Suspected) **Reason:** Depression
0.0
Batch: **Started:** L TERM **Stopped:** 28/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Tardive dyskinesia of lower limbs, buttocks, abdominal muscles confirmed by director of neurology.

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Report Detail

Case Number: 121290 **Seq:** 1
Reported: 02/10/1997
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: M
Weight:
Age: 71Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/05/1996 **Stopped:** 28/05/1997

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 121581 **Seq:** 1 **Gender:** M
Reported: 14/10/1997 **Weight:** 75.00
Hospitalisation: **Age:** 83Y
Onset Date: 09/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Atrial fibrillation Pulmonary oedema			Endep ceased, cardioversion

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **Stopped:** 09/10/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 121747 **Seq:** 1 **Gender:** F
Reported: 21/10/1997 **Weight:** 50.00
Hospitalisation: **Age:** 47Y
Onset Date: 09/08/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Urticaria			

Medicine Details:

ENDONE (Other drug) **Reason:** Pain
15.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ENDEP (Suspected) **Reason:** Pain
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 07/08/1997 **Stopped:** 09/08/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 122428 **Seq:** 1
Reported: 18/11/1997
Hospitalisation: Hospitalisation prolonged
Onset Date: 30/10/1997
Outcome: Recovered

Gender: M
Weight:
Age: 68Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Somnolence			

Medicine Details:

MS CONTIN (Other drug)			**Reason:**	Pain	
Oral application	120.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	28/10/1997
MAXOLON (Other drug)			**Reason:**	Nausea and vomiting	
Oral application	40.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	05/11/1997
SERENACE (Other drug)			**Reason:**	Pain	
Oral application	3.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	05/11/1997
ZANTAC (Other drug)			**Reason:**		
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	28/10/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 122428 **Seq:** 1
Reported: 18/11/1997
Hospitalisation: Hospitalisation prolonged
Onset Date: 30/10/1997
Outcome: Recovered

Gender: M
Weight:
Age: 68Y
DOB:
Causality: Causality probable

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:** Pain

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** 28/10/1997 **Stopped:** 30/10/1997

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 122731 **Seq:** 1 **Gender:** M
Reported: 21/11/1997 **Weight:**
Hospitalisation: **Age:** 43Y
Onset Date: 13/11/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paralysis Dysarthria		Paralysis of the legs and arms. Slurred speech.	

Medicine Details:

ENDEP (Suspected) **Reason:** Headache
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 13/11/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has recently received a head injury. similar adverse reaction to zoloft.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 123788 **Seq:** 1 **Gender:** F
Reported: 06/01/1998 **Weight:** 65.00
Hospitalisation: **Age:** 22Y
Onset Date: 24/12/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuropathy peripheral		In left ulnar nerve distributor	

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 20/12/1997 **Stopped:** 29/12/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 123816 **Seq:** 1 **Gender:** F
Reported: 07/01/1998 **Weight:** 70.00
Hospitalisation: **Age:** 40Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Polydipsia Weight increased Hyperhidrosis	Severe	Drinking upto 4-8 litres of water/day Lost 4kg rapidly	Tryptanol ceased, therapy changed to another

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 21/11/1997 **Stopped:** 10/12/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 123878 **Seq:** 1 **Gender:** U
Reported: 08/12/1997 **Weight:**
Hospitalisation: **Age:** 39
Onset Date: 11/11/1997 **DOB:** 19/05/1958
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction Hyperhidrosis Malaise Paraesthesia		Redness of legs, sunburn	

Medicine Details:

PROZAC (Other drug) **Reason:**
Capsule 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ZITHROMAX (Suspected) **Reason:**
1.0 Gram 1 time
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 08/11/1997 **Stopped:** 04/12/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 124561 **Seq:** 1 **Gender:** M
Reported: 02/02/1998 **Weight:** 61.00
Hospitalisation: **Age:** 37Y
Onset Date: 31/05/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lipodystrophy acquired		Fat wasting on face	

Medicine Details:

TRIMETHOPRIM-SULFAMETHOXAZOLE (Suspected) **Reason:**
0.0
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

CRIXIVAN (Suspected) **Reason:** Hiv infection
Capsule 2.4 Gram Daily Oral
Batch: **Started:** 24/08/1995 **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Otr periph nerve dis not auton
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 01/10/1995 **Stopped:** CONTIN

STAVUDINE (Suspected) **Reason:**
80.0 Milligram Daily
Batch: **Started:** 27/08/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 124561 **Seq:** 1
Reported: 02/02/1998
Hospitalisation:
Onset Date: 31/05/1997
Outcome: Not yet recovered

Gender: M
Weight: 61.00
Age: 37Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

3TC (Suspected)		**Reason:**	
	450.0 Milligram	Daily	
Batch:	**Started:** 27/08/1996	**Stopped:**	CONTIN
ACICLOVIR (Suspected)		**Reason:**	
	600.0 Milligram	Daily	
Batch:	**Started:** 01/01/1994	**Stopped:**	CONTIN
FLUCONAZOLE (Suspected)		**Reason:**	
	100.0 Milligram	Daily	
Batch:	**Started:** 01/10/1995	**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 124609 **Seq:** 1 **Gender:** M
Reported: 02/02/1998 **Weight:** 61.00
Hospitalisation: **Age:** 37Y
Onset Date: 31/05/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lipodystrophy acquired		Fat wasting on pt's face.	

Medicine Details:

TRIMETHOPRIM-SULFAMETHOXAZOLE (Suspected) **Reason:**
0.0 Milligram
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

CRIXIVAN (Suspected) **Reason:** Hiv infection
Capsule 2.4 Gram Daily Oral
Batch: **Started:** 24/08/1995 **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr periph nerve dis not auton
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 01/10/1995 **Stopped:** CONTIN

STAVUDINE (Suspected) **Reason:**
80.0 Milligram Daily
Batch: **Started:** 27/08/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

In may 1997 fat wasting was noted on the patient's face (hollow cheeks) and 'relative' central obesity (ie no change in abdomen). no action was taken with regard to indinavir sulfate. the patient's weight pre-protease treatment was 60 kg and post-protease treatment is 61kg. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 124609 **Seq:** 1
Reported: 02/02/1998
Hospitalisation:
Onset Date: 31/05/1997
Outcome: Not yet recovered

Gender: M
Weight: 61.00
Age: 37Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

3TC (Suspected)		**Reason:**	
	450.0 Milligram	Daily	
Batch:	**Started:** 27/08/1996	**Stopped:**	CONTIN
ACICLOVIR (Suspected)		**Reason:**	
	600.0 Milligram	Daily	
Batch:	**Started:** 01/01/1994	**Stopped:**	CONTIN
FLUCONAZOLE (Suspected)		**Reason:**	
	100.0 Milligram	Daily	
Batch:	**Started:** 01/10/1995	**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

In may 1997 fat wasting was noted on the patient's face (hollow cheeks) and 'relative' central obesity (ie no change in abdomen). no action was taken with regard to indinavir sulfate. the patient's weight pre-protease treatment was 60 kg and post-protease treatment is 61kg. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred

Report Detail

Case Number: 124612 **Seq:** 1 **Gender:** F
Reported: 02/02/1998 **Weight:** 44.00
Hospitalisation: **Age:** 63Y
Onset Date: 14/10/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

PULMICORT (Other drug) **Reason:**
Inhalation 250.0 Milligram Daily Inhalation
Batch: **Started:** 10/05/1994 **Stopped:** CONTIN

OGEN (Other drug) **Reason:**
Tablet 1.2 Milligram Daily Oral
Batch: **Started:** 17/01/1995 **Stopped:** CONTIN

ATROVENT (Other drug) **Reason:**
Inhalation 20.0 Milligram Daily Inhalation
Batch: **Started:** 22/02/1996 **Stopped:** CONTIN

ISOPTIN SR (Other drug) **Reason:**
Tablet 180.0 Milligram Daily Oral
Batch: **Started:** 09/05/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				Ast/alt both greater than 3 x uln

Additional Information:

Report Detail

Case Number: 124612 **Seq:** 1 **Gender:** F
Reported: 02/02/1998 **Weight:** 44.00
Hospitalisation: **Age:** 63Y
Onset Date: 14/10/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

ATORVASTATIN (Suspected)			**Reason:** Othr&unspec metabolic diseases	
Oral application	20.0 Milligram	Daily	Oral	
Batch:	**Started:** 11/06/1997		**Stopped:**	CONTIN
ENDEP (Suspected)			**Reason:**	
Tablet	10.0 Milligram	Daily	Oral	
Batch:	**Started:** 26/05/1997		**Stopped:**	CONTIN
CALTRATE (Suspected)			**Reason:**	
Tablet	600.0 Milligram	Daily	Oral	
Batch:	**Started:** 15/09/1997		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 124762 **Seq:** 1 **Gender:** M
Reported: 09/02/1998 **Weight:** 80.00
Hospitalisation: **Age:** 44Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			Aropax ceased

Medicine Details:

DUCENE (Other drug) **Reason:** Anxiety neurosis
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AROPAX (Suspected) **Reason:** Depression
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 01/07/1997 **Stopped:** 12/12/1997

TRYPTANOL (Suspected) **Reason:** Specific disorders of sleep
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/05/1997 **Stopped:** 15/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				05/11/97 09/01/98 alt 149 68 ast 112 - ggt 734 430

Additional Information:

Past history: hepatitis a - many years. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 125800 **Seq:** 1 **Gender:** F
Reported: 13/03/1998 **Weight:**
Hospitalisation: **Age:** 59Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Palpitations Tremor Hyperhidrosis		Symptons akin to hypoglycaemia Trembling	 Endep ceased

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 3D

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 125801 **Seq:** 1 **Gender:** F
Reported: 13/03/1998 **Weight:**
Hospitalisation: **Age:** 59Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Palpitations		Symptons akin to hypoglycaemia	
Tremor		Trembling	Endep ceased
Hyperhidrosis			

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

After taking endep 25 for 3 nights pt changed to triptine 25 with no ill effects. when dosage increased to 50 mg nocte, the same effects were experienced with endep 50. pt diagnosed with chronic fatigue syndrome, attacks of hypoglycaemia and allergies to various substances including most colours and flavours. generic substitution problem

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 125879 **Seq:** 1
Reported: 17/03/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blood creatine phosphokinase in			
Confusional state			
Hepatic function abnormal			
Hypertonia			
Hypochloraemia			
Hypomagnesaemia			
Hyponatraemia			
Neuroleptic malignant syndrome			Treatment: valium, gentamicin and ciproxin
Pyrexia			

Medicine Details:

RISPERIDONE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 25/02/1998

TRYPTANOL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 25/02/1998

HALOPERIDOL (Suspected) **Reason:** Reactive confusion
2.0 Milligram Daily
Batch: **Started:** 25/02/1998 **Stopped:** 26/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	<30 U/L	13/03/1994	8	
	ALT = SGPT	<30 U/L	22/02/1998	30	
	ALT = SGPT	<30 U/L	02/03/1998	42	
	AST = SGOT	<30 U/L	13/03/1994	17	
	AST = SGOT	<30 U/L	22/02/1998	48	
	AST = SGOT	<30 U/L	02/03/1998	70	
	Bilirubin	<20 UMOL/L	13/03/1994	5	

Additional Information:

Patient was treated for uti. died on 08/03/98.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 125879 **Seq:** 1
Reported: 17/03/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

RISPERIDONE (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
TRYPTANOL (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
HALOPERIDOL (Suspected)		**Reason:**	Reactive confusion
	2.0 Milligram	Daily	
Batch:	**Started:** 25/02/1998		**Stopped:** 26/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	<20 UMOL/L	22/02/1998	27	
	Bilirubin	<20 UMOL/L	02/03/1998	12	
	Chloride	100 - 111	13/03/1994	92	
	Chloride	100 - 111	22/02/1998	85	
	Chloride	100 - 111	02/03/1998	122	
	Creatine	<200 U/L	22/02/1998	1190	
	Creatine	<200 U/L	02/03/1998	1159	

Additional Information:

Patient was treated for uti. died on 08/03/98.

Report Detail

Case Number: 125879 **Seq:** 1
Reported: 17/03/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

RISPERIDONE (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
TRYPTANOL (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
HALOPERIDOL (Suspected)		**Reason:**	Reactive confusion
	2.0 Milligram	Daily	
Batch:	**Started:** 25/02/1998		**Stopped:** 26/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine [illegible]	<200 U/L	04/03/1998	1875	
	Creatine [illegible]	<200 U/L	01/03/1998	831	
	GGT = SGGT = [illegible]	<35 U/L	13/03/1994	25	
	GGT = SGGT = [illegible]	<35 U/L	02/03/1998	57	
	GGT = SGGT = [illegible]	<35 U/L	22/02/1998	65	
	LDH	<400 U/L	22/02/1998	652	
	LDH	<400 U/L	04/03/1998	677	

Additional Information:

Patient was treated for uti. died on 08/03/98.

Report Detail

Case Number: 125879 **Seq:** 1
Reported: 17/03/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: F
Weight:
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

RISPERIDONE (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
TRYPTANOL (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:** 25/02/1998
HALOPERIDOL (Suspected)		**Reason:**	Reactive confusion
	2.0 Milligram	Daily	
Batch:	**Started:** 25/02/1998		**Stopped:** 26/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Magnesium	0.70 - 1.00 MMOL/L	22/02/1998	0.63	
	SAP = ALP	35 - 120 U/L	13/03/1994	127	
	SAP = ALP	35 - 120 U/L	02/03/1998	68	
	SAP = ALP	35 - 120 U/L	22/02/1998	87	
	Sodium	136 - 145 MMOL/L	13/03/1994	134	
	Sodium	136 - 145 MMOL/L	22/02/1998	124	
	Sodium	136 - 145 MMOL/L	02/03/1998	156	

Additional Information:

Patient was treated for uti. died on 08/03/98.

Report Detail

Case Number: 126003 **Seq:** 1 **Gender:** F
Reported: 19/03/1998 **Weight:**
Hospitalisation: **Age:** 30
Onset Date: 07/02/1998 **DOB:** 01/12/1967
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Depression Migraine Rash		Feet, lower legs and hands feeling puffy.	Ceased panadeine forte for 2 days, swelling & rash subsided.re-introduced 09/02/98

Medicine Details:

INDOMETHACIN (Suspected) **Reason:** Migraine
Suppository 0.0 As necessary Rectal
Batch: **Started:** 30/01/1998 **Stopped:** CONTIN

STEMETIL (Suspected) **Reason:** Migraine
Suppository 0.0 As necessary Rectal
Batch: **Started:** 30/01/1998 **Stopped:** CONTIN

PANADEINE FORTE (Suspected) **Reason:** Migraine
Tablet 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 30/01/1998 **Stopped:** 03/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 126003 **Seq:** 1
Reported: 19/03/1998
Hospitalisation:
Onset Date: 07/02/1998
Outcome: Not yet recovered

Gender: F
Weight:
Age: 30
DOB: 01/12/1967
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** 03/02/1998 **Stopped:** CONTIN

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 126677 **Seq:** 1 **Gender:** F
Reported: 14/04/1998 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperthyroidism			Tryptanol ceased

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 126696 **Seq:** 1 **Gender:** M
Reported: 14/04/1998 **Weight:** 95.00
Hospitalisation: **Age:** 51Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention			Tryptanol ceased.

Medicine Details:

SULFASALAZINE (Other drug) **Reason:**
3.0 Gram Daily
Batch: **Started:** **Stopped:** CONTIN

ATENOLOL (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 01/01/1971 **Stopped:** 31/01/1991

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/01/1991 **Stopped:** 31/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				In 1988, ivp which showed a huge bladder without backpressure hydronephrosis. in1994 cystoscopy and urethroscopy showed some narrowing of the patient's bladder outlet but insufficient to cause large chronic retention.

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 126872 **Seq:** 1 **Gender:** M
Reported: 20/04/1998 **Weight:** 42.00
Hospitalisation: **Age:** 77Y
Onset Date: 05/02/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysarthria Confusional state Somnolence Urinary retention		Slurred speach.	Minipress, tryptanol ceased.

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** 03/02/1998 **Stopped:** 06/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 128085 **Seq:** 1
Reported: 05/06/1998
Hospitalisation:
Onset Date: 29/05/1998
Outcome: Not yet recovered

Gender: M
Weight: 92.00
Age: 27Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Renal impairment			Ceased likely drugs

Medicine Details:

GENTAMICIN SULPHATE (Suspected) **Reason:** Prophylaxis
Injection 240.0 Milligram 2 times Intravenous
Batch: **Started:** 28/05/1998 **Stopped:** 29/05/1998

DICLOXACILLIN SODIUM (Suspected) **Reason:** Prophylaxis
4.0 Gram Daily
Batch: **Started:** 28/05/1998 **Stopped:** 29/05/1998

INDOCID (Suspected) **Reason:** Other disturbance of sensation
Suppository 200.0 Milligram Daily Rectal
Batch: **Started:** **Stopped:** 01/06/1998

FRAGMIN (Suspected) **Reason:** Prophylaxis
Injection 5.0 Thousand Internat Daily Subcutaneous
Batch: **Started:** 29/05/1998 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatinine		29/05/1998	0.12	
	Creatinine		29/05/1998	0.17	
	Creatinine		30/05/1998	0.29	
	Creatinine		31/05/1998	0.31	
	Creatinine		01/06/1998	0.34	
	Creatinine		02/06/1998	0.36	

Additional Information:

Report Detail

Case Number: 128085 **Seq:** 1
Reported: 05/06/1998
Hospitalisation:
Onset Date: 29/05/1998
Outcome: Not yet recovered

Gender: M
Weight: 92.00
Age: 27Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Tablet 35.0 Milligram Daily Oral

Batch: **Started:** 26/05/1998 **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 128334 **Seq:** 1 **Gender:** F
Reported: 12/06/1998 **Weight:** 60.00
Hospitalisation: **Age:** 79
Onset Date: 01/05/1998 **DOB:** 08/10/1918
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Hallucinations at night	Ceased tryptanol and ms contin

Medicine Details:

WARFARIN SODIUM (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

LISINOPRIL (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRANSIDERM-NITRO (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

LANOXIN PG (Other drug) **Reason:**
3.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 128334 **Seq:** 1 **Gender:** F
Reported: 12/06/1998 **Weight:** 60.00
Hospitalisation: **Age:** 79
Onset Date: 01/05/1998 **DOB:** 08/10/1918
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

LASIX (Other drug)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	
PANADOL (Other drug)			**Reason:**		
	4.0	Gram	Daily		
Batch:	**Started:**			**Stopped:**	
TRYPTANOL (Suspected)			**Reason:**	Pain	
	10.0	Milligram	Daily		
Batch:	**Started:**	01/05/1998		**Stopped:**	05/05/1998
MS CONTIN (Suspected)			**Reason:**	Pain	
	10.0	Milligram	Daily		
Batch:	**Started:**	01/05/1998		**Stopped:**	05/05/1998

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 128510 **Seq:** 1
Reported: 18/06/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 73
DOB: 16/08/1924
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state		Confusion and falls.	Amitriptyline ceased

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Pain
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

PROZAC (Interaction) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

MS CONTIN (Other drug) **Reason:** Pain
90.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

KAPANOL (Other drug) **Reason:** Pain
100.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 128510 **Seq:** 1
Reported: 18/06/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 73
DOB: 16/08/1924
Causality: Causality possible

Reaction Detail

Medicine Details:

EPILIM (Other drug)		**Reason:**	Pain	
	1.0 Gram	Daily		
Batch:	**Started:**		**Stopped:**	CONTIN
TEGRETOL (Other drug)		**Reason:**	Other&unspecified epilepsy	
	200.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	CONTIN
TEMAZEPAM (Other drug)		**Reason:**	Specific disorders of sleep	
	30.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	CONTIN
DUPHALAC (Other drug)		**Reason:**	Constipation	
	30.0 Millilitre	Daily		
Batch:	**Started:**		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 128782 **Seq:** 1 **Gender:** M
Reported: 24/06/1998 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 10/06/1998 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation	Severe		
Urinary tract infection	Severe		Amitrptyline, thioridazine and benzotropine ceased.

Medicine Details:

RISPERDAL (Other drug) **Reason:**
0.0
Batch: **Started:** 01/04/1998 **Stopped:** 10/06/1998

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 29/05/1998

THIORIDAZINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 29/05/1998

BENZTROPINE MESYLATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 29/05/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 128939 **Seq:** 1 **Gender:** F
Reported: 06/07/1998 **Weight:**
Hospitalisation: **Age:** 70Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia	Required Specialist Consultation	Mild alopecia	

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 129149 **Seq:** 1
Reported: 13/07/1998
Hospitalisation: Admitted to hospital
Onset Date: 28/06/1998
Outcome: Recovered

Gender: F
Weight:
Age: 87Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation Dizziness			 Ceased medicaiton

Medicine Details:

ASPIRIN (Other drug) **Reason:**
0.5 Dose Unspecified Daily
Batch: **Started:** **Stopped:** CONTIN

GLYCERYL TRINITRATE (Suspected) **Reason:**
Ointment 0.0 Daily Topical
Batch: **Started:** **Stopped:** 28/06/1998

MODURETIC (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** 28/06/1998

TRYPTANOL (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:** 28/06/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 130635 **Seq:** 1 **Gender:** F
Reported: 31/08/1998 **Weight:**
Hospitalisation: **Age:** 77Y
Onset Date: 31/03/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Abdominal pain Drug ineffective		Huge loose bowel motions	Iv fluids, iv antibiotics and inotropic infusions.

Medicine Details:

ADRENALINE (Interaction) **Reason:**
Injection, intravenous infusion 0.0 Intravenous
Batch: **Started:** **Stopped:**

ENDEP (Interaction) **Reason:**
200.0 Milligram Daily
Batch: **Started:** **Stopped:**

MELLERIL (Interaction) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See seq 01 for other drugs. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 130773 **Seq:** 1
Reported: 02/09/1998
Hospitalisation:
Onset Date: 14/08/1998
Outcome: Not yet recovered

Gender: F
Weight:
Age: 42
DOB: 21/12/1955
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pain Headache Thirst		Dull pain left side of body	

Medicine Details:

ENDEP (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ALDACTONE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MICROGYNON (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ZYRTEC (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 130773 **Seq:** 1
Reported: 02/09/1998
Hospitalisation:
Onset Date: 14/08/1998
Outcome: Not yet recovered

Gender: F
Weight:
Age: 42
DOB: 21/12/1955
Causality: Causality possible

Reaction Detail

Medicine Details:

LITHICARB (Suspected)			**Reason:** Depression	
Tablet	500.0 Milligram	Daily	Oral	
Batch:	**Started:** 14/08/1998		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 131352 **Seq:** 1 **Gender:** F
Reported: 17/09/1998 **Weight:**
Hospitalisation: **Age:**
Onset Date: 02/07/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			Calamine lotion

Medicine Details:

TEGRETOL (Suspected) **Reason:**
400.0 Milligram Daily
Batch: **Started:** **Stopped:** 5W

PRO-BANTHINE (Suspected) **Reason:**
30.0 Milligram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 133235 **Seq:** 1
Reported: 18/11/1998
Hospitalisation: Admitted to hospital
Onset Date: 25/10/1998
Outcome: Unknown

Gender: M
Weight:
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension			Ceased deptran and endep.

Medicine Details:

COLOXYL (Other drug) **Reason:** Constipation
50.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

CAPADEX (Other drug) **Reason:** Pain
0.0 As necessary
Batch: **Started:** **Stopped:**

QUINSUL (Other drug) **Reason:** Pain in limb
300.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ASPIRIN (Other drug) **Reason:** Chron isch heart dis no hyper
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 133235 **Seq:** 1
Reported: 18/11/1998
Hospitalisation: Admitted to hospital
Onset Date: 25/10/1998
Outcome: Unknown

Gender: M
Weight:
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMFAMOX (Other drug)			**Reason:**	
	80.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
ENDEP (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**			**Stopped:**
DEPTRAN (Suspected)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 133878 **Seq:** 1 **Gender:** M
Reported: 03/12/1998 **Weight:**
Hospitalisation: **Age:** 47Y
Onset Date: 05/09/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia			Amitriptyline discontinued.

Medicine Details:

BISALAX (Other drug) **Reason:** Constipation
Suppository 1.0 Dose Unspecified Daily Rectal
Batch: **Started:** 11/06/1998 **Stopped:** CONTIN

COLOXYL WITH SENNA (Other drug) **Reason:** Other diseases of bone
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 26/05/1998 **Stopped:** CONTIN

BACLOFEN (Other drug) **Reason:** Abnormal involuntary movement
Tablet 80.0 Milligram Daily Oral
Batch: **Started:** 01/06/1998 **Stopped:** CONTIN

DANTROLENE SODIUM (Other drug) **Reason:** Abnormal involuntary movement
Capsule 150.0 Milligram Daily Oral
Batch: **Started:** 26/06/1998 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No previous adverse reation. patient commenced on mexilitine increaseing dose 100mg tds.

Report Detail

Case Number: 133878 **Seq:** 1 **Gender:** M
Reported: 03/12/1998 **Weight:**
Hospitalisation: **Age:** 47Y
Onset Date: 05/09/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
100.0 Milligram Daily
Batch: **Started:** 26/05/1998 **Stopped:** 05/09/1998

Laboratory Investigations:

Additional Information:

No previous adverse reation. patient commenced on mexilitine increaseing dose 100mg tds.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 134253 **Seq:** 1
Reported: 14/12/1998
Hospitalisation: Hospitalisation prolonged
Onset Date: 27/11/1998
Outcome: Recovered

Gender: M
Weight:
Age: 54Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arrhythmia			

Medicine Details:

PERICYAZINE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

PANADEINE (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** 23/11/1998 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient underwent electroconvulsive therapy in hospital and on 27 nov 98 after the 4th electroconvulsive treatment, the patient developed arrhythmia which lasted for four hours.

Report Detail

Case Number: 134289 **Seq:** 1 **Gender:** F
Reported: 14/12/1998 **Weight:**
Hospitalisation: **Age:** 64Y
Onset Date: 10/11/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion			

Medicine Details:

MS CONTIN (Other drug) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 05/11/1998

WARFARIN SODIUM (Suspected) **Reason:**
Tablet 3.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

CIPROFLOXACIN (Suspected) **Reason:**
Tablet 1.5 Gram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 134289 **Seq:** 1
Reported: 14/12/1998
Hospitalisation:
Onset Date: 10/11/1998
Outcome: Not yet recovered

Gender: F
Weight:
Age: 64Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

DEXAMETHASONE (Suspected) **Reason:**

Tablet 2.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 135295 **Seq:** 1 **Gender:** F
Reported: 25/01/1999 **Weight:** 56.00
Hospitalisation: **Age:** 22Y
Onset Date: 18/12/1998 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state		Increased confusion, disorientated	
Acidosis		Methylmalonic acidosis	Zantac, amitryptylene, & cogentin ceased.

Medicine Details:

POTASSIUM CHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** 18/12/1998 **Stopped:** CONTIN

ONDANSETRON HYDROCHLORIDE DIHYDRATE (Suspected) **Reason:**
8.0 Milligram As necessary
Batch: **Started:** 18/12/1998 **Stopped:** CONTIN

PARACETAMOL (Suspected) **Reason:**
500.0 Milligram As necessary
Batch: **Started:** 18/12/1998 **Stopped:**

ZANTAC (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 20/12/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Odansetron, vit. c, thiamine, carnitine, sodium nitrate, & paracetamol were all administered via gastrostomy. potassium chloride was added to feeds and the dose was 124mmol.

Report Detail

Case Number: 135295 **Seq:** 1 **Gender:** F
Reported: 25/01/1999 **Weight:** 56.00
Hospitalisation: **Age:** 22Y
Onset Date: 18/12/1998 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)		**Reason:**		
	75.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	20/12/1998
COGENTIN (Suspected)		**Reason:**		
	6.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	20/12/1998
ASCORBIC ACID (Suspected)		**Reason:**		
	500.0 Milligram	Daily		
Batch:	**Started:** 18/12/1998		**Stopped:**	CONTIN
THIAMINE (Suspected)		**Reason:**		
	100.0 Milligram	Daily		
Batch:	**Started:** 18/12/1998		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

Odansetron, vit. c, thiamine, carnitine, sodium nitrate, & paracetamol were all administered via gastrostomy. potassium chloride was added to feeds and the dose was 124mmol.

Report Detail

Case Number: 135295 **Seq:** 1 **Gender:** F
Reported: 25/01/1999 **Weight:** 56.00
Hospitalisation: **Age:** 22Y
Onset Date: 18/12/1998 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

CARNITINE (Suspected)		**Reason:**		
	9.9 Millilitre	Daily		
Batch:	**Started:** 18/12/1998		**Stopped:**	CONTIN
SODIUM CITRATE (Suspected)		**Reason:**		
	96.0 Millilitre	Daily		
Batch:	**Started:** 18/12/1998		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

Odansetron, vit. c, thiamine, carnitine, sodium nitrate, & paracetamol were all administered via gastrostomy. potassium chloride was added to feeds and the dose was 124mmol.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 135467 **Seq:** 1 **Gender:** F
Reported: 29/01/1999 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain		Burning sensation in chest.	
Depersonalisation		Feeling strange.	
Hypertonia		Stiffness in limbs.	
Agitation			Was given nitrazepam.
Hyperreflexia			
Nausea			

Medicine Details:

TRYPTANOL (Interaction) **Reason:** Depression
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

THIORIDAZINE HYDROCHLORIDE (Interaction) **Reason:** Unspecified psychosis
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

NEFAZODONE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 0.0 Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Medical journal of australia 1998, 169(10), 523-525. no washout period. heart rate of 70 bpm and blood pressure of 112/62 mmhg.

Report Detail

Case Number: 136232 **Seq:** 1 **Gender:** U
Reported: 15/02/1999 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 04/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Micturition disorder		Great dificulty urinating	
Dry mouth		Severe dry mouth	
Diplopia			
Hyperhidrosis			

Medicine Details:

MS CONTIN (Other drug) **Reason:** Pain
120.0 Milligram Daily
Batch: **Started:** 29/01/1999 **Stopped:** 08/02/1999

ENDONE (Other drug) **Reason:** Pain
5.0 Milligram Daily
Batch: **Started:** 01/01/1999 **Stopped:**

ASPIRIN (Other drug) **Reason:** Systemic lupus erythematosus
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1999 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Pain
25.0 Milligram Daily
Batch: **Started:** 04/02/1999 **Stopped:** 08/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has sle with arthritis. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 136559 **Seq:** 1 **Gender:** M
Reported: 22/02/1999 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: 14/01/1999 **DOB:**
Outcome: Recovered with sequelae **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion		Grand mal fits	

Medicine Details:

TRAMAL (Interaction) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 19/11/1998 **Stopped:** 14/01/1999

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Tramal commenced at 50mg tds and gradually increased to 150mg tds. experienced reactions and recovered with some short-term memory loss. no history of fitting.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 136591 **Seq:** 1 **Gender:** F
Reported: 25/02/1999 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 13/02/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Hypoaesthesia			

Medicine Details:

CLONAZEPAM (Other drug) **Reason:**
1.0 Gram Daily
Batch: **Started:** **Stopped:**

MS CONTIN (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

NAPROXEN (Other drug) **Reason:**
1.0 Gram Daily
Batch: **Started:** **Stopped:** 13/02/1999

TRYPTANOL (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Right hip & knee pain. mri scan l5/s1 disc bulging. nil allergies. pt started on tryptanol 25mg and increased to 50mg nocte.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 136815 **Seq:** 1 **Gender:** F
Reported: 26/02/1999 **Weight:**
Hospitalisation: **Age:** 75Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			Zoloft was discontinued and patient restarted the

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Depression

0.0

Batch: **Started:** 01/01/1968 **Stopped:** 30Y

ZOLOFT (Interaction) **Reason:** Depression

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 136860 **Seq:** 1 **Gender:** M
Reported: 01/03/1999 **Weight:** 0.00
Hospitalisation: **Age:** 99U
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast neoplasm			

Medicine Details:

TRYPTANOL (Suspected) **Reason:**
0.0 Oral
Batch: **Started:** 01/04/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 137525 **Seq:** 1
Reported: 18/03/1999
Hospitalisation: Hospitalisation prolonged
Onset Date: 27/11/1998
Outcome: Recovered

Gender: M
Weight:
Age: 54Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arrhythmia Tachycardia		Arrhythmia lasted 4 hours	

Medicine Details:

TRYPTINE (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** 24/11/1998 **Stopped:** CONTIN

NEULACTIL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

PANADEINE FORTE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Ect 27/11/98 and subsequently.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 138518 **Seq:** 1 **Gender:** F
Reported: 19/04/1999 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 10/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Rash after 8 days of vancomycin.	

Medicine Details:

VANCOMYCIN HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

GENTAMICIN SULPHATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

NEVIRAPINE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

3TC (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drug details - omeprazole, flumirazine, frusamide & quinine.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 138518 **Seq:** 1 **Gender:** F
Reported: 19/04/1999 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 10/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CHLORPROMAZINE HYDROCHLORIDE (Suspected)	**Reason:**	
0.0		
Batch:	**Started:**	**Stopped:**
AMITRIPTYLINE HYDROCHLORIDE (Suspected)	**Reason:**	
0.0		
Batch:	**Started:**	**Stopped:**

Laboratory Investigations:

Additional Information:

See original report for other drug details - omeprazole, flumirazine, frusamide & quinine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 138646 **Seq:** 1 **Gender:** F
Reported: 22/04/1999 **Weight:** 46.00
Hospitalisation: **Age:** 79
Onset Date: 14/09/1998 **DOB:** 22/04/1919
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Hypokalaemia Hyponatraemia Rhabdomyolysis		24 hours lying on the ground	Iv fluids - ceased indapamide

Medicine Details:

BRICANYL (Suspected) **Reason:** Asthma
Inhalation 0.0 Inhalation
Batch: **Started:** **Stopped:** 14/09/1998

INDAPAMIDE HEMIHYDRATE (Suspected) **Reason:**
2.5 Milligram Daily
Batch: **Started:** **Stopped:** 14/09/1998

ALENDRONATE SODIUM (Suspected) **Reason:** Other diseases of bone
10.0 Milligram Daily
Batch: **Started:** **Stopped:** 14/09/1998

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 14/09/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium		14/09/1998	3	
	Sodium		14/09/1998	125	

Additional Information:

See original report for details of other drugs - captopril, pulmicort. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 138646 **Seq:** 1
Reported: 22/04/1999
Hospitalisation:
Onset Date: 14/09/1998
Outcome: Recovered

Gender: F
Weight: 46.00
Age: 79
DOB: 22/04/1919
Causality: Causality possible

Reaction Detail

Medicine Details:

FERRO-GRADUMET (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1995 **Stopped:** 14/09/1998

Laboratory Investigations:

Additional Information:

See original report for details of other drugs - captopril, pulmicort. the dosage start date is not accurate but indicates that start occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 139194 **Seq:** 1 **Gender:** F
Reported: 03/05/1999 **Weight:** 79.00
Hospitalisation: **Age:** 53Y
Onset Date: 16/04/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle twitching Tremor Vomiting		Twitchy eye	 Nil

Medicine Details:

VOLTAREN (Suspected) **Reason:** Sciatica

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 15/04/1999 **Stopped:** 26/04/1999

TRYPTANOL (Suspected) **Reason:**

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** 15/04/1999 **Stopped:** 26/04/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date of recovery: 27/4/99. patient has hiatus hernia. see original report for details of other drugs - mersyndol forte, normison.

Report Detail

Case Number: 140292 **Seq:** 1 **Gender:** F
Reported: 07/06/1999 **Weight:**
Hospitalisation: **Age:** 83Y
Onset Date: 17/05/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			Diprosone cream

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
10.0 Milligram As necessary
Batch: **Started:** 14/05/1999 **Stopped:** 17/05/1999

LANSOPRAZOLE (Suspected) **Reason:** Other diseases of esophagus
60.0 Milligram Daily
Batch: **Started:** 01/04/1999 **Stopped:** 25/05/1999

PROPRANOLOL HYDROCHLORIDE (Suspected) **Reason:** Abnormal involuntary movement
80.0 Milligram Daily
Batch: **Started:** 01/04/1999 **Stopped:** 25/05/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 140777 **Seq:** 1 **Gender:** M
Reported: 22/06/1999 **Weight:** 80.00
Hospitalisation: **Age:** 35Y
Onset Date: 31/07/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia			Had to have surgical reduction.

Medicine Details:

ENDEP (Suspected) **Reason:** Specific disorders of sleep

0.0

Batch: **Started:** 01/07/1998 **Stopped:** 28/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drug details - murelax. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Report Detail

Case Number: 141388 **Seq:** 1
Reported: 12/07/1999
Hospitalisation:
Onset Date: 24/05/1999
Outcome: Not yet recovered

Gender: M
Weight: 70.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			Treated with hydrocortisone

Medicine Details:

CYCLOSPORIN-A (Suspected) **Reason:**
200.0 Milligram Daily
Batch: **Started:** 01/03/1999 **Stopped:** 08/06/1999

PROGOUT (Suspected) **Reason:** Gout
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:** 30/06/1999

CORDARONE X (Suspected) **Reason:** Chron isch heart dis no hyper
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **Stopped:** 30/06/1999

ENDEP (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 01/03/1999 **Stopped:** 30/06/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Skin biopsy:- suggestive of drug induced eruption

Additional Information:

Please see original report for details of other drugs - digoxin, hypnodorm. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 141601 **Seq:** 1
Reported: 15/07/1999
Hospitalisation: Hospitalisation prolonged
Onset Date:
Outcome: Recovered with sequelae

Gender: F
Weight:
Age: 81Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension		Bp decreased from 140/90 mmhg to 100/60	Tryptanol was discontinued

Medicine Details:

PARACETAMOL (Other drug) **Reason:**
1.0 Gram As necessary
Batch: **Started:** **Stopped:** CONTIN

CISAPRIDE (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 01/01/1999 **Stopped:** 28/07/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: patient fell, sustaining a further thoracic vertebral (osteoporotic) compression. patient's hospitalisation was prolonged because of reaction. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143143 **Seq:** 1 **Gender:** M
Reported: 27/08/1999 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/12/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		Decreased visual acuity	Tryptanol was discontinued

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
Tablet 0.0 Oral
Batch: **Started:** **Stopped:** 31/01/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143144 **Seq:** 1 **Gender:** M
Reported: 27/08/1999 **Weight:** 75.00
Hospitalisation: **Age:** 63Y
Onset Date: 23/07/1999 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gout			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis

Tablet 25.0 Milligram As necessary Oral

Batch: **Started:** 16/07/1999 **Stopped:** 18/07/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient does not take tryptanol very often, but when he does he develops gout. patient also taking amlodipine besylate and simvastatin.

Report Detail

Case Number: 143309 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 19/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperthyroidism			Ceased aratac. given prednisolone, and

Medicine Details:

ARATAC (Suspected) **Reason:**
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 23/02/1999

PRAVACHOL (Suspected) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 25/02/1999

TRYPTANOL (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 24/02/1999

CARDIZEM (Suspected) **Reason:**
Tablet 240.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 22/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Thyroid		19/02/1999	0.13	
	Thyroid		24/02/1999	<0.03	
	Thyroid		25/02/1999	<0.03	
	Thyroid		14/03/1999	0.54	
	Thyroid		09/03/1999	0.23	
	Thyroid		05/03/1999	0.04	
	Thyroxine		19/02/1999	20.4	

Additional Information:

See original report for details of other drugs: transiderm nitro, zantac, ventolin, becotide, lasix, marevan.

Report Detail

Case Number: 143309 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 19/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

SLOW-K (Suspected) **Reason:**
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 23/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Thyroxine		24/02/1999	36.4	
	Thyroxine		25/02/1999	27.8	
	Thyroxine		05/03/1999	18.8	
	Thyroxine		09/03/1999	18.5	
	Thyroxine		14/03/1999	16.7	
	Tri-iodo thyronine		19/02/1999	-	
	Tri-iodo thyronine		24/02/1999	4.1	

Additional Information:

See original report for details of other drugs: transiderm nitro, zantac, ventolin, becotide, lasix, marevan.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143309 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 19/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ARATAC (Suspected)			**Reason:**		
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	23/02/1999
PRAVACHOL (Suspected)			**Reason:**		
Tablet	10.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	25/02/1999
TRYPTANOL (Suspected)			**Reason:**		
Tablet	25.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	24/02/1999
CARDIZEM (Suspected)			**Reason:**		
Tablet	240.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	22/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Tri-iodo thyronine		25/02/1999	4.4	
	Tri-iodo thyronine		05/03/1999	3	
	Tri-iodo thyronine		09/03/1999	-	
	Tri-iodo thyronine		14/03/1999	-	

Additional Information:

See original report for details of other drugs: transiderm nitro, zantac, ventolin, becotide, lasix, marevan.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143309 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 19/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

SLOW-K (Suspected)			**Reason:**		
Tablet	3.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**			**Stopped:**	23/02/1999

Laboratory Investigations:

Additional Information:

See original report for details of other drugs: transiderm nitro, zantac, ventolin, becotide, lasix, marevan.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143311 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 24/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatocellular damage Prothrombin level decreased			

Medicine Details:

ARATAC (Suspected) **Reason:**
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 23/02/1999

PRAVACHOL (Suspected) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 25/02/1999

TRYPTANOL (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 24/02/1999

CARDIZEM (Suspected) **Reason:**
Tablet 240.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 22/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		24/02/1999	394	
	ALT = SGPT		26/02/1999	407	
	ALT = SGPT		02/03/1999	187	
	ALT = SGPT		08/03/1999	134	
	AST = SGOT		24/02/1999	-	
	AST = SGOT		08/03/1999	75	
	AST = SGOT		26/02/1999	217	

Additional Information:

See original report for details on other drugs: transiderm nitro, zantac, ventolin, becotide, lasix.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 143311 **Seq:** 1 **Gender:** F
Reported: 31/08/1999 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 24/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

SLOW-K (Suspected) **Reason:**
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 23/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT		02/03/1999	82	
	International [illegible]		24/02/1999	3.3	
	LDH		24/02/1999	-	
	LDH		26/02/1999	1295	
	LDH		02/03/1999	1010	
	LDH		08/03/1999	705	

Additional Information:

See original report for details on other drugs: transiderm nitro, zantac, ventolin, becotide, lasix.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 144205 **Seq:** 1 **Gender:** M
Reported: 29/09/1999 **Weight:**
Hospitalisation: **Age:** 37
Onset Date: 31/12/1999 **DOB:** 02/01/1962
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Dizziness Leukopenia Malaise			

Medicine Details:

CLOZARIL (Suspected) **Reason:** Unspecified schizophrenia
Tablet 0.0 Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	White blood cells				Decreasing white blood cell count from 6.0 - 4.0.

Additional Information:

- the date of onset is not accurate but indicates that onset occurred sometime during the year.

Report Detail

Case Number: 144448 **Seq:** 1 **Gender:** M
Reported: 07/10/1999 **Weight:**
Hospitalisation: **Age:** 66Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia Dry mouth Tongue ulceration		Bad taste	

Medicine Details:

PRINIVIL (Suspected) **Reason:**
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 01/01/1997 **Stopped:** CONTIN

LOSEC (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:** CONTIN

AVAPRO (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 01/01/1997 **Stopped:** CONTIN

ZOCOR (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/01/1997 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the

Report Detail

Case Number: 144448 **Seq:** 1
Reported: 07/10/1999
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: M
Weight:
Age: 66Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TAMBOCOR (Suspected)		**Reason:**		
	50.0 Milligram	Daily		
Batch:	**Started:** 01/01/1997		**Stopped:**	CONTIN
ZYLOPRIM (Suspected)		**Reason:**		
	300.0 Milligram	Daily		
Batch:	**Started:** 01/01/1997		**Stopped:**	CONTIN
ENDEP (Suspected)		**Reason:**		
	75.0 Milligram	Daily		
Batch:	**Started:** 01/01/1998		**Stopped:**	CONTIN
WARFARIN SODIUM (Suspected)		**Reason:**		
	3.0 Milligram	Daily		
Batch:	**Started:** 01/01/1997		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 144448 **Seq:** 1
Reported: 07/10/1999
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: M
Weight:
Age: 66Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

BURINEX (Suspected) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 01/01/1997 **Stopped:** CONTIN

Laboratory Investigations:

Additional Information:

the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 145620 **Seq:** 1
Reported: 05/11/1999
Hospitalisation:
Onset Date: 31/12/1998
Outcome: Not yet recovered

Gender: F
Weight: 80.00
Age: A6
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus		Buzzing in ears	
Headache		Pulsing in head	

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 01/01/1990 **Stopped:** 31/01/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for details of other drugs; atenolol, felodipine, fluvast- atin, nizatidine. for over 12 months patient has experienced symptoms with tryptanol, subsequently in 1999 therapy with tryptanol was interrupted for four days and although symptoms did not resolve, it did lessen. due to worsening symptoms of depression, patient restarted therapy with tryptanol and symptoms increased and still persist. - the date of onset is not accurate but indicates that onset occurred sometime during the year. the dosage start date is not accurate but indicates that start occurred sometime during the year. the dosage stop date is not accurate but indicates that stoppage occurred sometime during the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 147521 **Seq:** 1 **Gender:** M
Reported: 24/12/1999 **Weight:**
Hospitalisation: **Age:** 18Y
Onset Date: 04/11/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		Patient expereinced blurred vision.	Decreased night dose from 75mg to 50mg.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Tablet 125.0 Milligram Daily Oral

Batch: **Started:** 04/10/1999 **Stopped:** 04/11/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking coloxyl with senna, fludrocortisone, naproxen, enemax, baclofen, dantrolene, agarol and diazepam.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 148138 **Seq:** 1 **Gender:** M
Reported: 19/01/2000 **Weight:** 75.00
Hospitalisation: **Age:** 27Y
Onset Date: 13/01/2000 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia		Pt experienced stiffness, weakness & pain	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 09/01/2000 **Stopped:** 15/01/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 149035 **Seq:** 1 **Gender:** U
Reported: 11/02/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuropathy peripheral		Patient experienced peripheral neuropathy	

Medicine Details:

TASMAR (Suspected) **Reason:**
Oral application 0.0 Oral
Batch: **Started:** **Stopped:** 1M

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 0.0 Oral
Batch: **Started:** **Stopped:**

QUININE (Suspected) **Reason:**
Oral application 0.0 Oral
Batch: **Started:** **Stopped:**

PERGOLIDE (Suspected) **Reason:**
Oral application 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was treated with tasmar for one month before it was discontinued from the market.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 149411 **Seq:** 1 **Gender:** F
Reported: 18/02/2000 **Weight:**
Hospitalisation: **Age:** 52Y
Onset Date: 22/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension		Bp 100/60	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Migraine
25.0 Milligram Daily
Batch: **Started:** 18/02/1999 **Stopped:** 22/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 149657 **Seq:** 1
Reported: 25/02/2000
Hospitalisation: Admitted to hospital
Onset Date: 12/02/2000
Outcome: Not yet recovered

Gender: F
Weight:
Age: 47
DOB: 18/01/1953
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Peptic ulcer perforation	Severe Life		 Perforation oversewn.

Medicine Details:

INDOCID (Suspected) **Reason:** Unspecified osteomyelitis
Suppository 1.0 Dose Unspecified As necessary Rectal
Batch: **Started:** 28/04/1999 **Stopped:** 13/02/2000

ENDEP (Suspected) **Reason:** Unspecified osteomyelitis
20.0 Milligram Daily
Batch: **Started:** 12/01/2000 **Stopped:** 13/02/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking valium.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 151733 **Seq:** 1 **Gender:** M
Reported: 13/03/2000 **Weight:**
Hospitalisation: **Age:** 30
Onset Date: 04/09/1997 **DOB:** 11/11/1966
Outcome: Death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arrhythmia Sudden death			

Medicine Details:

BECONASE (Other drug) **Reason:**
Spray 0.0 Nasal
Batch: **Started:** 03/09/1997 **Stopped:**

BENZTROPINE MESYLATE (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 03/09/1997 **Stopped:**

OMEPRAZOLE (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

CLOZAPINE (Suspected) **Reason:**
350.0 Milligram 1 time
Batch: **Started:** 03/09/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level Serum drug level Serum drug level				Blood taken at the post mortem examination revealed the following:- - clozapine (1.3mg/l) - codeine___(0.27mg/l) - morphine - metabolite of codeine (0.03mg/l) - amitriptyline (0.18mg/l) Continued:- - nortriptyline - metabolite of amitriptyline (0.19mg/l) - trichloroethanol - metabolite of chloral hydrate (11mg/l) - benztropine (nil) - clonazepam (nil) Continued:- - 7-amino clonazepam - metabolite of clonazepam (nil) - valproic acid (44mg/l) - alcohol (nil)

Additional Information:

Date of death 4/9/1997. cause of death: "undetermined. unexpected death associated with high levels of antipsychotic medication and analgesia".

Report Detail

Case Number: 151733 **Seq:** 1 **Gender:** M
Reported: 13/03/2000 **Weight:**
Hospitalisation: **Age:** 30
Onset Date: 04/09/1997 **DOB:** 11/11/1966
Outcome: Death **Causality:** Causality possible

Reaction Detail

Medicine Details:

CLOZARIL (Suspected)			**Reason:**	Unspecified schizophrenia	
Tablet	500.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	04/09/1997
SODIUM VALPROATE (Suspected)			**Reason:**		
	1.8	Gram	Daily		
Batch:	**Started:**	03/09/1997		**Stopped:**	
CHLORAL HYDRATE (Suspected)			**Reason:**		
	1.0	Gram	Daily		
Batch:	**Started:**	03/09/1997		**Stopped:**	
CEPHALEXIN (Suspected)			**Reason:**		
	500.0	Milligram	Daily		
Batch:	**Started:**	03/09/1997		**Stopped:**	

Laboratory Investigations:

Additional Information:

Date of death 4/9/1997. cause of death: "undetermined. unexpected death associated with high levels of antipsychotic medication and analgesia".

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 151733 **Seq:** 1 **Gender:** M
Reported: 13/03/2000 **Weight:**
Hospitalisation: **Age:** 30
Onset Date: 04/09/1997 **DOB:** 11/11/1966
Outcome: Death **Causality:** Causality possible

Reaction Detail

Medicine Details:

PANADEINE FORTE (Suspected)		**Reason:**	
	60.0 Milligram	Daily	
Batch:	**Started:** 03/09/1997		**Stopped:**
CLONAZEPAM (Suspected)		**Reason:**	
	2.0 Milligram	Daily	
Batch:	**Started:**		**Stopped:**
AMITRIPTYLINE HYDROCHLORIDE (Suspected)		**Reason:**	
	50.0 Milligram	Daily	
Batch:	**Started:** 03/09/1997		**Stopped:**

Laboratory Investigations:

Additional Information:

Date of death 4/9/1997. cause of death: "undetermined. unexpected death associated with high levels of antipsychotic medication and analgesia".

Report Detail

Case Number: 151778 **Seq:** 1 **Gender:** M
Reported: 17/04/2000 **Weight:**
Hospitalisation: **Age:** 39
Onset Date: 21/03/2000 **DOB:** 14/10/1960
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cardiomyopathy		Hypertrophic obstructive cardiomyopathy	

Medicine Details:

CLOZARIL (Suspected) **Reason:** Unspecified schizophrenia
Tablet 500.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

DIAZEPAM (Suspected) **Reason:**
2.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 153097 **Seq:** 1
Reported: 25/05/2000
Hospitalisation: Admitted to hospital
Onset Date: 07/05/2000
Outcome: Recovered

Gender: F
Weight:
Age: 83Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing		Blanching on legs	
Rash		Butterfly rash over face	
Speech disorder		Unclear speech	
Confusional state			Amitriptyline and orudis ceased. date of recovery 17/05/2000

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 04/05/2000 **Stopped:** 07/05/2000

ORUDIS SR (Suspected) **Reason:** Unspecified arthritis
100.0 Milligram Daily
Batch: **Started:** 24/04/2000 **Stopped:** 07/05/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Valproate was reduced over 6 days prior to commencement of amitriptylline. prior to admission patient had mild dementia but was alert and able to clearly express herself.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 154639 **Seq:** 1
Reported: 12/07/2000
Hospitalisation:
Onset Date: 27/07/1999
Outcome: Not yet recovered

Gender: F
Weight: 74.00
Age: 50
DOB: 13/08/1948
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Salivary hypersecretion			Required a visit to doctor.

Medicine Details:

CLOZAPINE (Suspected) **Reason:** Unspecified psychosis
0.0
Batch: **Started:** 23/06/1999 **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** 09/07/1999 **Stopped:** 31/08/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 154645 **Seq:** 1
Reported: 12/07/2000
Hospitalisation:
Onset Date: 15/06/1999
Outcome: Not yet recovered

Gender: M
Weight: 85.00
Age: 58
DOB: 26/05/1941
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Salivary hypersecretion			Benztropine and terazosin. required a visit to doctor.

Medicine Details:

CLOZAPINE (Suspected) **Reason:** Schizophrenia
10.0 Milligram Daily
Batch: **Started:** 15/06/1999 **Stopped:** 23/06/1999

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Schizophrenia
25.0 Milligram Daily
Batch: **Started:** 15/06/1999 **Stopped:** 18/06/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 154969 **Seq:** 1 **Gender:** F
Reported: 19/07/2000 **Weight:**
Hospitalisation: **Age:** 75Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lacrimal disorder Constipation Dry mouth Somnolence Tremor	Severe	Reduced lacrimal flow Severe constipation.	Increased intake of fibre and

Medicine Details:

XANAX (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

TEMAZEPAM (Suspected) **Reason:** Depression
10.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Her constipation is so severe that manual evacuation of the bowel has been required.

Report Detail

Case Number: 156301 **Seq:** 1 **Gender:** F
Reported: 14/08/2000 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 01/08/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular		Rash on back and abdomen.	Diprosone cream. progout and urex ceased.

Medicine Details:

PROGOUT (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 25/07/2000 **Stopped:** 02/08/2000

UREX (Suspected) **Reason:** Edema and dropsy
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 01/08/2000 **Stopped:** 02/08/2000

MAXIPIME (Suspected) **Reason:** Agranulocytosis
Injection 4.0 Gram Daily Intravenous
Batch: **Started:** 25/07/2000 **Stopped:**

TOBRAMYCIN (Suspected) **Reason:** Agranulocytosis
Injection 240.0 Milligram Daily Intravenous
Batch: **Started:** 25/07/2000 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 156301 **Seq:** 1 **Gender:** F
Reported: 14/08/2000 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 01/08/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FLUCONAZOLE (Suspected)			**Reason:** Prophylaxis	
Oral application	200.0	Milligram	Daily	Oral
Batch:	**Started:** 25/07/2000			**Stopped:**
ENDEP (Suspected)			**Reason:**	
Tablet	10.0	Milligram	Daily	Oral
Batch:	**Started:** 27/07/2000			**Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 156991 **Seq:** 1 **Gender:** M
Reported: 25/08/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: 24/06/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			Ceased celebrex

Medicine Details:

CELEBREX (Suspected) **Reason:** Herpes zoster
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** 13/06/2000 **Stopped:** 16/06/2000

TRYPTANOL (Suspected) **Reason:** Herpes zoster
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 13/06/2000 **Stopped:** 27/06/2000

TEGRETOL (Suspected) **Reason:** Herpes zoster
Tablet 500.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 1M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient took celebrex for 4 days then ceased. eight days after ceasing celebrex patient developed symptoms.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 158045 **Seq:** 1 **Gender:** F
Reported: 25/09/2000 **Weight:** 58.00
Hospitalisation: **Age:** 20Y
Onset Date: 31/03/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blood creatine phosphokinase in		Mildly elevated ck	
Rash		Rash on feet, legs, hands.	
Alopecia			
Hyperhidrosis			
Mouth ulceration			
Myalgia			
Pyrexia			
Red blood cell sedimentation rate			
Systemic lupus erythematosus			Prednisolone 50mg/d. date of recovery 28/07/00
Weight decreased			

Medicine Details:

PANADEINE (Suspected) **Reason:** Pain
Oral application 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

MINOCYCLINE HYDROCHLORIDE (Suspected) **Reason:** Other acne of sebaceous glands
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 21/07/2000

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Anxiety neurosis
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 19/07/2000

MONOFEME 28 (Suspected) **Reason:** Contraception
Oral application 0.0 Daily Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 158223 **Seq:** 1 **Gender:** F
Reported: 03/10/2000 **Weight:** 47.00
Hospitalisation: **Age:** 56
Onset Date: 10/09/2000 **DOB:** 08/10/1943
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety		Increased anxiety state.	
Paraesthesia		Increased sensitivity to nerve endings.	
Drug ineffective			
Hyperkinesia			
Hypokinesia			

Medicine Details:

ENDEP (Suspected) **Reason:** Abnormal involuntary movement
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 08/09/2000 **Stopped:** 09/09/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking primolut n. one 25mg tablet did nothing to decrease pain of back spasm, increased to 2 tablets and brought on adr.

Report Detail

Case Number: 159597 **Seq:** 1 **Gender:** F
Reported: 29/11/2000 **Weight:**
Hospitalisation: **Age:** 59
Onset Date: 01/04/2000 **DOB:** 28/07/1940
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Deafness Gait disturbance Osteoarthritis Visual disturbance			Dose reduction of gabapentin to 300mg bd and amitriptyline to 50mg. ceased baclofen.

Medicine Details:

GABAPENTIN (Suspected) **Reason:**
900.0 Milligram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

BACLOFEN (Suspected) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 159826 **Seq:** 1 **Gender:** F
Reported: 11/12/2000 **Weight:** 63.00
Hospitalisation: **Age:** 57
Onset Date: 04/12/2000 **DOB:** 21/05/1943
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Stomatitis Rash Pruritus		Mucosal swelling. Pruritic skin rash, mucosal swelling	Required a visit to doctor.

Medicine Details:

ENDEP (Suspected) **Reason:**
30.0 Milligram Daily
Batch: **Started:** 29/11/2000 **Stopped:** 01/12/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 159922 **Seq:** 1 **Gender:** F
Reported: 11/12/2000 **Weight:**
Hospitalisation: **Age:** 83
Onset Date: 22/08/2000 **DOB:** 05/03/1917
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gastrointestinal haemorrhage			Prbc. warfarin ceased. start iron tablets.

Medicine Details:

ISORDIL (Suspected) **Reason:** Angina pectoris w/o hyperten
0.0 As necessary
Batch: **Started:** **Stopped:** 22/08/2000

WARFARIN SODIUM (Suspected) **Reason:** Phleb&thrombophleb lowr extrem
Oral application 3.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 22/08/2000

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 23/08/2000

IMDUR (Suspected) **Reason:** Angina pectoris w/o hyperten
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 22/08/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	International		22/08/2000	2.8	

Additional Information:

Patient also taking femara, prednisolone, coloxyl.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 159922 **Seq:** 1 **Gender:** F
Reported: 11/12/2000 **Weight:**
Hospitalisation: **Age:** 83
Onset Date: 22/08/2000 **DOB:** 05/03/1917
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ENALAPRIL MALEATE (Suspected) **Reason:**

Oral application	10.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:** 24/08/2000

Laboratory Investigations:

Additional Information:

Patient also taking femara, prednisolone, coloxyl.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 160197 **Seq:** 1
Reported: 22/12/2000
Hospitalisation:
Onset Date: 14/12/2000
Outcome: Not yet recovered

Gender: M
Weight: 80.00
Age: 28
DOB: 08/05/1972
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Rhabdomyolysis	Life	Life threatening reaction	Pt hospitalised icu--> ventilation

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CLOMIPRAMINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

TRAMAL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MERSYNDOL (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 160848 **Seq:** 1 **Gender:** M
Reported: 22/01/2001 **Weight:**
Hospitalisation: **Age:** 70
Onset Date: 05/01/2001 **DOB:** 02/01/1931
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pollakiuria		Increased urinary frequency - every half	Required a visit to doctor. ceased endep.
Urinary incontinence		Increased urinary leakage.	
Micturition disorder		Increased urinary urgency.	

Medicine Details:

LOSEC (Other drug) **Reason:** Ulcer of stomach w/o ment perf
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 01/01/1980 **Stopped:** CONTIN

ENDEP (Suspected) **Reason:** Migraine
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 24/11/2000 **Stopped:** 11/01/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Cystoscopy 30/11/2000. nil known allergies. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 161069 **Seq:** 1 **Gender:** F
Reported: 30/01/2001 **Weight:**
Hospitalisation: **Age:** 48
Onset Date: 21/01/2001 **DOB:** 18/10/1952
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia			

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 17/01/2001 **Stopped:** CONTIN

NEURONTIN (Suspected) **Reason:** Pain
Capsule 900.0 Milligram Daily Oral
Batch: **Started:** 31/12/2000 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 161330 **Seq:** 1
Reported: 12/02/2001
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: U
Weight: 100.00
Age: 31
DOB: 31/07/1969
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking - lithicarb, panadeine forte, panamax, voltaren and xanax.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 162180 **Seq:** 1
Reported: 05/03/2001
Hospitalisation: Admitted to hospital
Onset Date: 25/02/2001
Outcome: Recovered

Gender: F
Weight:
Age: 28
DOB: 26/02/1972
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome			Patient hospitalised. date of recovery 28/02/01.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ZYBAN SR (Interaction) **Reason:** Smoker
2.0 Dose Unspecified Daily
Batch: **Started:** 20/02/2001 **Stopped:** 25/02/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 162384 **Seq:** 1 **Gender:** F
Reported: 08/03/2001 **Weight:** 38.00
Hospitalisation: **Age:**
Onset Date: 11/01/2001 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Anorexia Confusional state Dry mouth Fatigue Hypertension Neuropathy peripheral		Involuntary jerking movements.	

Medicine Details:

BACLOFEN (Suspected) **Reason:** Abnormal involuntary movement
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** 07/01/2001 **Stopped:** 12/01/2001

QUININE (Suspected) **Reason:** Pain in limb
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 10/01/2001 **Stopped:** 11/01/2001

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** 07/01/2001 **Stopped:** 10/01/2001

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** 10/01/2001 **Stopped:** 11/01/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking trimethoprim, digesic, heparin.

Report Detail

Case Number: 162384 **Seq:** 1
Reported: 08/03/2001
Hospitalisation:
Onset Date: 11/01/2001
Outcome: Not yet recovered

Gender: F
Weight: 38.00
Age:
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

PERSANTIN SR (Suspected)			**Reason:**		
Capsule	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	11/01/2001
RISPERIDONE (Suspected)			**Reason:**	Abdominal pain	
	1.0	Milligram	Daily		
Batch:	**Started:**	10/01/2001		**Stopped:**	11/01/2001

Laboratory Investigations:

Additional Information:
Patient was also taking trimethoprim, digesic, heparin.

Report Detail

Case Number: 162593 **Seq:** 1 **Gender:** F
Reported: 13/03/2001 **Weight:**
Hospitalisation: **Age:** 62
Onset Date: 02/03/2001 **DOB:** 11/05/1938
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Insomnia			Stop endep. date of recovery: 3/3/2001.

Medicine Details:

ENDEP (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 26/02/2001 **Stopped:** 02/03/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking lipitor, caltrate, cartia, gopten and dapatabs. patient has allergy to iodine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 162604 **Seq:** 1 **Gender:** M
Reported: 13/03/2001 **Weight:** 65.00
Hospitalisation: **Age:** 83Y
Onset Date: 23/02/2001 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder		Involuntary muscle movements of arms	Endep ceased. date of recovery: 26/2/2001.

Medicine Details:

ENDEP (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient experienced adr for two successive nights. patient also taking androcur, cartia, temaze, zoladex, lasix and lanoxin pg.

Report Detail

Case Number: 163550 **Seq:** 1
Reported: 06/04/2001
Hospitalisation: Admitted to hospital
Onset Date: 20/02/2001
Outcome: Recovered

Gender: F
Weight: 45.00
Age: 35
DOB: 13/08/1965
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion		Grand mal seizure	Hospitalised

Medicine Details:

ENDEP (Interaction) **Reason:** Specific disorders of sleep
25.0 Milligram As necessary
Batch: **Started:** **Stopped:**

ZYBAN SR (Interaction) **Reason:** Smoker
300.0 Milligram Daily
Batch: **Started:** 14/02/2001 **Stopped:** 20/02/2001

EFEXOR-XR (Interaction) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** 16/01/2001 **Stopped:** 11/02/2001

METHADONE HYDROCHLORIDE (Other drug) **Reason:** Drug dependence
15.0 Millilitre Daily
Batch: **Started:** L TERM **Stopped:** 17/02/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient history of seizures on benzodiazepine withdrawal but with no eeg abnormalities at the time.

Report Detail

Case Number: 163727 **Seq:** 1
Reported: 19/04/2001
Hospitalisation:
Onset Date: 30/03/2001
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 43
DOB: 03/10/1957
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertonia		Involuntary spasmodic contraction,	Indep ceased. no change in spasms. restarted. required a

Medicine Details:

CAFERGOT (Other drug) **Reason:**
2.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:** CONTIN

THYROXINE SODIUM (Other drug) **Reason:**
150.0 Microgram Daily
Batch: **Started:** **Stopped:** CONTIN

ENDEP (Suspected) **Reason:** Migraine
25.0 Milligram Daily
Batch: **Started:** 01/01/2000 L TERM **Stopped:** 30/03/2001 CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Relevant medical history - hypothyroidism since 1992, migraine since 1994, no hypertension, no operations, no allergies, no other gastrointestinal symptoms. the dose start date for the drug endep is not necessarily accurate but indicates the drug was started sometime in the month. endep therapy was restarted on 06/04/2001.

Report Detail

Case Number: 163757 **Seq:** 1
Reported: 17/04/2001
Hospitalisation: Admitted to hospital
Onset Date: 05/04/2001
Outcome: Recovered with sequelae

Gender: F
Weight: 60.00
Age: 57
DOB: 09/03/1944
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion			Dilantin load. zyban and tramal ceased. patient hospitalised.

Medicine Details:

ZYBAN SR (Interaction) **Reason:** Smoker
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 17/03/2001 **Stopped:** 05/04/2001

TRAMAL (Interaction) **Reason:** Pain
600.0 Milligram Daily
Batch: **Started:** 02/04/2001 **Stopped:** 05/04/2001

METROGYL (Interaction) **Reason:** Diarrhea
Tablet 1.2 Gram Daily Oral
Batch: **Started:** 29/03/2001 **Stopped:** 08/04/2001

ENDEP (Interaction) **Reason:** Specific disorders of sleep
25.0 Milligram Daily
Batch: **Started:** 31/01/2001 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: broken colar bone (left clavicle).

Report Detail

Case Number: 163757 **Seq:** 1
Reported: 17/04/2001
Hospitalisation: Admitted to hospital
Onset Date: 05/04/2001
Outcome: Recovered with sequelae

Gender: F
Weight: 60.00
Age: 57
DOB: 09/03/1944
Causality: Causality possible

Reaction Detail

Medicine Details:

OROXINE (Other drug)			**Reason:**	Myxedema	
	0.2	Microgram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
BLACKMORES ACIDOPHILUS BIFIDUS (Other drug)			**Reason:**		
	3.0	Dose Unspecified	Daily		
Batch:	**Started:**	29/03/2001		**Stopped:**	09/04/2001

Laboratory Investigations:

Additional Information:

Sequelae: broken colar bone (left clavicle).

Report Detail

Case Number: 165061 **Seq:** 1
Reported: 01/06/2001
Hospitalisation: Admitted to hospital
Onset Date: 15/05/2001
Outcome: Not yet recovered

Gender: F
Weight: 48.00
Age: 50
DOB: 08/02/1951
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haemorrhage intracranial Syncope			Life threatening reaction. patient hospitalised

Medicine Details:

ENDEP (Interaction) **Reason:** Anorexia
4.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1986 **Stopped:**

ZYBAN SR (Interaction) **Reason:** Smoker
1.0 Dose Unspecified Daily
Batch: **Started:** 09/05/2001 **Stopped:** 15/05/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				15/05/01 - ct scan showed intracranial haemorrhage.

Additional Information:

Sequelae: neurological deficit. the dose start date for endep is not accurate but indicates the drug was start ed sometime in the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 165563 **Seq:** 1 **Gender:** F
Reported: 14/06/2001 **Weight:** 57.00
Hospitalisation: **Age:** 52Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Anhidrosis			
Anxiety			
Confusional state			
Micturition disorder			

Medicine Details:

TRYPTANOL (Suspected) **Reason:** Anxiety neurosis
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/06/2001 **Stopped:** 04/06/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recovered 6/6/01.

Report Detail

Case Number: 165593 **Seq:** 1 **Gender:** M
Reported: 20/06/2001 **Weight:** 90.00
Hospitalisation: **Age:** 48Y
Onset Date: 20/06/2001 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Salivary hypersecretion Gastrooesophageal reflux diseas		Waterbrash.	

Medicine Details:

ZYBAN SR (Suspected) **Reason:** Smoker
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/05/2001 S TERM **Stopped:**

VIOXX (Suspected) **Reason:** Lumbalgia
Tablet 12.5 Milligram Daily Oral
Batch: **Started:** 17/05/2001 **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/01/2000 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The date of onset is not necessarily accurate but indicates onset sometime in the month. the dose start date for zyban is not necessarily accurate but indicates the drug was started sometime in the month. the dose start date for amitriptyline is not accurate but indicates the drug was started sometime in the year. zyban was tolerated on 1 tab daily but had reaction when taken twice daily.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 166528 **Seq:** 1 **Gender:** M
Reported: 17/07/2001 **Weight:**
Hospitalisation: **Age:** 81
Onset Date: 05/04/2001 **DOB:** 19/05/1919
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gait disturbance Orthostatic hypotension			

Medicine Details:

COLOXYL WITH SENNA (Other drug) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

RANITIDINE (Suspected) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1999 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/01/1999 **Stopped:** 05/04/2001

ASASANTIN SR (Suspected) **Reason:**
Capsule 2.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dose start date is not necessarily accurate but inicates ranitidine and amitriptyline were started sometime in the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 166528 **Seq:** 1
Reported: 17/07/2001
Hospitalisation:
Onset Date: 05/04/2001
Outcome: Not yet recovered

Gender: M
Weight:
Age: 81
DOB: 19/05/1919
Causality: Causality possible

Reaction Detail

Medicine Details:

NITRAZEPAM (Suspected) **Reason:**

Oral application 5.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 05/04/2001

Laboratory Investigations:

Additional Information:

The dose start date is not necessarily accurate but inicates ranitidine and amitriptyline were started sometime in the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 167243 **Seq:** 1 **Gender:** F
Reported: 06/08/2001 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cardiomyopathy			

Medicine Details:

GABAPENTIN (Suspected) **Reason:** Pain
Oral application 1.6 Gram Daily Oral
Batch: **Started:** 01/06/2001 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ASPIRIN (Suspected) **Reason:**
Oral application 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

IMURAN (Suspected) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Echo on 3 july 2001 revealed left ventricular effusion fraction (lvef) of 30%, mitral regurgitation moderate level.

Additional Information:

The dose start date for gabapentin is not necessarily accurate but indicates the drug was started sometime in the month. patient has a history of abdominal pain secondary gastroparesis and a pethidine/narcotic dependancy.

Report Detail

Case Number: 167243 **Seq:** 1 **Gender:** F
Reported: 06/08/2001 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

LASIX (Suspected)			**Reason:**		
Tablet	40.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
MAGNESIUM ASPARTATE (Suspected)			**Reason:**		
Oral application	1.5	Gram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
METHADONE HYDROCHLORIDE (Suspected)			**Reason:**		
Oral Liquid	50.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
PETHIDINE HYDROCHLORIDE (Suspected)			**Reason:**		
	150.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

The dose start date for gabapentin is not necessarily accurate but indicates the drug was started sometime in the month. patient has a history of abdominal pain secondary gastroparesis and a pethidine/narcotic dependancy.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 167243 **Seq:** 1 **Gender:** F
Reported: 06/08/2001 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

PHENERGAN (Suspected)			**Reason:**		
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
ZOFRAN (Suspected)			**Reason:**		
Tablet	8.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

The dose start date for gabapentin is not necessarily accurate but indicates the drug was started sometime in the month. patient has a history of abdominal pain secondary gastroparesis and a pethidine/narcotic dependancy.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 167753 **Seq:** 1
Reported: 21/08/2001
Hospitalisation:
Onset Date: 06/08/2001
Outcome: Recovered

Gender: F
Weight: 60.00
Age: 82
DOB: 24/07/1919
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abnormal dreams		Vivid abnormal dreams	Required a visit to doctor and ceased amitriptyline

Medicine Details:

OMEPRAZOLE (Other drug) **Reason:** Peptic ulcer nos w/o ment perf
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AMLODIPINE (Other drug) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

METOPROLOL TARTRATE (Other drug) **Reason:** Essential benign hypertension
200.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

CLOPIDOGREL (Other drug) **Reason:** Acute,ill-def cerebrovascular disease w/o hyperte
75.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 167753 **Seq:** 1 **Gender:** F
Reported: 21/08/2001 **Weight:** 60.00
Hospitalisation: **Age:** 82
Onset Date: 06/08/2001 **DOB:** 24/07/1919
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

SIMVASTATIN (Other drug) **Reason:** Othr&unspec metabolic diseases
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Nervousness
10.0 Milligram Daily
Batch: **Started:** 01/08/2001 **Stopped:** 18/08/2001

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 167966 **Seq:** 1 **Gender:** F
Reported: 30/08/2001 **Weight:** 55.00
Hospitalisation: **Age:** 39
Onset Date: 23/07/2001 **DOB:** 04/04/1962
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation Agitation Anxiety		Dissociation. Panic attacks.	Date of recovery 19.8.01. required a visit to doctor.

Medicine Details:

SEREPAX (Other drug) **Reason:** Anxiety neurosis
0.2 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

CELEBREX (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
200.0 Milligram Daily
Batch: **Started:** 20/07/2001 **Stopped:** 13/08/2001

TRYPTANOL (Suspected) **Reason:**
75.0 Milligram Daily
Batch: **Started:** 20/07/2001 **Stopped:** 13/08/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 168122 **Seq:** 1 **Gender:** F
Reported: 05/09/2001 **Weight:** 47.00
Hospitalisation: **Age:** 35Y
Onset Date: 10/08/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Arthralgia Conjunctivitis Purpura		Generalised pruritic rash	

Medicine Details:

SLOW-K (Other drug) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

LASIX (Other drug) **Reason:**
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

FERRO-GRADUMET (Other drug) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 08/08/2001 **Stopped:** 10/08/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 168122 **Seq:** 1 **Gender:** F
Reported: 05/09/2001 **Weight:** 47.00
Hospitalisation: **Age:** 35Y
Onset Date: 10/08/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

NEURONTIN (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Oral application 600.0 Milligram Daily Oral
Batch: **Started:** 08/08/2001 **Stopped:** 10/08/2001

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 170287 **Seq:** 1
Reported: 26/11/2001
Hospitalisation: Hospitalisation prolonged
Onset Date: 11/08/2001
Outcome: Recovered

Gender: F
Weight:
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium Hyponatraemia			Patient had a prolonged stay in hospital, treated with potassium, restricted fluids. ceased indapamide, amitriptyline and sertraline.

Medicine Details:

ASPIRIN (Other drug)			**Reason:**		
Tablet	100.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
SIMVASTATIN (Other drug)			**Reason:**		
Tablet	10.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
IRBESARTAN (Other drug)			**Reason:**		
Tablet	150.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
RANITIDINE (Other drug)			**Reason:**		
Tablet	300.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium				Sodium = 125

Additional Information:

Report Detail

Case Number: 170287 **Seq:** 1
Reported: 26/11/2001
Hospitalisation: Hospitalisation prolonged
Onset Date: 11/08/2001
Outcome: Recovered

Gender: F
Weight:
Age: 78Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

VERAPAMIL HYDROCHLORIDE (Other drug)				**Reason:**		
Tablet	80.0	Milligram	Daily		Oral	
Batch:	**Started:**			**Stopped:**		CONTIN
INDAPAMIDE HEMIHYDRATE (Suspected)				**Reason:**		
Tablet	2.5	Milligram	Daily		Oral	
Batch:	**Started:**	L TERM		**Stopped:**	10/08/2001	
SERTRALINE HYDROCHLORIDE (Suspected)				**Reason:**	Depression	
Tablet	50.0	Milligram	Daily		Oral	
Batch:	**Started:**	28/07/2001		**Stopped:**	04/08/2001	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)				**Reason:**	Depression	
Tablet	2.0	Dose Unspecified	Daily		Oral	
Batch:	**Started:**	04/08/2001		**Stopped:**	11/08/2001	

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 170300 **Seq:** 1
Reported: 26/11/2001
Hospitalisation:
Onset Date: 01/11/2001
Outcome: Recovered

Gender: M
Weight: 121.00
Age: 64
DOB: 19/11/1936
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tongue oedema		Swollen tongue	

Medicine Details:

DIGOXIN (Other drug) **Reason:** Otr&nos disord of heart rhythm
1.0 Dose Unspecified Daily
Batch: **Started:** 05/02/1997 **Stopped:**

JEZIL (Other drug) **Reason:** Othr&unspec metabolic diseases
120.0 Milligram Daily
Batch: **Started:** 20/12/2000 **Stopped:**

METFORMIN HYDROCHLORIDE (Other drug) **Reason:** Diabetes mellitus
32.0 Unit Daily
Batch: **Started:** 03/09/1997 **Stopped:**

DIAMICRON (Other drug) **Reason:** Diabetes mellitus
30.0 Milligram Daily
Batch: **Started:** 10/12/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recovered 2/11/01.

Report Detail

Case Number: 170300 **Seq:** 1
Reported: 26/11/2001
Hospitalisation:
Onset Date: 01/11/2001
Outcome: Recovered

Gender: M
Weight: 121.00
Age: 64
DOB: 19/11/1936
Causality: Causality possible

Reaction Detail

Medicine Details:

CELEBREX (Other drug)			**Reason:**	Other diseases of bone
	200.0	Milligram	Daily	
Batch:	**Started:**	05/09/2000		**Stopped:**
PLAVIX (Suspected)			**Reason:**	Chron isch heart dis no hyper
Tablet	75.0	Milligram	Daily	Oral
Batch:	**Started:**	24/10/2001		**Stopped:**
ASASANTIN SR (Suspected)			**Reason:**	Transient cereb isch no hypert
Capsule	2.0	Dose Unspecified	Daily	Oral
Batch:	**Started:**	27/09/2001		**Stopped:**
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**	30/05/2001		**Stopped:**

Laboratory Investigations:

Additional Information:
Patient recovered 2/11/01.

Report Detail

Case Number: 170300 **Seq:** 1 **Gender:** M
Reported: 26/11/2001 **Weight:** 121.00
Hospitalisation: **Age:** 64
Onset Date: 01/11/2001 **DOB:** 19/11/1936
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TENORMIN (Suspected)			**Reason:**	Chron isch heart dis no hyper
Tablet	25.0 Milligram		Daily	Oral
Batch:	**Started:** 24/10/2001			**Stopped:**
COVERSYL (Suspected)			**Reason:**	Essential benign hypertension
	8.0 Milligram		Daily	
Batch:	**Started:** 05/01/2001			**Stopped:**

Laboratory Investigations:

Additional Information:
Patient recovered 2/11/01.

Report Detail

Case Number: 170546 **Seq:** 1 **Gender:** U
Reported: 29/11/2001 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/07/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state		Confused and vague.	Amitriptyline ceased.
Aphasia		Slurred speech.	
Delusion		Voices and sounds.	
Hallucination			

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram 3 times
Batch: **Started:** 31/07/2001 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 18/07/2001 **Stopped:** 31/07/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Non-hodgkins lymphoma with leptomeningeal disease.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 170605 **Seq:** 1
Reported: 29/11/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: M
Weight: 90.00
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain			

Medicine Details:

VENTOLIN (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 0.0 As necessary Inhalation
Batch: **Started:** L TERM **Stopped:**

EUHYPNOS (Other drug) **Reason:**
10.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

SEREVENT (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 2.0 Dose Unspecified Daily Inhalation
Batch: **Started:** L TERM **Stopped:**

SOLONE (Suspected) **Reason:** Otr respiratory systm diseases
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 10/06/2001 **Stopped:** 13/06/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient suffering with coad. the dose start date is not necessarily accurate but indicates endep was started sometime in the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 170605 **Seq:** 1
Reported: 29/11/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: M
Weight: 90.00
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:**

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** 01/12/2000 **Stopped:** 13/06/2001

Laboratory Investigations:

Additional Information:

Patient suffering with coad. the dose start date is not necessarily accurate but indicates endep was started sometime in the month.

Report Detail

Case Number: 170616 **Seq:** 1
Reported: 30/11/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 25Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia Psychotic disorder Salivary hypersecretion			Patient was hospitalised and treated with amitriptyline. it was decided to gradually reduce and ceased valproate. amitriptyline was discontinued.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CLOZAPINE (Suspected) **Reason:** Schizophrenia
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

SODIUM VALPROATE (Suspected) **Reason:**
800.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Neutrophils White blood cells				Nc 3.9 x 10?9/l, nc 1.8 x 10?9/l. Wbc 6.2 x 10?9/l, wbc 3.3 x 10?9/l.

Additional Information:

Journal: australian & new zealand journal of psychiatry author: pantelis c, adresanya a. tittle: increased risk of neutropenia and agranulocytosis with sodium valproat used adjunctively with clozapine. volume: 35 (4) year:2001 pages:544-545.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 170841 **Seq:** 1
Reported: 06/12/2001
Hospitalisation: Admitted to hospital
Onset Date: 27/11/2001
Outcome: Recovered

Gender: F
Weight:
Age: 70
DOB: 21/11/1931
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Delirium			Patient was hospitalised.

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** **Stopped:** 27/11/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Uti also diagnosed and thought to be a contributing factor in precipitating confusion. patient recovered 2/12/01.

Report Detail

Case Number: 170842 **Seq:** 1
Reported: 06/12/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 70
DOB: 21/11/1931
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Delirium			Patient was hospitalised.

Medicine Details:

ENDEP (Suspected) **Reason:**
0.0
Batch: **Started:** 29/11/2001 **Stopped:** 30/11/2001

TRAMAL (Suspected) **Reason:** Pain
Tablet 50.0 Milligram As necessary Oral
Batch: **Started:** **Stopped:** 27/11/2001

VIOXX (Suspected) **Reason:** Unspecified arthritis
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 27/11/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recovered 2/12/01.

Report Detail

Case Number: 172512 **Seq:** 1 **Gender:** M
Reported: 07/02/2002 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: 29/12/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence Vomiting			Treated with vancomycin 500mg iv daily, meropenem 500mg iv q8h and somac 40mg iv daily

Medicine Details:

BAXTER TPN SOLUTION (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

TRAMADOL HYDROCHLORIDE (Suspected) **Reason:**
50.0 Milligram As necessary
Batch: **Started:** **Stopped:** 29/12/2001

SUCRALFATE (Suspected) **Reason:**
Tablet 4.0 Gram Daily Oral
Batch: **Started:** **Stopped:** 29/12/2001

SOMAC (Suspected) **Reason:**
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 29/12/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 172512 **Seq:** 1 **Gender:** M
Reported: 07/02/2002 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: 29/12/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 29/12/2001

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 172524 **Seq:** 1
Reported: 07/02/2002
Hospitalisation: Admitted to hospital
Onset Date: 30/11/2001
Outcome: Not yet recovered

Gender: F
Weight: 73.00
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor		Tremor all over body.	Patient hospitalised. diazepam.

Medicine Details:

LOSEC (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MS CONTIN (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

VIOXX (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 173741 **Seq:** 1
Reported: 20/03/2002
Hospitalisation:
Onset Date: 25/02/2002
Outcome: Not yet recovered

Gender: M
Weight: 72.00
Age: 37
DOB: 19/11/1964
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia			Required a visit to doctor. cease amitriptyline.

Medicine Details:

COLOXYL WITH SENNA (Other drug) **Reason:** Constipation
4.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** CONTIN

MS CONTIN (Suspected) **Reason:** Pain
60.0 Milligram Daily
Batch: **Started:** 12/12/2001 **Stopped:**

DIAZEPAM (Suspected) **Reason:** Abnormal involuntary movement
20.0 Milligram Daily
Batch: **Started:** 10/10/2001 **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
25.0 Milligram Daily
Batch: **Started:** 01/09/2000 **Stopped:** 14/03/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous reaction - gynaecomastia thought to be due to baclofen. the dose start date for amitriptyline is not necessarily accurate but indicates amitriptyline was started sometime in the month.

Report Detail

Case Number: 173956 **Seq:** 1
Reported: 21/03/2002
Hospitalisation: Admitted to hospital
Onset Date: 21/02/2002
Outcome: Recovered

Gender: M
Weight:
Age: 43Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertonia		Increasing spasticity upper limbs.	
Chills			
Confusional state	Required Specialist Consultation		
Hyponatraemia			
Hypothermia	Severe		Tegretol ceased, withheld fluoxetine, sodium valproate, amitriptyline. fluid restrict 1.5l/day, pharmacological washout. patient was hospitalised. ic+ supplement.
Somnolence			

Medicine Details:

PRO-BANTHINE (Other drug) **Reason:**
45.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

CLOFEN (Other drug) **Reason:**
210.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

PAXAM (Other drug) **Reason:**
3.0 Microgram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

MYADEC (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium		21/02/2002	3.2	
	Sodium		14/03/2001	130	
	Sodium		21/02/2002	119	
	Sodium		22/02/2002	131	
	Sodium		23/02/2002	129	
	Sodium		25/02/2002	133	

Additional Information:

The dose start dates are not necessarily accurate but indicates valpro, endep, fluoxetine were started sometime in the year.

Report Detail

Case Number: 173956 **Seq:** 1
Reported: 21/03/2002
Hospitalisation: Admitted to hospital
Onset Date: 21/02/2002
Outcome: Recovered

Gender: M
Weight:
Age: 43Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MEGAFOL (Other drug)			**Reason:**	
	10.0 Milligram	Daily		
Batch:	**Started:** 01/01/1995		**Stopped:**	
ZINCAPS (Other drug)			**Reason:**	
	2.0 Dose Unspecified	Daily		
Batch:	**Started:** 01/01/1999		**Stopped:**	
TEGRETOL (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:** 15/02/2002		**Stopped:**	21/02/2002
VALPRO (Suspected)			**Reason:**	
	1.0 Gram	Daily		
Batch:	**Started:** 01/01/1999		**Stopped:**	

Laboratory Investigations:

Additional Information:

The dose start dates are not necessarily accurate but indicates valpro, endep, fluoxetine were started sometime in the year.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 173956 **Seq:** 1
Reported: 21/03/2002
Hospitalisation: Admitted to hospital
Onset Date: 21/02/2002
Outcome: Recovered

Gender: M
Weight:
Age: 43Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ENDEP (Suspected)			**Reason:**	
	25.0 Milligram		Daily	
Batch:	**Started:**	S TERM		**Stopped:**
FLUOXETINE HYDROCHLORIDE (Suspected)			**Reason:**	
	60.0 Milligram		Daily	
Batch:	**Started:** 01/01/1995			**Stopped:**

Laboratory Investigations:

Additional Information:

The dose start dates are not necessarily accurate but indicates valpro, endep, fluoxetine were started sometime in the year.

Report Detail

Case Number: 174125 **Seq:** 1 **Gender:** M
Reported: 02/04/2002 **Weight:** 110.00
Hospitalisation: **Age:** 23Y
Onset Date: 23/11/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis		Drug-induced hepatitis.	

Medicine Details:

CLOZAPINE (Other drug) **Reason:** Unspecified schizophrenia
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** 06/11/2001 **Stopped:**

SODIUM VALPROATE (Suspected) **Reason:** Unspecifid affective psychosis
Oral application 2.4 Gram Daily Oral
Batch: **Started:** 17/09/2001 **Stopped:** 19/12/2001

CHLORPROMAZINE HYDROCHLORIDE (Suspected) **Reason:** Other disturbance of sensation
Oral application 800.0 Milligram Daily Oral
Batch: **Started:** 17/09/2001 **Stopped:** 18/12/2001

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Salivary secretion disturbance
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 15/11/2001 **Stopped:** 17/12/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Nil allergies of relevance.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 174137 **Seq:** 1 **Gender:** F
Reported: 02/04/2002 **Weight:**
Hospitalisation: **Age:** 58
Onset Date: **DOB:** 30/09/1943
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Room full of strange people dressed in	Required a visit to doctor. sotacor ceased.

Medicine Details:

ESTRADERM (Other drug) **Reason:** Menopausal symptoms
0.0
Batch: **Started:** **Stopped:**

KAPANOL (Suspected) **Reason:** Pain
100.0 Milligram Daily
Batch: **Started:** 01/08/2000 **Stopped:**

TRYPTANOL (Suspected) **Reason:** Pain
25.0 Milligram Daily
Batch: **Started:** 01/04/2001 **Stopped:**

STILNOX (Suspected) **Reason:** Pain
10.0 Milligram Daily
Batch: **Started:** 01/04/2001 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dose start dates for kapanol, tryptanol and stilnox are not necessarily accurate but indicates these drugs were started sometime in the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 174137 **Seq:** 1
Reported: 02/04/2002
Hospitalisation:
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 58
DOB: 30/09/1943
Causality: Causality possible

Reaction Detail

Medicine Details:

SOTACOR (Suspected) **Reason:** Otr&nos disord of heart rhythm
80.0 Milligram Daily
Batch: **Started:** 18/03/2002 **Stopped:** 27/03/2002

Laboratory Investigations:

Additional Information:

The dose start dates for kapanol, tryptanol and stilnox are not necessarily accurate but indicates these drugs were started sometime in the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 175745 **Seq:** 1
Reported: 24/05/2002
Hospitalisation: Admitted to hospital
Onset Date: 28/04/2001
Outcome: Recovered

Gender: M
Weight:
Age: 43
DOB: 22/05/1957
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysuria Pollakiuria Urine analysis abnormal			Patient was hospitalised. ceased endep.

Medicine Details:

COGENTIN (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The onset date is not necessarily accurate but indicates onset sometime in the month. patient recovered 3/2.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 175963 **Seq:** 1 **Gender:** M
Reported: 06/06/2002 **Weight:**
Hospitalisation: **Age:** 52
Onset Date: 31/05/2002 **DOB:** 28/09/1949
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tongue oedema	Life	Progressive tongue swelling	Intubation, ventilation, defib.
Asphyxia	Life		
Cardiac arrest	Life		
Respiratory depression	Life		

Medicine Details:

TEMAZEPAM (Suspected) **Reason:** Specific disorders of sleep
10.0 Milligram Daily
Batch: **Started:** 31/05/2002 **Stopped:** 31/05/2002

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
10.0 Milligram Daily
Batch: **Started:** 31/05/2002 **Stopped:** 31/05/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Ph: cancer lungs, eaton-lambert syndrome.

Report Detail

Case Number: 176093 **Seq:** 1 **Gender:** F
Reported: 11/06/2002 **Weight:**
Hospitalisation: **Age:** 93Y
Onset Date: 26/04/2002 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Fluid restriction 1l - 1.2l daily. amitriptyline ceased.

Medicine Details:

HEPARIN SODIUM (Other drug) **Reason:**
Injection 10.0 Thousand Internat Daily Intravenous
Batch: **Started:** 26/04/2002 **Stopped:**

OMEPRAZOLE (Other drug) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

PARACETAMOL (Other drug) **Reason:**
Tablet 4.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 26/04/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Likely to have accidently overdosed (ie. forgotten she had taken dose and repeated).

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 176678 **Seq:** 1
Reported: 26/06/2002
Hospitalisation:
Onset Date: 28/11/2000
Outcome: Not yet recovered

Gender: F
Weight: 110.00
Age: 53Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination			Amitriphline ceased. required medical attention.

Medicine Details:

SERZONE (Other drug) **Reason:**
800.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

RANITIDINE (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
25.0 Milligram Daily
Batch: **Started:** 24/11/2000 **Stopped:** 27/11/2000

PANADEINE FORTE (Suspected) **Reason:** Otr&unsp vertebrogen pain synd
2.0 Dose Unspecified Daily
Batch: **Started:** 24/11/2000 **Stopped:** 27/11/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The date of onset is not necessarily accurate but indicates onset sometime in the month.

Report Detail

Case Number: 176678 **Seq:** 1
Reported: 26/06/2002
Hospitalisation:
Onset Date: 28/11/2000
Outcome: Not yet recovered

Gender: F
Weight: 110.00
Age: 53Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

TEGRETOL (Suspected)		**Reason:**	Otr&unsp vertebrogen pain synd	
	200.0 Milligram	Daily		
Batch:	**Started:** 28/11/2000		**Stopped:** 04/12/2000	
COLOXYL WITH SENNA (Suspected)		**Reason:**	Constipation	
	4.0 Dose Unspecified	Daily		
Batch:	**Started:** 25/11/2000		**Stopped:**	CONTIN
HEPARIN SODIUM (Suspected)		**Reason:**	Prophylaxis	
Injection	10.0 Unit	Daily	Subcutaneous	
Batch:	**Started:** 25/11/2000		**Stopped:** 06/12/2000	
PREDNISOLONE (Suspected)		**Reason:**		
	50.0 Milligram	Daily		
Batch:	**Started:** 25/11/2000		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

The date of onset is not necessarily accurate but indicates onset sometime in the month.

Report Detail

Case Number: 176678 **Seq:** 1
Reported: 26/06/2002
Hospitalisation:
Onset Date: 28/11/2000
Outcome: Not yet recovered

Gender: F
Weight: 110.00
Age: 53Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

KAPANOL (Suspected)			**Reason:**	Otr&unsp vertebrogen pain synd
	40.0 Milligram	Daily		
Batch:	**Started:** 25/11/2000		**Stopped:**	CONTIN
TRAMADOL HYDROCHLORIDE (Suspected)			**Reason:**	
	200.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	07/12/2000
SORBILAX (Suspected)			**Reason:**	Constipation
	40.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	CONTIN

Laboratory Investigations:

Additional Information:

The date of onset is not necessarily accurate but indicates onset sometime in the month.

Report Detail

Case Number: 177338 **Seq:** 1 **Gender:** F
Reported: 22/07/2002 **Weight:** 52.00
Hospitalisation: **Age:** 54Y
Onset Date: 10/07/2002 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paranoia Delusion		Increasing paranoria experiencing panic attacks	Amitryptyline dose reduced to 75mg

Medicine Details:

SINEMET CR (Other drug) **Reason:** Paralysis agitans
0.0 Daily
Batch: **Started:** **Stopped:** CONTIN

SINEMET (Other drug) **Reason:** Late effects intracranial abscess or pyogenic
Oral application 4.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** CONTIN

BROMOCRIPTINE MESYLATE (Other drug) **Reason:** Paralysis agitans
20.0 Milligram Daily
Batch: **Started:** **Stopped:** 10/07/2002

ALPRAZOLAM (Other drug) **Reason:**
1.5 Milligram Daily
Batch: **Started:** **Stopped:** 10/07/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Recent change in medications increased in amitryptilline and addition of entacapone.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 177338 **Seq:** 1
Reported: 22/07/2002
Hospitalisation:
Onset Date: 10/07/2002
Outcome: Not yet recovered

Gender: F
Weight: 52.00
Age: 54Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ENTACAPONE (Suspected)			**Reason:**	Paralysis agitans
	200.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	10/07/2002
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Anxiety neurosis
	150.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Additional Information:

Recent change in medications increased in amitryptilline and addition of entacapone.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 179805 **Seq:** 1 **Gender:** F
Reported: 11/10/2002 **Weight:** 70.00
Hospitalisation: **Age:** 61
Onset Date: 03/10/2002 **DOB:** 16/08/1941
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperhidrosis		All over body.	Patient required a visit to

Medicine Details:

VIOXX (Other drug) **Reason:** Pain
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

ZOCOR (Other drug) **Reason:** Othr&unspec metabolic diseases
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

MOBIC (Other drug) **Reason:** Pain
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:** Abnormal involuntary movement
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 179805 **Seq:** 1
Reported: 11/10/2002
Hospitalisation:
Onset Date: 03/10/2002
Outcome: Recovered

Gender: F
Weight: 70.00
Age: 61
DOB: 16/08/1941
Causality: Causality possible

Reaction Detail

Medicine Details:

TRAMAL (Suspected)			**Reason:**	Pain	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 180764 **Seq:** 1 **Gender:** M
Reported: 18/12/2002 **Weight:** 0.00
Hospitalisation: **Age:** 72
Onset Date: 15/04/2002 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue	Required Visit to Doctor	Extreme fatigue, sweating, palpitations, especially over the last month, worse in last few days.	Nil.
Hyperhidrosis	Required Visit to Doctor		
Palpitations	Required Visit to Doctor		
Serotonin syndrome	Required Visit to Doctor		

Medicine Details:

ENDEP (Interaction) **Reason:** Otr&unsp forms neuralg&neurit
Tablet 50.0 Milligram As necessary Oral
Batch: **Started:** L TERM **Stopped:**

TRAMAL SR (Interaction) **Reason:** Osteoarthritis
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** 03/12/2001 **Stopped:**

PAROXETINE HYDROCHLORIDE (Interaction) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 01/12/2001 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 183721 **Seq:** 1 **Gender:** M
Reported: 03/03/2003 **Weight:** 0.00
Hospitalisation: **Age:** 45Y
Onset Date: 29/10/2002 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Fall Vision blurred		Ataxic reaction, loss of coordination, increasing falls and blurred vision.	Dilantin withheld, Endep was recently ceased

Medicine Details:

DILANTIN (Interaction) **Reason:**
Capsule 400.0 Milligram Daily Oral
Batch: **Started:** 22/10/2002 **Stopped:** 29/10/2002

ENDEP (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 22/10/2002

LAMOTRIGINE (Other drug) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** Contin

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
03/03/2003	Pharmacokinetics				

Additional Information:

Hx: Brain tumor (1998) treated unsuccessfully with radiotherapy. Patient having approximately two seizures weekly so Dilantin was increased. Number of seizures decreased to nil.

Report Detail

Case Number: 184012 **Seq:** 1 **Gender:** F
Reported: 14/03/2003 **Weight:** 0.00
Hospitalisation: **Age:** 49
Onset Date: 03/03/2003 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion	Caused or prolonged inpatient hospitalisation	Patient experienced a fit.	Ceased Tryptanol, Clozaril,
Coordination abnormal	Caused or prolonged inpatient hospitalisation		
Depression	Caused or prolonged inpatient hospitalisation		

Medicine Details:

QUESTRAN (Other drug) **Reason:**
Batch: **Started:** **Stopped:** 05/03/2003

VALIUM (Other drug) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 05/03/2003 contin

TRYPTANOL (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 05/03/2003

CLOZARIL (Suspected) **Reason:** Unspecified schizophrenia
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 05/03/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 184012 **Seq:** 1 **Gender:** F
Reported: 14/03/2003 **Weight:** 0.00
Hospitalisation: **Age:** 49
Onset Date: 03/03/2003 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysarthria	Caused or prolonged inpatient hospitalisation		
Fall	Caused or prolonged inpatient hospitalisation		
Muscular weakness	Caused or prolonged inpatient hospitalisation		

Medicine Details:

LAMICTAL (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 05/03/2003

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 184360 **Seq:** 1 **Gender:** M
Reported: 21/03/2003 **Weight:** 0.00
Hospitalisation: **Age:** 55
Onset Date: 10/02/2003 **DOB:** 08/11/1947
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Night sweats Hypertension Tachycardia		Night sweats, tachycardia, hypertension	Ceased tramadol.

Medicine Details:

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:** Pain

Tablet 100.0 Milligram As necessary Oral

Batch: **Started:** 10/02/2003 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Pain

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 184961 **Seq:** 1 **Gender:** F
Reported: 14/04/2003 **Weight:** 0.00
Hospitalisation: **Age:** 89
Onset Date: 08/04/2003 **DOB:** 12/11/1913
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state	Caused or prolonged inpatient hospitalisation	Hallucination and increased confusion.	
Hallucination	Caused or prolonged inpatient hospitalisation	Hallucination and increased confusion.	

Medicine Details:

TRAMAL (Interaction) **Reason:**
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** 03/04/2003 **Stopped:** 08/04/2003

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 03/04/2003 **Stopped:** 08/04/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Keflex, rentec, pariet, Lasix, Epilm.

Report Detail

Case Number: 185128 **Seq:** 1 **Gender:** F
Reported: 17/04/2003 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 73
Onset Date: **DOB:** 22/04/1929
Outcome: Recovered 03/04/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome		"Serotonin Syndrome", hallucinations, confusions, feeling hot and cold, incoordination, sweating, hyperreflexia.	

Medicine Details:

CIPRAMIL (Interaction)			**Reason:**	Depression	
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Interaction)			**Reason:**		
Tablet	10.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
TRAMAL SR (Interaction)			**Reason:**		
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Hallucinations started 2/52 prior to admission, GP increased dose of Tramadol to 100mg SR bd and patient got worse. Other drugs: Perindopril, Fluticasone, Raniditine, Tiapiokenic Acid and Salbutamol.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 186104 **Seq:** 1
Reported: 22/05/2003
Hospitalisation: Admitted to hospital
Onset Date: 04/05/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 91
DOB: 02/02/1912
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Confusion, difficult behaviour, night terrors, shaking, cold and clammy, palpitations Dx delirium? serotonin syndrome secondary to addition of SSRI to long term TCA tx	Cease antidepressants, add risperidone and oxazepam prn, and IM droperidol for severe agitation with benztropine for epse. Commence mirtazapine once confusion resolved to tx depression. Psych r/v, d/c to
Abnormal behaviour	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRYPTANOL (Interaction) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 11/05/2003

ZOLOFT (Interaction) **Reason:** Depression
Tablet Daily Oral
Batch: **Started:** 04/05/2003 **Stopped:** 11/05/2003

VIOXX (Other drug) **Reason:**
Tablet 12.5 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

BACTRIM (Other drug) **Reason:** Othr diseases of urinary tract
Tablet 400.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:** 11/05/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 186104 **Seq:** 1
Reported: 22/05/2003
Hospitalisation: Admitted to hospital
Onset Date: 04/05/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 91
DOB: 02/02/1912
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chills	Caused or prolonged inpatient hospitalisation		
Confusional state	Caused or prolonged inpatient hospitalisation		
Delirium	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ASPIRIN (Other drug) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
Tablet 50.0 Microgram Daily Oral
Batch: **Started:** **Stopped:**

LISINOPRIL (Other drug) **Reason:** Essential benign hypertension
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ALLOPURINOL (Other drug) **Reason:** Gout
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 186104 **Seq:** 1
Reported: 22/05/2003
Hospitalisation: Admitted to hospital
Onset Date: 04/05/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 91
DOB: 02/02/1912
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Palpitations	Caused or prolonged inpatient hospitalisation		
Tremor	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRYPTANOL (Interaction)			**Reason:**	Depression	
Tablet	75.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	11/05/2003
ZOLOFT (Interaction)			**Reason:**	Depression	
Tablet			Daily		Oral
Batch:	**Started:**	04/05/2003		**Stopped:**	11/05/2003
VIOXX (Other drug)			**Reason:**		
Tablet	12.5	Milligram	1 time		Oral
Batch:	**Started:**			**Stopped:**	
BACTRIM (Other drug)			**Reason:**	Othr diseases of urinary tract	
Tablet	400.0	Milligram	1 time		Oral
Batch:	**Started:**			**Stopped:**	11/05/2003

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 186104 **Seq:** 1
Reported: 22/05/2003
Hospitalisation: Admitted to hospital
Onset Date: 04/05/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 91
DOB: 02/02/1912
Causality: Causality possible

Reaction Detail

Medicine Details:

ASPIRIN (Other drug)			**Reason:**		
Tablet	100.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
OROXINE (Other drug)			**Reason:**	Myxedema	
Tablet	50.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
LISINOPRIL (Other drug)			**Reason:**	Essential benign hypertension	
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
ALLOPURINOL (Other drug)			**Reason:**	Gout	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 186382 **Seq:** 1 **Gender:** M
Reported: 29/05/2003 **Weight:** 85.00
Hospitalisation: Admitted to hospital **Age:** 33
Onset Date: 22/05/2003 **DOB:** 16/09/1969
Outcome: Recovered 22/05/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion		Tonic-clonic seizure	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**

Oral application	25.0	Milligram	Total	Oral
Batch:	**Started:**		**Stopped:**	

TRAMAL SR (Interaction) **Reason:** Pain

Tablet	300.0	Milligram	Daily	Oral
Batch:	**Started:**		**Stopped:**	23/05/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 186779 **Seq:** 1 **Gender:** F
Reported: 10/06/2003 **Weight:** 0.00
Hospitalisation: **Age:** 39
Onset Date: 11/04/2003 **DOB:** 24/12/1963
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion		Grand Mal Seizure	Midazolam 1mg iv stat. replace electrolytes.

Medicine Details:

CYCLOSPORIN-A (Suspected) **Reason:**
500.0 Milligram Daily Oral
Batch: **Started:** 31/03/2003 **Stopped:** 10/04/2003

TRAMADOL HYDROCHLORIDE (Suspected) **Reason:** Pain
400.0 Milligram Daily
Batch: **Started:** 31/03/2003 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 10/04/2003 **Stopped:** 11/04/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
10/06/2003	Other data				

Additional Information:

Patient was taking magumin, Azathiopine, Heparin.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 186937 **Seq:** 1 **Gender:** M
Reported: 16/06/2003 **Weight:** 0.00
Hospitalisation: **Age:** 85
Onset Date: 26/05/2003 **DOB:** 05/05/1918
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium	Caused or prolonged inpatient hospitalisation	Hyponatraemia and delirium	
Hyponatraemia	Caused or prolonged inpatient hospitalisation	Hyponatraemia and delirium	Demeclocycline

Medicine Details:

NATRILIX SR (Suspected) **Reason:**
Batch: **Started:** **Stopped:** 26/05/2003

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 26/05/2003

TRITACE (Suspected) **Reason:**
Tablet 1.2 Milligram Daily Oral
Batch: **Started:** **Stopped:** 26/05/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
16/06/2003	Sodium		26/05/2003	117	
16/06/2003	Sodium		11/06/2003	138	

Additional Information:

Patient was also taking Aspirin, Seretidine, Ventolin, Atrovent.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 188269 **Seq:** 1 **Gender:** M
Reported: 11/07/2003 **Weight:** 83.00
Hospitalisation: **Age:** 74
Onset Date: 10/07/2003 **DOB:** 11/04/1929
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Serotonin syndrome:- Agitation, Disorientation, Hypertension (BP 210/93)	Stopped Amitriptyline. ? Parkinson's disease. Stop Haloperidol and Metaclopramide
Coordination abnormal	Caused or prolonged inpatient hospitalisation	Unsteadiness and Dizziness, Tremor and ataxia	
Agitation	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 12.5 Milligram 1 time Oral
Batch: **Started:** 04/06/2003 **Stopped:** 11/07/2003 0

MIRTAZAPINE (Interaction) **Reason:**
Tablet 45.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

PARACETAMOL (Other drug) **Reason:**
Tablet 1.0 Gram 4 times Oral
Batch: **Started:** 04/06/2003 **Stopped:**

METOCLOPRAMIDE HYDROCHLORIDE (Other drug) **Reason:**
Tablet 10.0 Milligram 3 times Oral
Batch: **Started:** 04/06/2003 **Stopped:** 11/07/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 188269 **Seq:** 1 **Gender:** M
Reported: 11/07/2003 **Weight:** 83.00
Hospitalisation: **Age:** 74
Onset Date: 10/07/2003 **DOB:** 11/04/1929
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Disorientation	Caused or prolonged inpatient hospitalisation		
Dizziness	Caused or prolonged inpatient hospitalisation		
Hypertension	Caused or prolonged inpatient hospitalisation		

Medicine Details:

HALOPERIDOL (Other drug) **Reason:**
Tablet 500.0 Microgram 2 times Oral
Batch: **Started:** 03/06/2003 **Stopped:** 10/07/2003

FENTANYL TRANSDERMAL (Other drug) **Reason:**
Patch, dermal 50.0 Microgram Per hour Transdermal
Batch: **Started:** 06/06/2003 **Stopped:**

KETAMINE HYDROCHLORIDE (Other drug) **Reason:**
Injection 150.0 Milligram Daily Subcutaneous
Batch: **Started:** 06/06/2003 **Stopped:** 11/07/2003

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 188269 **Seq:** 1 **Gender:** M
Reported: 11/07/2003 **Weight:** 83.00
Hospitalisation: **Age:** 74
Onset Date: 10/07/2003 **DOB:** 11/04/1929
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 12.5 Milligram 1 time Oral
Batch: **Started:** 04/06/2003 **Stopped:** 11/07/2003 0

MIRTAZAPINE (Interaction) **Reason:**
Tablet 45.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

PARACETAMOL (Other drug) **Reason:**
Tablet 1.0 Gram 4 times Oral
Batch: **Started:** 04/06/2003 **Stopped:**

METOCLOPRAMIDE HYDROCHLORIDE (Other drug) **Reason:**
Tablet 10.0 Milligram 3 times Oral
Batch: **Started:** 04/06/2003 **Stopped:** 11/07/2003

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 188269 **Seq:** 1 **Gender:** M
Reported: 11/07/2003 **Weight:** 83.00
Hospitalisation: **Age:** 74
Onset Date: 10/07/2003 **DOB:** 11/04/1929
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

HALOPERIDOL (Other drug)			**Reason:**		
Tablet	500.0	Microgram	2 times		Oral
Batch:	**Started:**	03/06/2003		**Stopped:**	10/07/2003
FENTANYL TRANSDERMAL (Other drug)			**Reason:**		
Patch, dermal	50.0	Microgram	Per hour		Transdermal
Batch:	**Started:**	06/06/2003		**Stopped:**	
KETAMINE HYDROCHLORIDE (Other drug)			**Reason:**		
Injection	150.0	Milligram	Daily		Subcutaneous
Batch:	**Started:**	06/06/2003		**Stopped:**	11/07/2003

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 188389 **Seq:** 1 **Gender:** M
Reported: 11/07/2003 **Weight:** 0.00
Hospitalisation: **Age:** 50
Onset Date: 30/06/2003 **DOB:** 08/07/1952
Outcome: Recovered 30/06/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination, visual		Visual hallicantions of people in room	Oxazepam.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 100.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient here had taken 3 extra tablets.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 188436 **Seq:** 1 **Gender:** F
Reported: 11/07/2003 **Weight:** 0.00
Hospitalisation: **Age:** 77Y
Onset Date: 28/03/2003 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Left ventricular failure Confusional state Dyspnoea Hyponatraemia Oedema peripheral		Shortness of breath, LVF, hyponatraemia, ankle oedema, confusion.	Pleural tap. Fluid restricted to

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Osteoarthritis

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 30/03/2003

MELOXICAM (Suspected) **Reason:** Osteoarthritis

Tablet 7.5 Milligram Daily Oral

Batch: **Started:** **Stopped:** 29/03/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
11/07/2003	Creatinine				
11/07/2003	Potassium				
11/07/2003	Sodium				

Additional Information:

Amitriptyline and Meloxica were started sometime in September 2001.
patient was also taking Fosinopril.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 188547 **Seq:** 1 **Gender:** F
Reported: 17/07/2003 **Weight:** 0.00
Hospitalisation: **Age:** 71
Onset Date: 30/06/2003 **DOB:** 30/09/1931
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cough	Caused or prolonged inpatient hospitalisation	Coughing, fever, sweats.	
Hyperhidrosis	Caused or prolonged inpatient hospitalisation		
Pyrexia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRYPTANOL (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

EFEXOR (Interaction) **Reason:**
Tablet 37.5 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/07/2003	Chest X-ray				

Additional Information:

Tryptenol was started sometime in jan 2003 and stopped sometime in July 2003.
Efexor was started sometime in 2000.
Serotonin syndrome.
Patient was also taking Serepax and Lipitor.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 190213 **Seq:** 1 **Gender:** F
Reported: 04/09/2003 **Weight:** 58.00
Hospitalisation: **Age:** 17
Onset Date: 30/08/2003 **DOB:** 20/06/1986
Outcome: Recovered 31/08/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache	Required Visit to Doctor	Severe headache (? atypical migraine), blanching mascular rash proximal to IV site.	Cessation of Chlorpromazine.
Rash macular	Required Visit to Doctor		

Medicine Details:

Ritalin LA (Other drug) **Reason:**
Capsule 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

MAXOLON (Suspected) **Reason:**
Tablet 10.0 Milligram 3 times Oral
Batch: **Started:** 30/08/2003 **Stopped:** 31/08/2003

ASPIRIN (Suspected) **Reason:**
Tablet 600.0 Milligram 1 time Oral
Batch: **Started:** 30/08/2003 **Stopped:**

PANADEINE FORTE (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified 1 time Oral
Batch: **Started:** 30/08/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Nil previous known ADRs.

Report Detail

Case Number: 190213 **Seq:** 1 **Gender:** F
Reported: 04/09/2003 **Weight:** 58.00
Hospitalisation: **Age:** 17
Onset Date: 30/08/2003 **DOB:** 20/06/1986
Outcome: Recovered 31/08/2003 **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	50.0	Milligram	1 time		Oral
Batch:	**Started:**	30/08/2003		**Stopped:**	
DIHYDERGOT (Suspected)			**Reason:**		
	3.0	Dose Unspecified	Daily		
Batch:	**Started:**	30/08/2003		**Stopped:**	31/08/2003
LARGACTIL (Suspected)			**Reason:**		
Injection	12.5	Milligram	Daily		Intravenous
Batch:	**Started:**	30/08/2003		**Stopped:**	30/08/2003

Laboratory Investigations:

Additional Information:
Nil previous known ADRs.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 190727 **Seq:** 1 **Gender:** M
Reported: 22/09/2003 **Weight:** 0.00
Hospitalisation: **Age:** 56
Onset Date: 03/09/2003 **DOB:** 25/07/1947
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety	Caused or prolonged inpatient hospitalisation	Serotonin syndrome, anxiety, sweating, agitation, tremor, rigidity, nystagmus.	Cyproheptadine 16mg QID.
Agitation	Caused or prolonged inpatient hospitalisation		
Hyperhidrosis	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRAMAL (Interaction) **Reason:** Pain
Ointment 100.0 Milligram Daily Rectal
Batch: **Started:** 01/03/2003 **Stopped:** 03/09/2003

ENDEP (Interaction) **Reason:** Pain
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 01/09/2003 **Stopped:** 03/09/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Karvezide 300/12.5 and Nexium.

Report Detail

Case Number: 190727 **Seq:** 1 **Gender:** M
Reported: 22/09/2003 **Weight:** 0.00
Hospitalisation: **Age:** 56
Onset Date: 03/09/2003 **DOB:** 25/07/1947
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle rigidity	Caused or prolonged inpatient hospitalisation		
Nystagmus	Caused or prolonged inpatient hospitalisation		
Serotonin syndrome	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRAMAL (Interaction) **Reason:** Pain
Ointment 100.0 Milligram Daily Rectal
Batch: **Started:** 01/03/2003 **Stopped:** 03/09/2003

ENDEP (Interaction) **Reason:** Pain
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 01/09/2003 **Stopped:** 03/09/2003

Laboratory Investigations:

Additional Information:
Patient also taking Karvezide 300/12.5 and Nexium.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 190727 **Seq:** 1
Reported: 22/09/2003
Hospitalisation:
Onset Date: 03/09/2003
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 56
DOB: 25/07/1947
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor	Caused or prolonged inpatient hospitalisation		

Medicine Details:

TRAMAL (Interaction) **Reason:** Pain

Ointment	100.0 Milligram	Daily		Rectal
Batch:	**Started:** 01/03/2003		**Stopped:**	03/09/2003

ENDEP (Interaction) **Reason:** Pain

Tablet	150.0 Milligram	Daily		Oral
Batch:	**Started:** 01/09/2003		**Stopped:**	03/09/2003

Laboratory Investigations:

Additional Information:

Patient also taking Karvezide 300/12.5 and Nexium.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 190904 **Seq:** 1 **Gender:** F
Reported: 26/09/2003 **Weight:** 75.00
Hospitalisation: **Age:** 65
Onset Date: **DOB:** 15/07/1938
Outcome: Recovered 04/09/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dermatitis psoriasiform		Psoriasiform drug eruption.	

Medicine Details:

ENDEP (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** Contin

ZANTAC (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 02/06/2003

NICOTINIC ACID (Suspected) **Reason:** Othr&unspec metabolic diseases
1.5 Gram Daily
Batch: **Started:** L TERM **Stopped:** 02/06/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No clear temporal relationships for any one drug.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191095 **Seq:** 1 **Gender:** F
Reported: 07/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 83
Onset Date: **DOB:** 30/12/1919
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium		Delerium	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Pain
50.0 Milligram Daily
Batch: **Started:** 09/07/2003 **Stopped:** 21/07/2003

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:** Pain
150.0 Milligram Daily
Batch: **Started:** 09/07/2003 **Stopped:** 11/07/2003

OXYCONTIN (Interaction) **Reason:** Pain
20.0 Milligram Daily
Batch: **Started:** 11/07/2003 **Stopped:** 08/08/2003

MORPHINE SULPHATE (Interaction) **Reason:** Pain
Injection 5.0 Milligram Daily Subcutaneous
Batch: **Started:** 11/07/2003 **Stopped:** 13/07/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also administered: Coloxyl, Celebrex, Lipitor, Imdur, Prednisone, Actonel, Panamax, Flucloxacillin, Flagyl, Heparin, Ranitidine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191113 **Seq:** 1 **Gender:** F
Reported: 07/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 64
Onset Date: **DOB:** 02/04/1939
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention		Urinary retention, failed trial of void, constipation, mild stump pain, minimum phantom pain.	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 31/08/2003 **Stopped:**

VANCOMYCIN HYDROCHLORIDE (Suspected) **Reason:**
Injection 2.0 Gram Daily Intravenous
Batch: **Started:** 22/07/2003 **Stopped:** 02/09/2003

TIMENTIN (Suspected) **Reason:**
12.4 Gram Daily
Batch: **Started:** 25/07/2003 **Stopped:** 02/09/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs taken: Flagyl, Atenolol, Simvastatin, Ramipril, Glizlazide, Aspirin, Heparin, Paracetamol, Oxycontin, Coloxyl + Senna. Tof V, 1/9, 5/9, 8/9.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191114 **Seq:** 1 **Gender:** F
Reported: 07/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 76
Onset Date: 10/09/2003 **DOB:** 24/01/1927
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation		Faecal loading	Cease Amitriptyline. Add

Medicine Details:

FENTANYL (Suspected) **Reason:**
75.0 Milligram Daily
Batch: **Started:** 15/08/2003 **Stopped:** 16/09/2003

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 16/08/2003 **Stopped:** 03/09/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs taken: Aspirin, Amlodipine, GTN Patch, Metoprolol, Pantoprozole, Frusemide, Magmin, Panadol, Timentin, Dorbepoletin.
Previous allergy / ADR:
Morphine and Tramal - n.v. Prednisone-rash.
History (ischemic foot) after prev revascularization

Report Detail

Case Number: 191216 **Seq:** 1 **Gender:** F
Reported: 10/10/2003 **Weight:** 69.00
Hospitalisation: **Age:** 84Y
Onset Date: 03/07/2003 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pancytopenia		Pancytopenia	Ceased thalidomide

Medicine Details:

Thalidomide (Suspected) **Reason:** Multiple myeloma
200.0 Milligram Daily
Batch: **Started:** **Stopped:** 03/07/2003

THYROXINE SODIUM (Suspected) **Reason:**
150.0 Microgram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 50.0 Milligram Daily
Batch: **Started:** **Stopped:**

PREDNISOLONE (Suspected) **Reason:**
12.5 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
10/10/2003	Other data				

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191233 **Seq:** 1 **Gender:** F
Reported: 13/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 44
Onset Date: 15/09/2003 **DOB:** 26/02/1959
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Seizure	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Batch: **Started:** **Stopped:**

PAROXETINE HYDROCHLORIDE (Interaction) **Reason:**
Batch: **Started:** **Stopped:**

CLOZARIL (Interaction) **Reason:**
Tablet 12.0 Milligram Daily Oral
Batch: **Started:** 31/05/1993 **Stopped:** contin

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
13/10/2003	Electroencephalog				
13/10/2003	Other data				

Additional Information:

History of three seisures over the last 30 years but was not receiving anti-epileptic treatment.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191391 **Seq:** 1 **Gender:** F
Reported: 17/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 45
Onset Date: **DOB:** 17/08/1958
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash	Required Visit to Doctor	Rash on abdomen, nausea	
Nausea	Required Visit to Doctor		

Medicine Details:

ALLOPURINOL (Other drug) **Reason:** Gout
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:** Specific disorders of sleep
25.0 Milligram Daily
Batch: **Started:** 17/02/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/10/2003	Other data				

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 191473 **Seq:** 1 **Gender:** M
Reported: 21/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 75
Onset Date: 30/07/2003 **DOB:** 07/11/1927
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Fall Hallucination		Dizziness, falls x 2, hallucinating for 3 days	Phenergan and Neurontin

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Capsule 75.0 Milligram Daily Oral
Batch: **Started:** 22/07/2003 **Stopped:** 26/08/2003

PHENERGAN (Suspected) **Reason:** Other puritis conditions
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 23/07/2003 **Stopped:** 05/08/2003

NEURONTIN (Suspected) **Reason:** Pain
Capsule 900.0 Milligram Daily Oral
Batch: **Started:** 29/07/2003 **Stopped:** 04/08/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also administered: Panadol, Atrovent Forte, Salbutamol, Endone, Multivite, Coloxyl/Senna.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 193907 **Seq:** 1 **Gender:** F
Reported: 15/01/2004 **Weight:** 0.00
Hospitalisation: **Age:** 99U
Onset Date: 21/12/2003 **DOB:**
Outcome: Recovered 23/12/2003 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Oedema peripheral		Swelling of feet and slight swelling of	

Medicine Details:

ENDEP (Suspected) **Reason:** Pain
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 20/12/2003 **Stopped:** 22/12/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Lipitor, Zoton, Fosamax.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 193930 **Seq:** 1
Reported: 16/01/2004
Hospitalisation:
Onset Date: 03/01/2004
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 52Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea Agitation Dizziness Dysarthria Headache Medication error Nausea Panic attack		Within 10 mins of taking Endep , difficulty breathing, nausea, dizziness and extreme panic/agitation, headache and slurred speech.	An ambulance attended but no treatment given.

Medicine Details:

ENDEP (Suspected) **Reason:** Specific disorders of sleep

Tablet 25.0 Milligram 1 time Oral

Batch: **Started:** 03/01/2004 **Stopped:** 03/01/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Within 10 minutes of taking her husband's tablet experienced symptoms.

Report Detail

Case Number: 194311 **Seq:** 1 **Gender:** M
Reported: 04/02/2004 **Weight:** 101.00
Hospitalisation: **Age:** 37
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Agitation Hyperhidrosis Lacrimation increased Muscle twitching Restlessness		Insomnia, restlessnes, sweating, Watery eyes, agitation, had 2 episodes sudden back twitches.	

Medicine Details:

TRAMAL (Suspected) **Reason:** Pain
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

LOPID (Suspected) **Reason:** Othr&unspec metabolic diseases
Tablet 1200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 01/10/2003 **Stopped:**

OXYCODONE HYDROCHLORIDE (Suspected) **Reason:** Pain
40.0 Milligram Daily
Batch: **Started:** 01/10/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking paracetamol, Omeprazole, Coloxyl with senna, Lactulose.
Serotonin syndrome suspected.

Report Detail

Case Number: 194437 **Seq:** 1 **Gender:** M
Reported: 09/02/2004 **Weight:** 0.00
Hospitalisation: **Age:** 83
Onset Date: 10/11/2003 **DOB:** 08/05/1920
Outcome: Recovered 11/11/2003 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Angioedema	Caused or prolonged inpatient hospitalisation	Significant tongue swelling but not respiratory compromise. No dyspnoea, no stridor and able to swallow.	Phenergan 25mg oral initially; In ED - Adrenaline 0.3mg s/c, Hydrocortisone 200mg IV, Phenergan 25mg IM; In ward - Phenergan 10mg tds, Hydrocortisone 100mg qid (4 doses). Tritace and Endep ceased.

Medicine Details:

TRITACE (Suspected) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ENDEP (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

2 previous episodes of angioedema while taking Ramipril. Patient also taking Celebrex, Dilantin, Simvastatin and Frusemide. Allergies: 'sensitivity' to Pethidine and Benzodiazepines.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 194684 **Seq:** 1 **Gender:** F
Reported: 17/02/2004 **Weight:** 70.00
Hospitalisation: **Age:** 50
Onset Date: **DOB:** 02/10/1953
Outcome: Recovered 03/02/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state	Required Visit to Doctor	Confusion, night terrors, sleep walking, tremor, sweats, fear and shoulder cramps.	
Muscle spasms	Required Visit to Doctor	Leg and shoulder cramps.	
Fear	Required Visit to Doctor		
Hyperhidrosis	Required Visit to Doctor		
Sleep terror	Required Visit to Doctor		

Medicine Details:

LITHIUM CARBONATE (Suspected) **Reason:**
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SEROQUEL (Suspected) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Onset date: 09/03. Dose start and stop dates for all drugs: 06/03 an 07/03.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 194684 **Seq:** 1 **Gender:** F
Reported: 17/02/2004 **Weight:** 70.00
Hospitalisation: **Age:** 50
Onset Date: **DOB:** 02/10/1953
Outcome: Recovered 03/02/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sleep walking	Required Visit to Doctor		
Tremor	Required Visit to Doctor		

Medicine Details:

LITHIUM CARBONATE (Suspected) **Reason:**
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SEROQUEL (Suspected) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Onset date: 09/03. Dose start and stop dates for all drugs: 06/03 an 07/03.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 195119 **Seq:** 1 **Gender:** F
Reported: 04/03/2004 **Weight:** 0.00
Hospitalisation: **Age:** 79
Onset Date: 15/12/2003 **DOB:** 07/02/1924
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fall Orthostatic hypotension		Fall and postural hypotension.	Endep dose was decreased to 25mg nocte then ceased.

Medicine Details:

ENDEP (Suspected) **Reason:** Otr&unsp forms neuralg&neurit
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Oxycontin, Oroxine, Actonel, Coloxyl with Senna.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 195274 **Seq:** 1 **Gender:** U
Reported: 11/03/2004 **Weight:** 51.00
Hospitalisation: **Age:** 64
Onset Date: 22/02/2004 **DOB:** 20/08/1939
Outcome: Recovered 26/02/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thrombocytopenia	Caused or prolonged inpatient hospitalisation	Thrombocytopenia and petechiae rash.	Given Prednisolone 50mg.
Petechiae	Caused or prolonged inpatient hospitalisation		

Medicine Details:

PANADOL (Suspected) **Reason:** Pain
1.0 Gram As necessary Oral
Batch: **Started:** 13/02/2004 **Stopped:** 24/02/2004

DI-GESIC (Suspected) **Reason:** Pain
Tablet 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

TEGRETOL (Suspected) **Reason:** Herpes zoster
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 13/02/2004 **Stopped:** 24/02/2004

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 13/02/2004 **Stopped:** 24/02/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
11/03/2004	Platelets		25/02/2004	10	

Additional Information:

Report Detail

Case Number: 195274 **Seq:** 1 **Gender:** U
Reported: 11/03/2004 **Weight:** 51.00
Hospitalisation: **Age:** 64
Onset Date: 22/02/2004 **DOB:** 20/08/1939
Outcome: Recovered 26/02/2004 **Causality:** Causality possible

Reaction Detail

Medicine Details:

INDOCID (Suspected)			**Reason:**		
Capsule	75.0	Milligram	Daily		Oral
Batch:	**Started:**	13/02/2004		**Stopped:**	18/02/2004
VALTREX (Suspected)			**Reason:**	Herpes zoster	
Tablet	3.0	Gram	Daily		Oral
Batch:	**Started:**	30/01/2004		**Stopped:**	18/02/2004

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 195292 **Seq:** 1 **Gender:** M
Reported: 11/03/2004 **Weight:** 45.00
Hospitalisation: **Age:** 12
Onset Date: 13/09/2003 **DOB:** 20/01/1991
Outcome: Recovered 14/09/2003 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Hallucinations of people in room.	Dose reduced to 25mg , reaction experienced when 37.5mg dose was taken.

Medicine Details:

PARADEX (Other drug) **Reason:**
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** 10/09/2003 **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
37.5 Milligram Daily
Batch: **Started:** 13/09/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Apparently dose related, rechallenge planned.

Report Detail

Case Number: 196160 **Seq:** 1
Reported: 02/04/2004
Hospitalisation:
Onset Date: 20/02/2004
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 27
DOB: 03/12/1976
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tachycardia		Patient developed tachycardia with a heart rate of 120 bpm.	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

CLOZARIL (Interaction) **Reason:** Schizophrenia
Tablet 800.0 Milligram Daily Oral
Batch: **Started:** 20/08/2001 **Stopped:** 01/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
02/04/2004	Other data		20/02/2004	Heart rate of	

Additional Information:

Clozaril was ceased in approx 01/2002 and was recommenced on 27/10/2003

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 196478 **Seq:** 1 **Gender:** M
Reported: 15/04/2004 **Weight:** 60.00
Hospitalisation: **Age:** 46Y
Onset Date: 15/05/2003 **DOB:**
Outcome: Unrelated death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tardive dyskinesia	Required Visit to Doctor	Tardive dyskinesia	Lopinavir/Ritonavir bd, Lamivudine 150g bd

Medicine Details:

OLANZAPINE (Suspected) **Reason:** Depression
10.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient died from HIV 9/12/03. Patient was also taking Valaciclovir, Abacavir, Bactrim and Gabapentin.

Report Detail

Case Number: 196665 **Seq:** 1 **Gender:** M
Reported: 21/04/2004 **Weight:** 0.00
Hospitalisation: **Age:** 52
Onset Date: 09/11/2003 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome		Serotonin syndrome with hallucinating hyperreflexia increased sweating - hallucination, confusion and jerky movements	Ceased Fluoxetine and Amitriptyline. Gabapentin restarted

Medicine Details:

FLUOXETINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 11/11/2003 7 years

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 06/11/2003 **Stopped:** 11/11/2003

MORPHINE SULPHATE (Suspected) **Reason:**
Tablet 160.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** Months

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs: Ipratropium, Salbutamol, Calcium, Prednisolone, Omeprazole, Roxithromycin, Ampicillin and Gabapentin. History: Ca lung/emphysema/asbestos exposure. Shingles (mid thoracic) developed and diagnosed 5/11.

Report Detail

Case Number: 196682 **Seq:** 1 **Gender:** F
Reported: 21/04/2004 **Weight:** 0.00
Hospitalisation: **Age:** 84Y
Onset Date: 13/11/2003 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium Fall Hyponatraemia		Delirium secondary to hyponatremia	Ceased Atacand Plus and Amitriptyline

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Specific disorders of sleep
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 13/11/2003 1 -2 years

ATACAND PLUS 16/12.5 (Suspected) **Reason:** Essential benign hypertension
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:** 13/11/2003 1 - 2 years

PARACETAMOL (Suspected) **Reason:** Pain
3.0 Gram Daily Oral
Batch: **Started:** **Stopped:** 13/11/2003

AMLODIPINE (Suspected) **Reason:** Essential benign hypertension
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 13/11/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
21/04/2004	Sodium				

Additional Information:

Other drugs: Betamethasone, Latanoprost eye drops, Rofecoxib, Atenolol, Carbamazepine and Aspirin.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 196754 **Seq:** 1
Reported: 23/04/2004
Hospitalisation:
Onset Date: 07/10/2003
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 57
DOB: 07/02/1946
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting	Required Visit to Doctor	Nausea, vomiting and dizziness, poor	
Balance disorder	Required Visit to Doctor		
Dizziness	Required Visit to Doctor		

Medicine Details:

ENDEP (Suspected) **Reason:** Pain
10.0 Milligram Daily
Batch: **Started:** 06/10/2003 **Stopped:** 07/10/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 196958 **Seq:** 1 **Gender:** U
Reported: 03/05/2004 **Weight:** 0.00
Hospitalisation: **Age:** 64
Onset Date: 28/04/2004 **DOB:** 02/03/1940
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hyponatraemia		Acute confusional state after routine out patient colonoscopy and hyponatraemia.	

Medicine Details:

CIPRAMIL (Suspected) **Reason:** Depression
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 15/06/2003 **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 15/06/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Xanax, Micardis, Somac and Zocor.
Patient also had a bowel preparation (not specified).

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 197079 **Seq:** 1 **Gender:** F
Reported: 06/05/2004 **Weight:** 0.00
Hospitalisation: **Age:** 73
Onset Date: 08/03/2004 **DOB:** 10/07/1930
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fall	Caused or prolonged inpatient hospitalisation	Falls, slurred speech and was unable to communicate.	Amitriptyline ceased.
Communication disorder	Caused or prolonged inpatient hospitalisation		
Dysarthria	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 100.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 09/03/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Losec, Metoprolol, Aspirin and Amiodarone.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 197583 **Seq:** 1 **Gender:** M
Reported: 26/05/2004 **Weight:** 0.00
Hospitalisation: **Age:** 54Y
Onset Date: 20/12/2003 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination, visual		Visual hallucinations.	Amitriptyline ceased. Trial and error with ceasing and changing Beta blockers and other causative drug isolated.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 28/12/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Metoprolol, Omeprazole, Folic Acid, Metformin, Glimepiride and Perindopril.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 197673 **Seq:** 1 **Gender:** M
Reported: 26/05/2004 **Weight:** 0.00
Hospitalisation: **Age:** 79Y
Onset Date: 07/04/2004 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention		Urinary retention.	Ceased Amitriptyline for 4 days, then recommenced at lower dose (10mg/daily), then finally ceased 02/02/04.

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 20.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 02/02/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs taken: Paracetamol, Salbutamol, Ramipril,Tiotropium, valproate and Tramadol HCL.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 197889 **Seq:** 1 **Gender:** M
Reported: 04/06/2004 **Weight:** 0.00
Hospitalisation: **Age:** 25
Onset Date: 17/03/2004 **DOB:** 06/03/1979
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness	Required Visit to Doctor	Tonic clonic seizure, dizziness and	
Loss of consciousness	Required Visit to Doctor	Tonic clonic seizure, dizziness and	
Tonic convulsion	Required Visit to Doctor	Tonic clonic seizure, dizziness and	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs taken: Flupenthixol, Guetiapine and Diazepam. Recent changes to medications eg: Prozac, Endep, Diazepam ceased. History of drug and alcohol abuse. Schizophrenia and cognitive impairment.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199387 **Seq:** 1
Reported: 26/07/2004
Hospitalisation:
Onset Date: 15/01/2004
Outcome: Not yet recovered

Gender: M
Weight: 52.40
Age: 76
DOB: 05/03/1927
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea Musculoskeletal chest pain Pleural fibrosis Pulmonary fibrosis		Increasing shortness of breath and mild chest wall pain.	Pergolide and Comtan ceased.

Medicine Details:

PERGOLIDE (Suspected) **Reason:** Paralysis agitans
Tablet 2.2 Gram Daily Oral
Batch: **Started:** L TERM **Stopped:** 22/07/2004

COMTAN (Suspected) **Reason:** Paralysis agitans
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 22/07/2004

MADOPAR (Suspected) **Reason:** Paralysis agitans
1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:** Continuing

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
50.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** Continuing

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Sinemet and Moduretic.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199505 **Seq:** 1 **Gender:** F
Reported: 30/07/2004 **Weight:** 0.00
Hospitalisation: **Age:** 23
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suicidal ideation		Suicidal ideation	

Medicine Details:

TRYPTANOL (Suspected) **Reason:**

Batch: **Started:** **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199596 **Seq:** 1
Reported: 02/08/2004
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 99u
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Patient developed serotonin syndrome	Remeron discontinued
Drug interaction	Caused or prolonged inpatient hospitalisation	Possible serotonin syndrome but symptoms not described	All medications ceased. Remeron discontinued

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:** Pain

Batch: **Started:** **Stopped:** 0

REMERON (Interaction) **Reason:**

Tablet 15.0 Milligram Daily

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Same symptoms on restarting Remeron a few days later. Again ceased and has since recovered

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199796 **Seq:** 1
Reported: 09/08/2004
Hospitalisation: Admitted to hospital
Onset Date: 15/10/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 38Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor	Caused or prolonged inpatient hospitalisation	Initially noticed involuntary movement of head/neck muscles while watching TV. The base of her spine became sore and then tremors started - not just tremors but parkinsonian-like movements. Tremors escalated over next couple of days - on day 5 had almost spastic movements with negative control over limbs.	Alprim discontinued. Discharged 3 hours later.
Back pain	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ALPRIM (Interaction) **Reason:** Othr diseases of urinary tract
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** S TERM **Stopped:**

ENDEP (Interaction) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

LOSEC (Other drug) **Reason:** Other diseases of esophagus
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
Tablet 100.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Underlying fever and UTI. Cases (attached) which document Trimethoprim inhibiting. Dihydrofolate reductase causing decreased ED Dopamine production which may lead to Parkinsonian symptoms.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199796 **Seq:** 1
Reported: 09/08/2004
Hospitalisation: Admitted to hospital
Onset Date: 15/10/2003
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 38Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ALPRIM (Interaction) **Reason:** Othr diseases of urinary tract
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** S TERM **Stopped:**

ENDEP (Interaction) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

LOSEC (Other drug) **Reason:** Other diseases of esophagus
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
Tablet 100.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Additional Information:

Underlying fever and UTI. Cases (attached) which document Trimethoprim inhibiting. Dihydrofolate reductase causing decreased ED Dopamine production which may lead to Parkinsonian symptoms.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 199988 **Seq:** 1 **Gender:** M
Reported: 13/08/2004 **Weight:** 0.00
Hospitalisation: **Age:** 79
Onset Date: 07/01/2004 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention		Patient experienced urinary retention on [illegible]	Ceased Amitriptyline for 4 days, then recommecned at 10mg/nocte then ceased again [illegible]

Medicine Details:

SODIUM VALPROATE (Other drug)			**Reason:**	
Tablet	600.0	Milligram		Oral
Batch:	**Started:**		**Stopped:**	
PARACETAMOL (Other drug)			**Reason:**	
Tablet	4.0	Gram		Oral
Batch:	**Started:**		**Stopped:**	
SALBUTAMOL (Other drug)			**Reason:**	
Inhalation	5.0	Milligram		Inhalation
Batch:	**Started:**		**Stopped:**	
RAMIPRIL (Other drug)			**Reason:**	
Capsule	10.0	Milligram		Oral
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 199988 **Seq:** 1 **Gender:** M
Reported: 13/08/2004 **Weight:** 0.00
Hospitalisation: **Age:** 79
Onset Date: 07/01/2004 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

GLICLAZIDE (Other drug)			**Reason:**		
Tablet	160.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
TIOTROPIUM (Suspected)			**Reason:**		
Capsule	18.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
TRAMADOL HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	200.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	07/01/2004

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 200190 **Seq:** 1
Reported: 20/08/2004
Hospitalisation: Admitted to hospital
Onset Date: 01/08/2004
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 80
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Epilepsy	Caused or prolonged inpatient hospitalisation	Patient developed insomnia, hallucinations, increased agitation and aggression. He had a high alcohol intake and experienced two epileptic fits within 72 hours.	Aricept ceased.
Aggression	Caused or prolonged inpatient hospitalisation		
Agitation	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ALCOHOL (Suspected) **Reason:**
Oral
Batch: **Started:** **Stopped:**

TOLVON (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

TEMAZEPAM (Suspected) **Reason:**
Oral
Batch: **Started:** 01/08/2004 **Stopped:**

ARICEPT (Suspected) **Reason:** Presenile dementia
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 09/04/2004 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 200190 **Seq:** 1
Reported: 20/08/2004
Hospitalisation: Admitted to hospital
Onset Date: 01/08/2004
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 80
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination	Caused or prolonged inpatient hospitalisation		
Insomnia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 200342 **Seq:** 1 **Gender:** M
Reported: 27/08/2004 **Weight:** 69.00
Hospitalisation: **Age:** 64
Onset Date: 13/08/2004 **DOB:** 09/11/1939
Outcome: Recovered 18/08/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Inappropriate antidiuretic hormon		SIADH suspiciousness.	Stop suspected drugs.

Medicine Details:

MAGNESIUM ASPARTATE (Other drug) **Reason:** Pain in limb
Tablet 1.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

CEPHALEXIN (Other drug) **Reason:** Othr diseases of urinary tract
2.0 Gram Daily
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Specific disorders of sleep
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

IRBESARTAN-HYDROCHLOROTHIAZIDE (Suspected) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
27/08/2004	Sodium				Na+ 125 mmol/l, 26 mmol/l, 137 mmol/l, 2 days.
27/08/2004	Urine osmolality				Ur osm 280 mmol/kg, 495 mmol/kg. 2 days.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 200342 **Seq:** 1 **Gender:** M
Reported: 27/08/2004 **Weight:** 69.00
Hospitalisation: **Age:** 64
Onset Date: 13/08/2004 **DOB:** 09/11/1939
Outcome: Recovered 18/08/2004 **Causality:** Causality possible

Reaction Detail

Medicine Details:

MELOXICAM (Suspected) **Reason:** Other diseases of joint
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 200396 **Seq:** 1 **Gender:** F
Reported: 30/08/2004 **Weight:** 65.00
Hospitalisation: **Age:** 99U
Onset Date: 15/08/2004 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Swollen tongue		Patient experienced tongue swelling.	

Medicine Details:

TEMAZE (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 23/04/2004 **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 11/08/2000 **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 200979 **Seq:** 1 **Gender:** U
Reported: 21/09/2004 **Weight:** 0.00
Hospitalisation: **Age:** 77Y
Onset Date: 20/07/2004 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sedation Sleep disorder		Excessive sedation, causes patient to sleep thru urge to urinate.	

Medicine Details:

QVAR AUTOHALER (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

SEREVENT (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

VENTOLIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

DOTHEP (Other drug) **Reason:**
Tablet 75.0 Milligram Oral
Batch: **Started:** 01/07/1998 **Stopped:** 20/07/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Problem commenced when patient changed from dothiepin to amitriptyline (same dose).

Report Detail

Case Number: 200979 **Seq:** 1
Reported: 21/09/2004
Hospitalisation:
Onset Date: 20/07/2004
Outcome: Not yet recovered

Gender: U
Weight: 0.00
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

KALMA (Other drug)			**Reason:**		
	0.5	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	contin
ENDEP (Suspected)			**Reason:**		
Tablet	75.0	Milligram	Daily	Oral	
Batch:	**Started:**	20/07/2004		**Stopped:**	0

Laboratory Investigations:

Additional Information:

Problem commenced when patient changed from dothiepin to amitriptyline (same dose).

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 201589 **Seq:** 1
Reported: 11/10/2004
Hospitalisation: Admitted to hospital
Onset Date: 27/02/2004
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 34
DOB: 12/01/1970
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Jaundice	Caused or prolonged inpatient hospitalisation	Patient developed jaundice and liver function tests abnormal nos.	Amitriptyline ceased.
Chromaturia	Caused or prolonged inpatient hospitalisation		
Faeces discoloured	Caused or prolonged inpatient hospitalisation		

Medicine Details:

VOLTAREN (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
Batch: **Started:** **Stopped:**

DIAZEPAM (Other drug) **Reason:**
5.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

CAPADEX (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
Capsule As necessary Oral
Batch: **Started:** **Stopped:**

PANADEINE FORTE (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
Tablet As necessary Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
11/10/2004	Other data		03/03/2004	CT abdo: Mild	

Additional Information:

Report Detail

Case Number: 201589 **Seq:** 1
Reported: 11/10/2004
Hospitalisation: Admitted to hospital
Onset Date: 27/02/2004
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 34
DOB: 12/01/1970
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Liver function test abnormal	Caused or prolonged inpatient hospitalisation		
Rash	Caused or prolonged inpatient hospitalisation		

Medicine Details:

CEPHALEXIN (Other drug) **Reason:** Pulmonary collapse
2.0 Gram Daily
Batch: **Started:** 24/02/2004 **Stopped:** 05/03/2004

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 15/12/2003 **Stopped:** 15/02/2004

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 204761 **Seq:** 1 **Gender:** F
Reported: 01/02/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 38
Onset Date: 18/11/2004 **DOB:**
Outcome: Recovered 07/12/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Electrocardiogram QT prolonged	Caused or prolonged inpatient hospitalisation	Prolonged QT interval - collapse, drowsiness, jerking movements previous few days	Stop Quetiapine and
Circulatory collapse	Caused or prolonged inpatient hospitalisation		
Dyskinesia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

HUMULIN NPH (Other drug) **Reason:** Diabetes mellitus
Injection 10.0 Unit Subcutaneous
Batch: **Started:** **Stopped:**

IRBESARTAN (Other drug) **Reason:** Essential benign hypertension
150.0 Milligram Oral
Batch: **Started:** **Stopped:**

ATORVASTATIN (Other drug) **Reason:** Othr&unspec metabolic diseases
40.0 Milligram
Batch: **Started:** **Stopped:**

METHADONE HYDROCHLORIDE (Other drug) **Reason:**
20.0 Milligram Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
01/02/2005	Electrocardiograph		18/11/2004	499	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 204761 **Seq:** 1 **Gender:** F
Reported: 01/02/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 38
Onset Date: 18/11/2004 **DOB:**
Outcome: Recovered 07/12/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ACTRAPID (Other drug) **Reason:** Diabetes mellitus
Injection 6.0 Unit Subcutaneous
Batch: **Started:** **Stopped:**

MOTILIUM (Other drug) **Reason:**
3.0 Dose Unspecified Oral
Batch: **Started:** **Stopped:**

METOCLOPRAMIDE HYDROCHLORIDE (Other drug) **Reason:**
30.0 Milligram Oral
Batch: **Started:** **Stopped:**

QUETIAPINE (Suspected) **Reason:** Depression
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 15/02/2004 **Stopped:** 22/11/2004 0

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 204761 **Seq:** 1 **Gender:** F
Reported: 01/02/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 38
Onset Date: 18/11/2004 **DOB:**
Outcome: Recovered 07/12/2004 **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression

Tablet 250.0 Milligram Daily Oral

Batch: **Started:** 15/02/2004 **Stopped:** 19/11/2004 0

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 205132 **Seq:** 1 **Gender:** M
Reported: 09/02/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 80
Onset Date: **DOB:** 11/10/1924
Outcome: Recovered 27/10/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	serotonin syndrome suspected (tremor and "almost aphasic")	ceased both citalopram and amitriptyline
Tremor	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 25.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:** 27/10/2004

CITALOPRAM HYDROBROMIDE (Suspected) **Reason:**
Tablet 30.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:** 21/10/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 205275 **Seq:** 1
Reported: 15/02/2005
Hospitalisation: Admitted to hospital
Onset Date: 30/12/2004
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 41
DOB: 08/07/1963
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination	Caused or prolonged inpatient hospitalisation	"Agitated, Delusional, Hallucinating, Confused, uncooperative,combative, Jerking movements of arms and legs"	Diazepam 5mg iv ; Midazolam 5mg iv.; Symptoms persisted patient transferred to St George Hospital Sydney,
Abnormal behaviour	Caused or prolonged inpatient hospitalisation		
Aggression	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ENDEP (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

TRAMAL (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 205275 **Seq:** 1
Reported: 15/02/2005
Hospitalisation: Admitted to hospital
Onset Date: 30/12/2004
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 41
DOB: 08/07/1963
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation	Caused or prolonged inpatient hospitalisation		
Confusional state	Caused or prolonged inpatient hospitalisation		
Delusional disorder, unspecified t	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ENDEP (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

TRAMAL (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 205275 **Seq:** 1
Reported: 15/02/2005
Hospitalisation: Admitted to hospital
Onset Date: 30/12/2004
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 41
DOB: 08/07/1963
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ENDEP (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

TRAMAL (Suspected) **Reason:**
Dose Unspecified Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 205621 **Seq:** 1
Reported: 28/02/2005
Hospitalisation:
Onset Date: 15/09/2004
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 59
DOB: 12/05/1945
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chronic obstructive pulmonary di		Patient developed chronic obstructive airways disease exacerbated.	Atrovent puffer plus nebuliser and Ventolin given.

Medicine Details:

CALTRATE (Other drug) **Reason:**
10.0 Milligram
Batch: **Started:** **Stopped:**

NAPROSYN SR (Other drug) **Reason:**
40.0 Milligram
Batch: **Started:** **Stopped:**

PREMARIN (Other drug) **Reason:** Menopausal symptoms
Batch: **Started:** **Stopped:**

LIPITOR (Other drug) **Reason:** Othr&unspec metabolic diseases
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has been on symbicort but this was stopped whilst she trialled spiriva which didn't make much difference

Report Detail

Case Number: 205621 **Seq:** 1
Reported: 28/02/2005
Hospitalisation:
Onset Date: 15/09/2004
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 59
DOB: 12/05/1945
Causality: Causality possible

Reaction Detail

Medicine Details:

LASIX (Other drug)				**Reason:**	Essential benign hypertension
Batch:	**Started:**				**Stopped:**
ENDONE (Suspected)				**Reason:**	Pain
Tablet	5.0	Milligram	As necessary		Oral
Batch:	**Started:**				**Stopped:**
PETHIDINE HYDROCHLORIDE (Suspected)				**Reason:**	Pain
	100.0	Milligram	As necessary		
Batch:	**Started:**				**Stopped:**
ALODORM (Suspected)				**Reason:**	Specific disorders of sleep
Tablet	5.0	Milligram	Daily		Oral
Batch:	**Started:**				**Stopped:**

Laboratory Investigations:

Additional Information:

Patient has been on symbicort but this was stopped whilst she trialled spiriva which didn't make much difference

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 205621 **Seq:** 1
Reported: 28/02/2005
Hospitalisation:
Onset Date: 15/09/2004
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 59
DOB: 12/05/1945
Causality: Causality possible

Reaction Detail

Medicine Details:

OXYCONTIN (Suspected)			**Reason:**	Pain	
Tablet	80.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
ENDEP (Suspected)			**Reason:**	Pain	
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
MURELAX (Suspected)			**Reason:**	Abnormal involuntary movement	
Tablet	30.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Patient has been on symbicort but this was stopped whilst she trialled spiriva which didn't make much difference

Report Detail

Case Number: 205854 **Seq:** 1 **Gender:** M
Reported: 08/03/2005 **Weight:** 80.00
Hospitalisation: **Age:** 79
Onset Date: **DOB:** 12/02/1926
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness	Required Visit to Doctor	Patient experienced dizziness, nausea, slurred speech and anorexia.	
Anorexia	Required Visit to Doctor		
Dysarthria	Required Visit to Doctor		
Nausea	Required Visit to Doctor		

Medicine Details:

STEMETIL (Interaction) **Reason:** Nausea and vomiting
Batch: **Started:** **Stopped:** 0

CARDIZEM CD (Interaction) **Reason:** Otr&unsp vertebrogen pain synd
Capsule 360.0 Milligram Oral
Batch: **Started:** **Stopped:**

ENDEP (Interaction) **Reason:** Specific disorders of sleep
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 03/11/2004 **Stopped:** 05/11/2004 0

LANOXIN (Other drug) **Reason:** Otr&nos disord of heart rhythm
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 205854 **Seq:** 1
Reported: 08/03/2005
Hospitalisation:
Onset Date:
Outcome: Recovered

Gender: M
Weight: 80.00
Age: 79
DOB: 12/02/1926
Causality: Causality possible

Reaction Detail

Medicine Details:

SOMAC (Other drug)		**Reason:**	Other diseases of esophagus	
Batch:	**Started:**		**Stopped:**	0
ANGININE (Other drug)		**Reason:**	Angina pectoris w/o hyperten	
Batch:	**Started:**		**Stopped:**	0
DAPA-TABS (Other drug)		**Reason:**	Essential benign hypertension	
Batch:	**Started:**		**Stopped:**	
ATACAND (Other drug)		**Reason:**	Essential benign hypertension	
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 205854 **Seq:** 1 **Gender:** M
Reported: 08/03/2005 **Weight:** 80.00
Hospitalisation: **Age:** 79
Onset Date: **DOB:** 12/02/1926
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ISCOVER (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
VALIUM (Other drug)		**Reason:**	Anxiety neurosis
Batch:	**Started:**		**Stopped:**
ZYLOPRIM (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
STILNOX (Other drug)		**Reason:**	Specific disorders of sleep
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 205854 **Seq:** 1
Reported: 08/03/2005
Hospitalisation:
Onset Date:
Outcome: Recovered

Gender: M
Weight: 80.00
Age: 79
DOB: 12/02/1926
Causality: Causality possible

Reaction Detail

Medicine Details:

DOLOXENE (Other drug) **Reason:** Pain

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 206217 **Seq:** 1 **Gender:** M
Reported: 22/03/2005 **Weight:** 75.00
Hospitalisation: Required a visit to the doctor **Age:** 58
Onset Date: 15/01/2005 **DOB:** 19/01/1946
Outcome: Recovered 15/03/2005 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash		The patient developed a hot, itchy, red	

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 206347 **Seq:** 1 **Gender:** F
Reported: 30/03/2005 **Weight:** 65.00
Hospitalisation: **Age:** 64
Onset Date: 03/03/2005 **DOB:** 24/04/1940
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Flushing Malaise		Arm swollen red hot, flushed later in the day, felt sick after ingestion.	

Medicine Details:

ENDEP (Suspected)			**Reason:**	
Tablet	10.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**
OROXINE (Suspected)			**Reason:**	
Tablet	100.0	Microgram	Daily	Oral
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 206482 **Seq:** 1 **Gender:** M
Reported: 01/04/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 21Y
Onset Date: 15/04/2004 **DOB:**
Outcome: Recovered 18/04/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination, visual	Caused or prolonged inpatient hospitalisation	Visual hallucinations, confusion and	Decrease dose of Oxycodone, withheld Tramadol and restarted at lower dose, stop Amitriptyline
Confusional state	Caused or prolonged inpatient hospitalisation		
Disorientation	Caused or prolonged inpatient hospitalisation		

Medicine Details:

MAGMIN (Other drug) **Reason:** Acute heart failure,undefined
Batch: **Started:** **Stopped:**

CREON FORTE (Other drug) **Reason:** Acute heart failure,undefined
Batch: **Started:** **Stopped:**

OMEPRAZOLE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

CENTRUM TABLETS (Other drug) **Reason:** Acute heart failure,undefined
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 206482 **Seq:** 1 **Gender:** M
Reported: 01/04/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 21Y
Onset Date: 15/04/2004 **DOB:**
Outcome: Recovered 18/04/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug interaction	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AUGMENTIN DUO FORTE (Other drug) **Reason:** Otr&nos infec&parasit diseases
Batch: **Started:** **Stopped:**

AMANTADINE HYDROCHLORIDE (Other drug) **Reason:** Other disturbance of sensation
Batch: **Started:** **Stopped:**

SALBUTAMOL (Other drug) **Reason:** Acute heart failure,undefined
Batch: **Started:** **Stopped:**

OXYCODONE HYDROCHLORIDE (Suspected) **Reason:** Pain
10.0 Milligram As necessary Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 206482 **Seq:** 1 **Gender:** M
Reported: 01/04/2005 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 21Y
Onset Date: 15/04/2004 **DOB:**
Outcome: Recovered 18/04/2004 **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRAMADOL HYDROCHLORIDE (Suspected)			**Reason:** Pain			
Tablet, modified release	400.0	Milligram	Daily		Oral	
Batch:	**Started:** 08/04/2004			**Stopped:** 15/04/2004		0
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:** Pain			
	25.0	Milligram	Daily		Oral	
Batch:	**Started:**			**Stopped:** 17/04/2004		

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 208190 **Seq:** 1
Reported: 26/05/2005
Hospitalisation: Required a visit to the doctor
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 58.00
Age: 68
DOB: 22/12/1936
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Periorbital oedema		Patient experienced puffy hands and periorbital oedema.	

Medicine Details:

ADVANTAN (Other drug) **Reason:** Otr ecz&der due otr&unsp cause
Batch: **Started:** **Stopped:**

CALTRATE + VITAMIN D (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

PANADOL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ANDRODERM (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

On low oxalat diet too.

Report Detail

Case Number: 208190 **Seq:** 1
Reported: 26/05/2005
Hospitalisation: Required a visit to the doctor
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 58.00
Age: 68
DOB: 22/12/1936
Causality: Causality possible

Reaction Detail

Medicine Details:

OVESTIN (Other drug)			**Reason:** Vaginitis&vulvitis	
Batch:	**Started:**			**Stopped:**
ENDEP (Suspected)			**Reason:**	
Tablet	25.0 Milligram		Daily	Oral
Batch:	**Started:** 01/05/2005			**Stopped:**

Laboratory Investigations:

Additional Information:
On low oxalat diet too.

Report Detail

Case Number: 208670 **Seq:** 1 **Gender:** M
Reported: 12/06/2005 **Weight:** 64.00
Hospitalisation: **Age:** 34Y
Onset Date: 10/06/2005 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Incapacity/disability	many features of Serotonin Syndrome (as outlined in online Australian T.G.A. & U.S.A. N.I.H. data) experienced for approximately 4 hours: Mixed cognitive-behavioural changes: confusion, alternating brief periods of agitation & long periods of elation, slowed perception of time; Autonomic dysfunction: hypertension, & Neuromuscular dysfunction: dizziness, loss of balance & uncoordination, myoclonus: mostly positive but some negative sporadic sleep-hypermyoclonus, & automatic dysreflexia: decreased	discontinuation of/no further use of Tramadol (also no pain relief received from drug anyway, so doubly a bad experience)
Agitation	Incapacity/disability		

Medicine Details:

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:**
Capsule 100.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 2.5 Milligram Daily Oral
Batch: 5A182 **Started:** 01/02/2004 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 208670 **Seq:** 1 **Gender:** M
Reported: 12/06/2005 **Weight:** 64.00
Hospitalisation: **Age:** 34Y
Onset Date: 10/06/2005 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Autonomic nervous system imbal	Incapacity/disability		
Balance disorder	Incapacity/disability		
Cognitive disorder	Incapacity/disability		
Confusional state	Incapacity/disability		
Coordination abnormal	Incapacity/disability		
Dizziness	Incapacity/disability		

Medicine Details:

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:**
Capsule 100.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 2.5 Milligram Daily Oral
Batch: 5A182 **Started:** 01/02/2004 **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 208670 **Seq:** 1 **Gender:** M
Reported: 12/06/2005 **Weight:** 64.00
Hospitalisation: **Age:** 34Y
Onset Date: 10/06/2005 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective	Incapacity/disability		
Heart rate decreased	Incapacity/disability		
Hypertension	Incapacity/disability		
Inappropriate affect	Incapacity/disability		
Myoclonus	Incapacity/disability		

Medicine Details:

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:**
Capsule 100.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Interaction) **Reason:**
Tablet 2.5 Milligram Daily Oral
Batch: 5A182 **Started:** 01/02/2004 **Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 209485 **Seq:** 1
Reported: 05/07/2005
Hospitalisation: Required a visit to the doctor
Onset Date: 01/06/2005
Outcome: Recovered

Gender: M
Weight: 90.00
Age: 48Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lethargy Dry mouth		The patient became totally lethargic, "can't move at all' and had a dry mouth.	

Medicine Details:

ENDONE (Other drug) **Reason:** Pain
1.0 Dosage Unit not a
Batch: **Started:** **Stopped:**

PARIET (Other drug) **Reason:** Otr disorders stomach function
20.0 Milligram
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:** Pain
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 31/05/2005 **Stopped:** 02/06/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Suspect drug certain rechallenge.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

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Report Detail

Case Number: 210086 **Seq:** 1 **Gender:** F
Reported: 25/07/2005 **Weight:** 70.00
Hospitalisation: **Age:** 47
Onset Date: 09/07/2005 **DOB:** 02/02/1958
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Therapy regimen changed	Life	Clouding of consciousness, tachycardia and myoclonic jerks. Patient not used with this dose - ok with 75mg Amitriptyline.	
Depressed level of consciousnes	Life		
Myoclonus	Life		
Tachycardia	Life		

Medicine Details:

IMIGRAN (Suspected) **Reason:** Migraine
Tablet 50.0 Milligram 1 time Oral
Batch: **Started:** 09/07/2005 **Stopped:**

ENDEP (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 17/06/2005 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 210123 **Seq:** 1 **Gender:** F
Reported: 26/07/2005 **Weight:** 0.00
Hospitalisation: **Age:** 36Y
Onset Date: 24/06/2004 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence Somnolence		Patient ok during the day but increased drowsiness as day progresses, fell asleep at he dinner table	

Medicine Details:

COLOFAC (Other drug) **Reason:**
135.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

GAVISCON (Other drug) **Reason:**
As necessary
Batch: **Started:** L TERM **Stopped:**

UREMIDE (Other drug) **Reason:**
80.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ARTANE (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 210123 **Seq:** 1 **Gender:** F
Reported: 26/07/2005 **Weight:** 0.00
Hospitalisation: **Age:** 36Y
Onset Date: 24/06/2004 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

NEXIUM (Other drug)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
MOTILIUM (Other drug)			**Reason:**		
	10.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
ENDEP (Suspected)			**Reason:**	Depression	
Tablet	25.0	Milligram	Daily		Oral
Batch:	**Started:**	14/05/2004		**Stopped:**	
VALIUM (Suspected)			**Reason:**		
Tablet	15.0	Milligram	Daily		Oral
Batch:	**Started:**		L TERM	**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 210123 **Seq:** 1 **Gender:** F
Reported: 26/07/2005 **Weight:** 0.00
Hospitalisation: **Age:** 36Y
Onset Date: 24/06/2004 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

ERYTHROMYCIN (Suspected) **Reason:** Ac upr resp inf,mult,unsp stes
1.6 Gram Daily Oral
Batch: **Started:** 20/06/2004 **Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 210214 **Seq:** 1 **Gender:** F
Reported: 28/07/2005 **Weight:** 0.00
Hospitalisation: **Age:** 74Y
Onset Date: 30/11/2004 **DOB:**
Outcome: Death 30/11/2004 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sudden death	Death	Patient had died suddenly, the cause of death stated as 'Endep toxicity'.	

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 210412 **Seq:** 1
Reported: 05/08/2005
Hospitalisation: Admitted to hospital
Onset Date: 05/06/2005
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 45
DOB: 22/04/1960
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Toxic epidermal necrolysis	Caused or prolonged inpatient hospitalisation	The patient was diagnosed with mild toxic epidermal necrolysis.	Treated with Celestone M, wet wraps, IV Hydrocortisone, Lignocaine viscous, Promethazine oral and Amosan mouthwash.

Medicine Details:

ESOMEPRAZOLE MAGNESIUM TRIHYDRATE (Other drug) **Reason:**
Batch: **Started:** L TERM **Stopped:**

SALBUTAMOL (Other drug) **Reason:**
Batch: **Started:** L TERM **Stopped:**

SERETIDE ACCUHALER NOS (Other drug) **Reason:**
Batch: **Started:** L TERM **Stopped:**

CARBAMAZEPINE (Suspected) **Reason:** Pain
Oral
Batch: **Started:** 13/05/2005 **Stopped:** 05/06/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
05/08/2005	Histology				Histology: Complete epidermal necrosis and superficial dermal inflammaction, could represent either steven johnson syndrome or toxic epidermal necrolysis. Final diagnosis: Mild TENS (proximal and distal distribution, atypical targetoid lesions).

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 210412 **Seq:** 1
Reported: 05/08/2005
Hospitalisation: Admitted to hospital
Onset Date: 05/06/2005
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 45
DOB: 22/04/1960
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Batch: **Started:** 13/05/2005 **Stopped:** 05/06/2005

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212062 **Seq:** 1
Reported: 28/09/2005
Hospitalisation:
Onset Date: 01/09/2005
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 56
DOB: 29/01/1949
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Musculoskeletal stiffness Agitation Clonus Hyperhidrosis Hypertonia Serotonin syndrome		Patient experienced stiffness, agitation, sweating, clonus ++ increased tone, serotonin syndrome or NMS.	Ceased all medication, ICU.

Medicine Details:

OLANZAPINE (Suspected) **Reason:** Manic-depression psycho,manic
Batch: **Started:** 18/08/2005 **Stopped:** 08/09/2005

DOXEPIN HYDROCHLORIDE (Suspected) **Reason:**
Batch: **Started:** **Stopped:** 08/09/2005

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Batch: **Started:** **Stopped:** 08/09/2005

MELLERIL (Suspected) **Reason:**
Batch: **Started:** **Stopped:** 08/09/2005 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 212062 **Seq:** 1
Reported: 28/09/2005
Hospitalisation:
Onset Date: 01/09/2005
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 56
DOB: 29/01/1949
Causality: Causality possible

Reaction Detail

Medicine Details:

RISPERIDONE (Suspected)			**Reason:**	Otr spec symp psychopathol nec
Batch:	**Started:** 28/08/2005		**Stopped:**	08/09/2005
LITHIUM CARBONATE (Suspected)			**Reason:**	Manic-depression psycho,manic
Tablet	1.0 Gram	Daily		Oral
Batch:	**Started:** 18/08/2005		**Stopped:**	08/09/2005 0

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212367 **Seq:** 1
Reported: 10/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 27/01/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 76
DOB: 19/01/1929
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue	Caused or prolonged inpatient hospitalisation	The patient experienced tiredness and	Gabapentin and Amitriptyline
Confusional state	Caused or prolonged inpatient hospitalisation		

Medicine Details:

GABAPENTIN (Suspected) **Reason:** Pain

Capsule 300.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

Tablet 20.0 Milligram Daily Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report 212366 is the first sequence to this report.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212405 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 12/03/2005
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 43
DOB: 21/04/1961
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence	Caused or prolonged inpatient hospitalisation	The patient experienced drowsiness and	Cyclosporin ceased.
Headache	Caused or prolonged inpatient hospitalisation		

Medicine Details:

CYCLOSPORIN-A (Interaction) **Reason:** Otr dis of arteries&arterioles
Capsule 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

CARBAMAZEPINE (Interaction) **Reason:**
Tablet 800.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ENDEP (Interaction) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

GEMFIBROZIL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 212405 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 12/03/2005
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 43
DOB: 21/04/1961
Causality: Causality probable

Reaction Detail

Medicine Details:

AZATHIOPRINE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
DEPTRAN (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PREDNISOLONE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
KAPANOL (Other drug)		**Reason:**	
	100.0 Milligram		
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212405 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 12/03/2005
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 43
DOB: 21/04/1961
Causality: Causality probable

Reaction Detail

Medicine Details:

KAPANOL (Other drug) **Reason:**
60.0 Milligram
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

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Report Detail

Case Number: 212406 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 05/04/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 99u
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension	Caused or prolonged inpatient hospitalisation	The patient developed hypotension and mild hyponatraemia.	Dose of Endep reduced. Accupril and Natrilix ceased.
Hyponatraemia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ENDEP (Suspected) **Reason:** Specific disorders of sleep
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

NATRILIX SR (Suspected) **Reason:** Acute heart failure,undefined
Tablet 1.5 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ACCUPRIL (Suspected) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Mild hyponatraemia.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212418 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 01/05/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 86
DOB: 03/06/1918
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Caused or prolonged inpatient hospitalisation	The patient experienced overflow diarrhoea and vomiting.	Rehydrated. Methadone, Gabapentin and Amitriptyline ceased.
Constipation	Caused or prolonged inpatient hospitalisation		
Faecal incontinence	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ISOSORBIDE MONONITRATE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
ASPIRIN (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
AMIODARONE HYDROCHLORIDE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
RAMIPRIL (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 212418 **Seq:** 1
Reported: 11/10/2005
Hospitalisation: Admitted to hospital
Onset Date: 01/05/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 86
DOB: 03/06/1918
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting	Caused or prolonged inpatient hospitalisation		

Medicine Details:

PHYSEPTONE (Suspected) **Reason:** Otr&unsp forms neuralg&neurit

Tablet	20.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**

	20.0	Milligram	Daily	

Batch: **Started:** **Stopped:**

GABAPENTIN (Suspected) **Reason:**

Capsule	600.0	Milligram	Daily	Oral

Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 214540 **Seq:** 1
Reported: 04/01/2006
Hospitalisation:
Onset Date:
Outcome: Unknown

Gender: F
Weight: 0.00
Age: 46
DOB: 01/04/1959
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Restlessness Agitation Dyspnoea Insomnia Palpitations		The patient became restless, had some difficulty breathing, had some palpitations, was agitated and she could not sleep.	

Medicine Details:

DIABEX (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

DIAMICRON (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

PANADEINE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

DOLOBID (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 214540 **Seq:** 1 **Gender:** F
Reported: 04/01/2006 **Weight:** 0.00
Hospitalisation: **Age:** 46
Onset Date: **DOB:** 01/04/1959
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

TRAMAL (Suspected)			**Reason:**	
Batch:	**Started:** 29/11/2003			**Stopped:**
PROZAC (Suspected)			**Reason:**	
				Oral
Batch:	**Started:** 15/05/2005			**Stopped:**
AVAPRO (Suspected)			**Reason:**	
Tablet				Oral
Batch:	**Started:**			**Stopped:**
ENDEP (Suspected)			**Reason:**	
Tablet	50.0 Milligram		Daily	Oral
Batch:	**Started:** 15/03/2003			**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 214654 **Seq:** 1
Reported: 06/01/2006
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 66Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	The patient developed suspected serotonin syndrome.	Zoloft ceased.

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

ZOLOFT (Suspected) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 10/10/2000 **Stopped:** 16/10/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Had occurred previously whilst using Cipramil.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 215131 **Seq:** 1 **Gender:** U
Reported: 20/01/2006 **Weight:** 57.00
Hospitalisation: **Age:** 16Y
Onset Date: 03/12/2004 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state	Caused or prolonged inpatient hospitalisation	Patient experienced a confusional state and delirium.	Supportive treatment.
Delirium	Caused or prolonged inpatient hospitalisation		

Medicine Details:

PARACETAMOL (Other drug) **Reason:**
1.0 Gram Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 50.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 215320 **Seq:** 1 **Gender:** M
Reported: 27/01/2006 **Weight:** 90.00
Hospitalisation: **Age:** 62
Onset Date: **DOB:** 01/09/1943
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Pyrexia Serotonin syndrome		The patient is unable to sleep at night and spends time in bed during the day. He feels useless, has less interest in everything and has suicidal thoughts now and again. He has a poor appetite, has lost weight, has nerves, restlessness, tiredness, memory impairment, fevers, stiffness in his joints and vision problems. He has suspected	

Medicine Details:

METFORMIN HYDROCHLORIDE (Other drug) **Reason:**
2.0 Gram
Batch: **Started:** **Stopped:**

DAONIL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

GLUCOBAY (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 215586 **Seq:** 1 **Gender:** M
Reported: 06/02/2006 **Weight:** 150.00
Hospitalisation: **Age:** 47Y
Onset Date: 18/01/2006 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia		The patient developed neutropenia.	Ceased Gabapentin, Pantoprazole, Ranitidine, Gliclazide and Amitriptyline.

Medicine Details:

PARACETAMOL (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
METOPROLOL TARTRATE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
FRUSEMIDE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
CHLORVESCENT (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/02/2006	Eosinophils	<0.4	12/01/2006	0.7	
06/02/2006	Eosinophils	<0.4	13/01/2006	0.7	
06/02/2006	Eosinophils	<0.4	15/01/2006	0.8	
06/02/2006	Eosinophils	<0.4	14/01/2006	0.9	
06/02/2006	Eosinophils	<0.4	18/01/2006	0.3	
06/02/2006	Eosinophils	<0.4	20/01/2006	0.6	
06/02/2006	Eosinophils	<0.4	19/01/2006	0.5	

Additional Information:

Patient also taking Vitamin C, Zinc, Enoxaparin, Temazepam, Yakult, Iperamide, Omeprazole and Morphine.

Report Detail

Case Number: 215586 **Seq:** 1
Reported: 06/02/2006
Hospitalisation:
Onset Date: 18/01/2006
Outcome: Not yet recovered

Gender: M
Weight: 150.00
Age: 47Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

NILSTAT (Other drug)			**Reason:**	
Batch:	**Started:**		**Stopped:**	
PANTOPRAZOLE (Suspected)			**Reason:**	
	80.0 Milligram	Daily		
Batch:	**Started:** 12/12/2005		**Stopped:**	
GABAPENTIN (Suspected)			**Reason:**	
	300.0 Milligram	Daily	Oral	
Batch:	**Started:** 13/01/2006		**Stopped:** 18/01/2006	
GLIPIZIDE (Suspected)			**Reason:**	
Tablet	5.0 Milligram	Daily	Oral	
Batch:	**Started:** 16/01/2006		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/02/2006	Eosinophils	<0.4	17/01/2006	0.3	
06/02/2006	Eosinophils	<0.4	16/01/2006	0.8	
06/02/2006	Neutrophils	2-7.5	12/01/2006	5.1	
06/02/2006	Neutrophils	2-7.5	13/01/2006	4.5	
06/02/2006	Neutrophils	2-7.5	14/01/2006	3.2	
06/02/2006	Neutrophils	2-7.5	15/01/2006	2.4	
06/02/2006	Neutrophils	2-7.5	16/01/2006	1.5	

Additional Information:

Patient also taking Vitamin C, Zinc, Enoxaparin, Temazepam, Yakult, Iperamide, Omeprazole and Morphine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 215586 **Seq:** 1
Reported: 06/02/2006
Hospitalisation:
Onset Date: 18/01/2006
Outcome: Not yet recovered

Gender: M
Weight: 150.00
Age: 47Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:** Specific disorders of sleep	
	75.0 Milligram		Daily	Oral
Batch:	**Started:** 14/01/2006			**Stopped:** 18/01/2006
OMEPRAZOLE (Suspected)			**Reason:**	
Injection	80.0 Milligram		Daily	Intravenous
Batch:	**Started:** 16/01/2006			**Stopped:** 18/01/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/02/2006	Neutrophils	2-7.5	17/01/2006	0.9	
06/02/2006	Neutrophils	2-7.5	18/01/2006	0.3	
06/02/2006	Neutrophils	2-7.5	19/01/2006	0.9	
06/02/2006	Neutrophils	2-7.5	20/01/2006	1.3	

Additional Information:

Patient also taking Vitamin C, Zinc, Enoxaparin, Temazepam, Yakult, Iperamide, Omeprazole and Morphine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 216006 **Seq:** 1
Reported: 20/02/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 17/07/2005
Outcome: Recovered

Gender: M
Weight: 70.00
Age: 85Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation Dry mouth Hangover		The patient experienced constipation, had a very parched, dry mouth, and a hangover feeling	

Medicine Details:

TEMAZE (Other drug) **Reason:** Specific disorders of sleep
10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AROPAX (Other drug) **Reason:** Depression
20.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

DIAZEPAM (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ENDEP (Suspected) **Reason:** Specific disorders of sleep
Tablet 12.5 Milligram Daily Oral
Batch: **Started:** 15/07/2005 **Stopped:** 20/07/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 216655 **Seq:** 1 **Gender:** F
Reported: 23/03/2006 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 82Y
Onset Date: 26/12/2005 **DOB:**
Outcome: Recovered 28/12/2005 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Electrocardiogram QT prolonged	Caused or prolonged inpatient hospitalisation	Patient experienced prolonged QT, bradycardia and polymorphic ventricular tachycardia (Torsades), collapse.	
Circulatory collapse	Caused or prolonged inpatient hospitalisation		
Torsade de pointes	Caused or prolonged inpatient hospitalisation		

Medicine Details:

SOTALOL HYDROCHLORIDE (Suspected) **Reason:** Otr&nos disord of heart rhythm
Tablet 320.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 27/12/2005

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 05/12/2005 **Stopped:** 27/12/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 216789 **Seq:** 1
Reported: 29/03/2006
Hospitalisation: Admitted to hospital
Onset Date: 01/09/2005
Outcome: Not yet recovered

Gender: M
Weight: 106.00
Age: 74
DOB: 26/04/1931
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hot flush	Caused or prolonged inpatient hospitalisation	Patient presented with hot fluhes, tingling in the legs, falls racing heart, shortness of breath, dyskinetic movements, developed postural hypotension	Serotoninergic
Dyskinesia	Caused or prolonged inpatient hospitalisation		
Dyspnoea	Caused or prolonged inpatient hospitalisation		

Medicine Details:

SIMVASTATIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ASPIRIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

OMEPRAZOLE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ATENOLOL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 216789 **Seq:** 1
Reported: 29/03/2006
Hospitalisation: Admitted to hospital
Onset Date: 01/09/2005
Outcome: Not yet recovered

Gender: M
Weight: 106.00
Age: 74
DOB: 26/04/1931
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension	Caused or prolonged inpatient hospitalisation		
Paraesthesia	Caused or prolonged inpatient hospitalisation		
Tachycardia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AMIODARONE HYDROCHLORIDE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ALLOPURINOL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SERTRALINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 217062 **Seq:** 1 **Gender:** M
Reported: 06/04/2006 **Weight:** 80.00
Hospitalisation: **Age:** 99u
Onset Date: 18/02/2006 **DOB:**
Outcome: Recovered 20/02/2006 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Agitation Confusional state Hypomania		Signs and symptoms of suspected serotonin syndrome, as diagnosed by emergency room doctor. Patient quite agitated, confused and hypomanic.	Tramal ceased, replaced with Oxycontin.

Medicine Details:

ENDEP (Interaction) **Reason:** Pain
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

PAXTINE (Interaction) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

TRAMAL SR (Interaction) **Reason:** Pain
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 23/02/2006 0

TRAMAL (Interaction) **Reason:** Pain
100.0 Milligram Daily
Batch: **Started:** **Stopped:** 28/02/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 217062 **Seq:** 1 **Gender:** M
Reported: 06/04/2006 **Weight:** 80.00
Hospitalisation: **Age:** 99u
Onset Date: 18/02/2006 **DOB:**
Outcome: Recovered 20/02/2006 **Causality:** Causality possible

Reaction Detail

Medicine Details:

CELEBREX (Other drug)			**Reason:**	Pain	
Capsule	200.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	0
CLOFEN (Other drug)			**Reason:**	Pain	
Tablet	50.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 217554 **Seq:** 1 **Gender:** M
Reported: 20/04/2006 **Weight:** 105.00
Hospitalisation: Required a visit to the doctor **Age:** 72
Onset Date: **DOB:** 12/11/1933
Outcome: Recovered 04/04/2006 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cellulitis Rash		Patient developed cellulitis left arm picc site, rash inner aspect right arm, rash on both legs and lower back	Amitripryline ceased and iv Ceftriaxone given 2gm. The next day rash more evident, ceftriaxone ceased, the next day rash completely gone. Recommenced on Amitripryline

Medicine Details:

CIPROFLOXACIN (Other drug) **Reason:**
1.0 Gram
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 01/04/2006

CEFTRIAXONE (Suspected) **Reason:**
Injection 2.0 Megabecquerel Daily Intravenous
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 219210 **Seq:** 1
Reported: 19/06/2006
Hospitalisation: Admitted to hospital
Onset Date: 10/12/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 47
DOB: 21/06/1958
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Serotonin syndrome - myoclonic jerks, shivering, sweating as a result of combination Tramadol, Amitriptyline and narcotics	Tramadol ceased.
Chills	Caused or prolonged inpatient hospitalisation		
Hyperhidrosis	Caused or prolonged inpatient hospitalisation		

Medicine Details:

PARACETAMOL (Other drug) **Reason:** Pain
Batch: **Started:** **Stopped:**

METFORMIN HYDROCHLORIDE (Other drug) **Reason:** Diabetes mellitus
Batch: **Started:** **Stopped:**

GLICLAZIDE (Other drug) **Reason:** Diabetes mellitus
Batch: **Started:** **Stopped:**

AZATHIOPRINE (Other drug) **Reason:** Regional enteritis
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was on Prednisone for Crohns disease, this caused increased weight and sugar levels. Prednisone has been substituted with Azathioprine.

Report Detail

Case Number: 219210 **Seq:** 1
Reported: 19/06/2006
Hospitalisation: Admitted to hospital
Onset Date: 10/12/2005
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 47
DOB: 21/06/1958
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myoclonus	Caused or prolonged inpatient hospitalisation		

Medicine Details:

OXYBUTYNIN HYDROCHLORIDE (Other drug) **Reason:** Incontinence of urine
Batch: **Started:** **Stopped:**

TRAMADOL HYDROCHLORIDE (Suspected) **Reason:** Other disturbance of sensation
Tablet 50.0 Milligram As necessary Oral
Batch: **Started:** 10/12/2004 **Stopped:** 11/12/2005

OXYCODONE HYDROCHLORIDE (Suspected) **Reason:** Pain
Tablet 180.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 10/12/2004 **Stopped:**

Laboratory Investigations:

Additional Information:

Patient was on Prednisone for Crohns disease, this caused increased weight and sugar levels. Prednisone has been substituted with Azathioprine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 219724 **Seq:** 1
Reported: 05/07/2006
Hospitalisation: Admitted to hospital
Onset Date: 01/03/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 66
DOB: 10/01/1940
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular	Caused or prolonged inpatient hospitalisation	Patient presented with a maculopapular rash involving the whole skin and mucous membrane with exfoliation. Started as itch only approximately 3.5 months ago then developed rash. Admitted with hypoglycaemia, rash and renal failure.	All medications ceased.
Renal impairment	Caused or prolonged inpatient hospitalisation	suspected renal failure	

Medicine Details:

BACTRIM DS (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified Weekly Oral
Batch: **Started:** **Stopped:** 15/06/2006

AVANDIA (Suspected) **Reason:** Diabetes mellitus
Tablet, film coated 4.0 Milligram Daily Oral
Batch: **Started:** 01/03/2006 **Stopped:** 15/06/2006

AVAPRO (Suspected) **Reason:** Essential benign hypertension
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 15/06/2006

DIAMICRON MR (Suspected) **Reason:** Diabetes mellitus
Tablet 120.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 15/06/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 219724 **Seq:** 1
Reported: 05/07/2006
Hospitalisation: Admitted to hospital
Onset Date: 01/03/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 66
DOB: 10/01/1940
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus	Caused or prolonged inpatient hospitalisation		
Skin exfoliation	Caused or prolonged inpatient hospitalisation		

Medicine Details:

NORVASC (Suspected) **Reason:** Essential benign hypertension
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 15/06/2006

AZATHIOPRINE (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:** 15/06/2006

NOTEN (Suspected) **Reason:** Essential benign hypertension
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 15/06/2006

CATAPRES (Suspected) **Reason:** Essential benign hypertension
200.0 Microgram Daily
Batch: **Started:** **Stopped:** 15/06/2006

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 219724 **Seq:** 1
Reported: 05/07/2006
Hospitalisation: Admitted to hospital
Onset Date: 01/03/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 66
DOB: 10/01/1940
Causality: Causality possible

Reaction Detail

Medicine Details:

ENDEP (Suspected) **Reason:** Depression

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 15/06/2006

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 220567 **Seq:** 1 **Gender:** M
Reported: 04/08/2006 **Weight:** 0.00
Hospitalisation: **Age:** 50Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vision blurred Dry mouth Liver function test abnormal		Blured vision. dry mouth, liver function tests	

Medicine Details:

Lyrica (Suspected) **Reason:** Pain
Capsule Oral
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
04/08/2006	ALT = SGPT				Jul 2006: ALT - 80.
04/08/2006	AST = SGOT				Jul 2006: AST - 80.
04/08/2006	GGT = SGGT =				Jul 2006: GGT - 200.
04/08/2006	Other data				Mar 2006 - liver function test normal. Jul 2006 - liver function test raised.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 220654 **Seq:** 1 **Gender:** F
Reported: 08/08/2006 **Weight:** 68.00
Hospitalisation: Admitted to hospital **Age:** 26
Onset Date: 05/07/2006 **DOB:** 15/08/1979
Outcome: Recovered 21/07/2006 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral	Caused or prolonged inpatient hospitalisation	Peripheral oedema of legs and arms, pharyngeal oedema, malaise, anorexia, vomiting and poor sleep	
Anorexia	Caused or prolonged inpatient hospitalisation		
Malaise	Caused or prolonged inpatient hospitalisation		

Medicine Details:

KALMA (Other drug) **Reason:** Anxiety neurosis
4.0 Milligram
Batch: **Started:** 01/07/2006 **Stopped:**

FLIXOTIDE (Other drug) **Reason:** Asthma
Inhalation 2.0 Dose Unspecified Inhalation
Batch: **Started:** 18/04/2006 **Stopped:**

BETNOVATE (Other drug) **Reason:** Otr ecz&der due otr&unsp cause
Batch: **Started:** 15/03/2006 **Stopped:**

ENDEP (Suspected) **Reason:** Anxiety neurosis
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 02/07/2006 **Stopped:** 20/07/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 220654 **Seq:** 1 **Gender:** F
Reported: 08/08/2006 **Weight:** 68.00
Hospitalisation: Admitted to hospital **Age:** 26
Onset Date: 05/07/2006 **DOB:** 15/08/1979
Outcome: Recovered 21/07/2006 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pharyngeal oedema	Caused or prolonged inpatient hospitalisation		
Poor quality sleep	Caused or prolonged inpatient hospitalisation		
Vomiting	Caused or prolonged inpatient hospitalisation		

Medicine Details:

KALMA (Other drug) **Reason:** Anxiety neurosis
4.0 Milligram
Batch: **Started:** 01/07/2006 **Stopped:**

FLIXOTIDE (Other drug) **Reason:** Asthma
Inhalation 2.0 Dose Unspecified Inhalation
Batch: **Started:** 18/04/2006 **Stopped:**

BETNOVATE (Other drug) **Reason:** Otr ecz&der due otr&unsp cause
Batch: **Started:** 15/03/2006 **Stopped:**

ENDEP (Suspected) **Reason:** Anxiety neurosis
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 02/07/2006 **Stopped:** 20/07/2006

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 220864 **Seq:** 1 **Gender:** M
Reported: 17/08/2006 **Weight:** 0.00
Hospitalisation: **Age:** 59Y
Onset Date: 10/07/2006 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysphagia	Caused or prolonged inpatient hospitalisation	Barium swollen (14/7): severe failure of lower oesophageal sphincter opening-not seen to open with peristalsis. Oesophagus mildly dilated and demonstrated impaired motility. High risk of reflux regurgitation. Oropharyngeal dysphagia	Ceased amitriptyline,

Medicine Details:

COLOXYL WITH SENNA (Other drug)		**Reason:**	Constipation
Batch:	**Started:**		**Stopped:**
LACTULOSE (Other drug)		**Reason:**	Constipation
Batch:	**Started:**		**Stopped:**
MOVICOL (Other drug)		**Reason:**	Constipation
Batch:	**Started:**		**Stopped:**
METOCLOPRAMIDE HYDROCHLORIDE (Other drug)		**Reason:**	Nausea and vomiting
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Tay Sachs disease:autonomic neuropathy, previous reflux symptoms when eating too quickly.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 220864 **Seq:** 1
Reported: 17/08/2006
Hospitalisation:
Onset Date: 10/07/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 05/07/2006 **Stopped:** 14/07/2006

Laboratory Investigations:

Additional Information:

Tay Sachs disease:autonomic neuropathy, previous reflux symptoms when eating too quickly.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221296 **Seq:** 1
Reported: 01/09/2006
Hospitalisation:
Onset Date: 22/04/2006
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 83
DOB: 22/05/1922
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium	Caused or prolonged inpatient hospitalisation	Delerium secondary to serotonin syndrome, also myoclonic jerking and fever	
Myoclonus	Caused or prolonged inpatient hospitalisation		
Pyrexia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

WARFARIN SODIUM (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

FRUSEMIDE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

CELECOXIB (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

DIGOXIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 221296 **Seq:** 1
Reported: 01/09/2006
Hospitalisation:
Onset Date: 22/04/2006
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 83
DOB: 22/05/1922
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation		

Medicine Details:

PANTOPRAZOLE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

SPIRONOLACTONE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

RAMIPRIL (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

TRAMADOL HYDROCHLORIDE (Suspected) **Reason:**
Capsule 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 02/05/2006

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221296 **Seq:** 1
Reported: 01/09/2006
Hospitalisation:
Onset Date: 22/04/2006
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 83
DOB: 22/05/1922
Causality: Causality possible

Reaction Detail

Medicine Details:

AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	100.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	02/05/2006
PAROXETINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	01/05/2006

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221386 **Seq:** 1 **Gender:** M
Reported: 05/09/2006 **Weight:** 0.00
Hospitalisation: **Age:** 86Y
Onset Date: 22/08/2006 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination, auditory Headache		Experienced auditory (wind blowing very loudly) and severe headache after taking endep 10mg at approx 9pm	

Medicine Details:

FISH OIL NOS (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Glucosamine Nos (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 10.0 Milligram 1 time Oral
Batch: **Started:** 21/08/2006 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previosuly prescibed an antidepressant for back pain and tingling. It gave him terrible noises in the head upon waking,

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221404 **Seq:** 1 **Gender:** F
Reported: 05/09/2006 **Weight:** 0.00
Hospitalisation: **Age:** 78
Onset Date: 19/08/2006 **DOB:** 09/05/1928
Outcome: Recovered 19/08/2006 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tongue oedema Paraesthesia		Tongue swelled and lip tingly	Endone ceased.

Medicine Details:

ISCOVER (Other drug) **Reason:**
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

MICARDIS (Other drug) **Reason:**
80.0 Milligram Daily
Batch: **Started:** **Stopped:**

NORVASC (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

ASPIRIN (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 221404 **Seq:** 1 **Gender:** F
Reported: 05/09/2006 **Weight:** 0.00
Hospitalisation: **Age:** 78
Onset Date: 19/08/2006 **DOB:** 09/05/1928
Outcome: Recovered 19/08/2006 **Causality:** Causality possible

Reaction Detail

Medicine Details:

ENDONE (Suspected)			**Reason:**	Pain	
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**	18/08/2006		**Stopped:**	19/08/2006
ENDEP (Suspected)			**Reason:**	Pain	
Tablet	25.0	Milligram	Daily		Oral
Batch:	**Started:**	15/08/2006		**Stopped:**	19/08/2006

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221596 **Seq:** 1
Reported: 13/09/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 19/08/2006
Outcome: Unknown

Gender: F
Weight: 80.00
Age: 36
DOB: 03/01/1970
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Drug ineffective Headache Rash		Itching rash (worse in 12 hours following dose) and worsening of headache.	

Medicine Details:

LOGYNON ED (Other drug) **Reason:** Contraception
Batch: **Started:** **Stopped:**

TRAMAL SR (Other drug) **Reason:** Headache
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 27/06/2006 **Stopped:**

ENDEP (Suspected) **Reason:** Headache
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 18/08/2006 **Stopped:** 24/08/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Hayfever and eczema in history.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 221995 **Seq:** 1 **Gender:** M
Reported: 27/09/2006 **Weight:** 0.00
Hospitalisation: Required a visit to the doctor **Age:** 54
Onset Date: 02/06/2006 **DOB:** 23/06/1951
Outcome: Recovered 11/09/2006 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Abnormal behaviour Anger Anxiety Dizziness Headache		Agitation, anxiety, headaches, dizziness, strange behaviour and anger.	Endep.

Medicine Details:

PARIET (Other drug) **Reason:** Other diseases of esophagus
Batch: **Started:** **Stopped:**

LIPITOR (Other drug) **Reason:** Othr&unspec metabolic diseases
Batch: **Started:** **Stopped:**

PANADEINE (Other drug) **Reason:** Pain
Batch: **Started:** **Stopped:**

MOBIC (Other drug) **Reason:** Osteoarthritis
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 221995 **Seq:** 1 **Gender:** M
Reported: 27/09/2006 **Weight:** 0.00
Hospitalisation: Required a visit to the doctor **Age:** 54
Onset Date: 02/06/2006 **DOB:** 23/06/1951
Outcome: Recovered 11/09/2006 **Causality:** Causality possible

Reaction Detail

Medicine Details:

CIPRAMIL (Suspected)			**Reason:** Depression	
Tablet	60.0 Milligram		Daily	Oral
Batch:	**Started:** 09/10/2005		**Stopped:**	
ENDEP (Suspected)			**Reason:** Headache	
Tablet	12.5 Milligram		Daily	Oral
Batch:	**Started:** 02/06/2006		**Stopped:** 08/09/2006	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 223697 **Seq:** 1
Reported: 01/12/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 25/10/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 83
DOB: 02/04/1923
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting Liver function test abnormal Nausea		Patient complained of vomiting and nausea, elevated LFTs.	Suspected agents withdrawn, supportive care.

Medicine Details:

CLOPIDOGREL (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
SPIRIVA (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
Symbicort Turbuhaler (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PROGOUT (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
01/12/2006	ALT = SGPT	1-40	23/10/2006	59	
01/12/2006	ALT = SGPT	1-40	25/10/2006	1628	
01/12/2006	ALT = SGPT	1-40	26/10/2006	2090	
01/12/2006	AST = SGOT	1-35	23/10/2006	45	
01/12/2006	AST = SGOT	1-35	26/10/2006	1897	
01/12/2006	AST = SGOT	1-35	25/10/2006	2245	
01/12/2006	Bilirubin	<20	23/10/2006	19	

Additional Information:

Patient was also taking Solprin, Quit-X, Mult-B Forte, Novomix-30.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 223697 **Seq:** 1
Reported: 01/12/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 25/10/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 83
DOB: 02/04/1923
Causality: Causality possible

Reaction Detail

Medicine Details:

FRUSEMIDE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
KENACOMB (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PROTAPHANE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
KARVEZIDE 300/12.5 (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
01/12/2006	Bilirubin	<20	25/10/2006	39	
01/12/2006	Bilirubin	<20	26/10/2006	38	
01/12/2006	GGT = SGGT =	<55	23/10/2006	53	
01/12/2006	GGT = SGGT =	<55	25/10/2006	67	
01/12/2006	GGT = SGGT =	<55	26/10/2006	94	
01/12/2006	SAP = ALP		23/10/2006	63	35-115
01/12/2006	SAP = ALP		25/10/2006	82	35-115

Additional Information:

Patient was also taking Solprin, Quit-X, Mult-B Forte, Novomix-30.

Report Detail

Case Number: 223697 **Seq:** 1
Reported: 01/12/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 25/10/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 83
DOB: 02/04/1923
Causality: Causality possible

Reaction Detail

Medicine Details:

Profloxin (Suspected)			**Reason:**	Pulmonary collapse	
Tablet	1.0	Gram	Daily	Oral	
Batch:	**Started:**	12/10/2006		**Stopped:**	17/10/2006 0
RULIDE (Suspected)			**Reason:**	Pulmonary collapse	
Tablet	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	12/10/2006		**Stopped:**	16/10/2006
CLEXANE (Suspected)			**Reason:**		
Injection	30.0	Milligram	Daily		
Batch:	**Started:**	14/10/2006		**Stopped:**	18/10/2006
CLEXANE (Suspected)			**Reason:**		
Injection	60.0	Milligram	Daily		
Batch:	**Started:**	18/10/2006		**Stopped:**	24/10/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
01/12/2006	SAP = ALP		26/10/2006	136	35-115

Additional Information:

Patient was also taking Solprin, Quit-X, Mult-B Forte, Novomix-30.

Report Detail

Case Number: 223697 **Seq:** 1
Reported: 01/12/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 25/10/2006
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 83
DOB: 02/04/1923
Causality: Causality possible

Reaction Detail

Medicine Details:

ALDACTONE (Suspected)			**Reason:**			
Tablet	25.0	Milligram	Daily		Oral	
Batch:	**Started:**	19/10/2006		**Stopped:**	25/10/2006	
ENDEP (Suspected)			**Reason:**			
Tablet	25.0	Milligram	Daily		Oral	
Batch:	**Started:**	21/10/2006		**Stopped:**	22/10/2006	0
ENDEP (Suspected)			**Reason:**			
Tablet	10.0	Milligram	Daily		Oral	
Batch:	**Started:**	23/10/2006		**Stopped:**	25/10/2006	

Laboratory Investigations:

Additional Information:
Patient was also taking Solprin, Quit-X, Mult-B Forte, Novomix-30.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 223841 **Seq:** 1 **Gender:** F
Reported: 06/12/2006 **Weight:** 74.00
Hospitalisation: **Age:** 27
Onset Date: 05/09/2006 **DOB:** 10/07/1979
Outcome: Recovered 02/10/2006 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor	Life	Tremor, confusion, flushed twitches, nausea, myalgia, abnormal liver function tests	Ceased Tramal.
Blood creatine phosphokinase in	Life		
Confusional state	Life		
Flushing	Life		
Liver function test abnormal	Life		
Muscle twitching	Life		
Myalgia	Life		
Nausea	Life		
Serotonin syndrome	Life		

Medicine Details:

TRAMAL (Interaction) **Reason:** Pain in joint
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** 18/07/2006 **Stopped:** 05/09/2006 Tramal ceased.

ENDEP (Interaction) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 18/07/2006 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/12/2006	Creatine		05/09/2006	1300	High ALTand LDH.
06/12/2006	Creatine		02/11/2006	94	High ALTand LDH.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 224202 **Seq:** 1
Reported: 19/12/2006
Hospitalisation: Admitted to hospital
Onset Date: 26/09/2006
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 82
DOB: 04/09/1924
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Ventricular tachycardia	Caused or prolonged inpatient hospitalisation	Tachycardia ventricular and QT interval	Ceased amitriptyline and
Electrocardiogram QT prolonged	Caused or prolonged inpatient hospitalisation		

Medicine Details:

ENDEP (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

LOVAN (Interaction) **Reason:**
Capsule 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ASPIRIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

INDAPAMIDE HEMIHYDRATE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 224202 **Seq:** 1
Reported: 19/12/2006
Hospitalisation: Admitted to hospital
Onset Date: 26/09/2006
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 82
DOB: 04/09/1924
Causality: Causality possible

Reaction Detail

Medicine Details:

PANTOPRAZOLE (Other drug)		**Reason:**	
Batch:	**Started:**	**Stopped:**	
LISINOPRIL (Other drug)		**Reason:**	
Batch:	**Started:**	**Stopped:**	0
DIAZEPAM (Other drug)		**Reason:**	
Batch:	**Started:**	**Stopped:**	
GRAPEFRUIT JUICE (Other drug)		**Reason:**	
Batch:	**Started:**	**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 224404 **Seq:** 1
Reported: 08/01/2007
Hospitalisation: Required a visit to the doctor
Onset Date: 07/10/2006
Outcome: Recovered

Gender: F
Weight: 0.00
Age: 18Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Polyuria		Increased freqency of urination.	Ceased.

Medicine Details:

Microgynon 50 ED (Other drug) **Reason:**

Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:** Disturbance of sleep

Tablet 10.0 Milligram Daily Oral

Batch: **Started:** 05/10/2006 **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 225117 **Seq:** 1 **Gender:** M
Reported: 27/01/2007 **Weight:** 80.00
Hospitalisation: **Age:** 72
Onset Date: 23/01/2007 **DOB:** 04/10/1934
Outcome: Unknown 27/01/2007 **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest discomfort	Incapacity/disability	Tightness of chest, complaining of hard to breathe, lasts for approx 2 hours post dose is dose related	diazepam sometimes helps, but not lately
Dyspnoea	Incapacity/disability		

Medicine Details:

ENDEP (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** 21/01/2007 **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 225813 **Seq:** 1
Reported: 14/02/2007
Hospitalisation: Required a specialist consultation
Onset Date: 30/01/2007
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 50Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fall Hypersomnia Muscular weakness Tremor		Falling down and sleeping for 20 hours a day; fine tremor in both arms, more pronounced in the left; a few days later this was a whole body tremor, with weakness of arms.	

Medicine Details:

NEURONTIN (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Batch: **Started:** **Stopped:**

ZOLOFT (Suspected) **Reason:**
Batch: **Started:** **Stopped:**

Lyrica (Suspected) **Reason:**
300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 225900 **Seq:** 1 **Gender:** M
Reported: 16/02/2007 **Weight:** 0.00
Hospitalisation: **Age:** 38
Onset Date: **DOB:** 04/03/1968
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		LFT's rasied.	Decreased dose of Carbamazepine. Changed from Methadone to Hydromorphone and Fentanyl

Medicine Details:

PARACETAMOL (Suspected)			**Reason:**		
Tablet	2000.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	
CARBAMAZEPINE (Suspected)			**Reason:**		
Tablet	400.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	
TRAMADOL HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	200.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**		
Tablet	50.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	15/10/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
16/02/2007	ALT = SGPT		10/10/2006	38	
16/02/2007	ALT = SGPT		12/10/2006	317	
16/02/2007	ALT = SGPT		17/10/2006	96	
16/02/2007	ALT = SGPT		16/10/2006	125	
16/02/2007	AST = SGOT		10/10/2006	19	
16/02/2007	AST = SGOT		12/10/2006	238	
16/02/2007	AST = SGOT		17/10/2006	18	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 225900 **Seq:** 1 **Gender:** M
Reported: 16/02/2007 **Weight:** 0.00
Hospitalisation: **Age:** 38
Onset Date: **DOB:** 04/03/1968
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

METHADONE HYDROCHLORIDE (Suspected) **Reason:**
180.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 15/10/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
16/02/2007	AST = SGOT		16/10/2006	23	
16/02/2007	GGT = SGGT =		10/10/2006	226	
16/02/2007	GGT = SGGT =		12/10/2006	663	
16/02/2007	GGT = SGGT =		16/10/2006	497	
16/02/2007	GGT = SGGT =		17/10/2006	468	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 225921 **Seq:** 1 **Gender:** M
Reported: 19/02/2007 **Weight:** 0.00
Hospitalisation: **Age:** 25Y
Onset Date: 15/06/2004 **DOB:**
Outcome: Recovered 15/04/2005 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Patient described seizures as "involuntary muscle movements and jerks of legs and arms, like intense muscle spasm and cramp". GP witnessed one episode and described it as a seizure, no loss of conciousness	Epilim. All drugs ceased by

Medicine Details:

DI-GESIC (Other drug) **Reason:**
8.0 Dose Unspecified
Batch: **Started:** **Stopped:**

VIOXX (Other drug) **Reason:**
25.0 Milligram
Batch: **Started:** **Stopped:** 15/10/2004

VALIUM (Other drug) **Reason:** Pain
8.0 Milligram Daily
Batch: **Started:** **Stopped:** 15/02/2005

TRAMADOL HYDROCHLORIDE (Suspected) **Reason:** Pain
200.0 Milligram Daily
Batch: **Started:** **Stopped:** 15/02/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 225921 **Seq:** 1 **Gender:** M
Reported: 19/02/2007 **Weight:** 0.00
Hospitalisation: **Age:** 25Y
Onset Date: 15/06/2004 **DOB:**
Outcome: Recovered 15/04/2005 **Causality:** Causality possible

Reaction Detail

Medicine Details:

EFEXOR (Suspected)			**Reason:**	Pain
	150.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	15/03/2005
ENDEP (Suspected)			**Reason:**	Pain
	25.0	Milligram	Daily	
Batch:	**Started:**		**Stopped:**	15/02/2005

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 225967 **Seq:** 1
Reported: 20/02/2007
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 42
DOB: 10/08/1964
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Atrioventricular block	Life	Patient experienced syncope/presyncope and AV block during the use of	

Medicine Details:

ROACCUTANE (Suspected) **Reason:**
Oral
Batch: **Started:** 15/08/2006 **Stopped:** 16/01/2007 0

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
10.0 Milligram Oral
Batch: **Started:** 15/06/2003 **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 226134 **Seq:** 1 **Gender:** M
Reported: 27/02/2007 **Weight:** 60.00
Hospitalisation: **Age:** 64
Onset Date: 02/02/2007 **DOB:** 24/08/1942
Outcome: Recovered 05/02/2007 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Serotonin Syndrome	Supportive treatment

Medicine Details:

CIPRAMIL (Interaction) **Reason:**
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 19/01/2007 **Stopped:**

SINEMET (Interaction) **Reason:** Paralysis agitans
Tablet 750.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ENDEP (Interaction) **Reason:**
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 02/01/2007

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 226387 **Seq:** 1
Reported: 06/03/2007
Hospitalisation: Required a visit to the doctor
Onset Date: 15/12/2006
Outcome: Not yet recovered

Gender: F
Weight: 79.00
Age: 60
DOB: 15/10/1946
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hirsutism		Hirsutism - facial, arms, abdomen, legs.	Reduced dose Cyclosporin.

Medicine Details:

CYCLOSPORIN-A (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

ATACAND (Suspected) **Reason:**
Tablet 32.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ISOPTIN (Suspected) **Reason:**
Tablet 240.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 226387 **Seq:** 1
Reported: 06/03/2007
Hospitalisation: Required a visit to the doctor
Onset Date: 15/12/2006
Outcome: Not yet recovered

Gender: F
Weight: 79.00
Age: 60
DOB: 15/10/1946
Causality: Causality possible

Reaction Detail

Medicine Details:

TEMAZEPAM (Suspected)			**Reason:**		
Tablet	10.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 226975 **Seq:** 1 **Gender:** M
Reported: 26/03/2007 **Weight:** 0.00
Hospitalisation: **Age:** 39Y
Onset Date: **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Brugada syndrome Electrocardiogram QT prolonged Sudden death		brugada syndrome, prolonged QT interval, sudden death	

Medicine Details:

HALOPERIDOL (Suspected) **Reason:**
30.0 Milligram Daily
Batch: **Started:** **Stopped:** 0

AMITRIPTYLINE HYDROCHLORIDE (Suspected) **Reason:**
200.0 Milligram Daily
Batch: **Started:** **Stopped:** 0

SODIUM VALPROATE (Suspected) **Reason:**
1.0 Gram Daily
Batch: **Started:** **Stopped:** 0

LITHIUM CARBONATE (Suspected) **Reason:**
400.0 Milligram Daily
Batch: **Started:** **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
26/03/2007	Electrocardiograph				Electrocardiogram ST-T segment abnormal, ECG QT prolonged, Bundle branch block right, ECG ST segment elevation, ECG T wave inversion
26/03/2007	Serum drug level				Amitriptyline level was 1.36 mmol/L (NR 0.27 - 0.72), nortriptyline was 3.92 (NR 0.19 - 0.57) and Lithium level was 0.2 (NR 0.5 - 1.0)
26/03/2007	Troponin				Troponin I was undetectable.

Additional Information:

The authors considered the QT prolongation related to treatment with amitriptyline and haloperidol. The authors comment that Brugada syndrome "has been associated with mutations in the sodium channel, SCN5A and may be unmasked by medications with sodium channel blocking activity". Furthermore, "in this case, all four medications may inactivate the cardiac sodium channels".
Roberts-Thomson KC et al. Drug-induced Brugada syndrome with ST-T wave alternans and long QT. Intern Med J 2007; 37(3):199-200.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 228219 **Seq:** 1 **Gender:** M
Reported: 04/05/2007 **Weight:** 0.00
Hospitalisation: **Age:** 72
Onset Date: 12/02/2007 **DOB:** 05/04/1934
Outcome: Recovered 13/03/2007 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Scrotal oedema	Caused or prolonged inpatient hospitalisation	Patient developed a maculopapular eruption, had a generalised itchy rash, eosinophilia and peripheral hand and scrotal oedema	Celestone wet wraps, prednisolone reducing dose (50mg and tapered), cessation of drugs
Drug eruption	Caused or prolonged inpatient hospitalisation		
Eosinophilia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

FOSAMAX (Other drug) **Reason:**
Tablet 70.0 Milligram Oral
Batch: **Started:** 11/08/2006 **Stopped:** 15/09/2006

DIAMICRON MR (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

RENITEC (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

LASIX (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
04/05/2007	Skin biopsy				Skin biopsy consistent with drug eruption.

Additional Information:

Report Detail

Case Number: 228219 **Seq:** 1 **Gender:** M
Reported: 04/05/2007 **Weight:** 0.00
Hospitalisation: **Age:** 72
Onset Date: 12/02/2007 **DOB:** 05/04/1934
Outcome: Recovered 13/03/2007 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral	Caused or prolonged inpatient hospitalisation		
Pruritus	Caused or prolonged inpatient hospitalisation		
Rash maculo-papular	Caused or prolonged inpatient hospitalisation		

Medicine Details:

CALTRATE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

LIPITOR (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

ENDEP (Suspected) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 02/02/2007 **Stopped:** 05/03/2007

TEGRETOL (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** 02/02/2007 **Stopped:** 05/03/2007

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 228438 **Seq:** 1 **Gender:** F
Reported: 11/05/2007 **Weight:** 78.00
Hospitalisation: **Age:** 80
Onset Date: 13/02/2007 **DOB:** 20/11/1926
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia	Life	Patient developed decreased neutrophil count - neutropenia and leukopenia.	Endep stopped. Monitored
Leukopenia	Life		

Medicine Details:

LIPITOR (Other drug) **Reason:**
20.0 Milligram
Batch: **Started:** **Stopped:**

CARVEDILOL HYDROCHLORIDE (Other drug) **Reason:**
50.0 Milligram
Batch: **Started:** **Stopped:**

DUROGESIC (Other drug) **Reason:**
25.0 Microgram
Batch: **Started:** **Stopped:**

KEFLEX (Other drug) **Reason:**
1.5 Gram
Batch: **Started:** 30/01/2007 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
11/05/2007	Neutrophils		08/02/2007	2.52	1.8-8.0x10^9/L
11/05/2007	Neutrophils		13/02/2007	1.47	1.8-8.0x10^9/L
11/05/2007	Neutrophils		16/02/2007	1.45	1.8-8.0x10^9/L
11/05/2007	Neutrophils		06/03/2007	1.94	1.8-8.0x10^9/L
11/05/2007	Neutrophils		02/03/2007	1.57	1.8-8.0x10^9/L
11/05/2007	Neutrophils		23/02/2007	1.19	1.8-8.0x10^9/L
11/05/2007	White blood cells		08/02/2007	4.0	4-11x10^9/L

Additional Information:

PHx: Recurrent UTIs, IHD/AMI, laminectomy, CCF, PVD, Narcolepsy, HT, sciatica.

Report Detail

Case Number: 228438 **Seq:** 1 **Gender:** F
Reported: 11/05/2007 **Weight:** 78.00
Hospitalisation: **Age:** 80
Onset Date: 13/02/2007 **DOB:** 20/11/1926
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

THYROXINE SODIUM (Other drug)			**Reason:**		
	50.0	Microgram	Daily		
Batch:	**Started:**			**Stopped:**	0
LASIX (Other drug)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	0
ENDEP (Suspected)			**Reason:**		
Tablet	50.0	Milligram	Daily	Oral	
Batch:	**Started:**	22/01/2007		**Stopped:**	28/02/2007 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
11/05/2007	White blood cells		13/02/2007	3.5	4-11x10^9/L
11/05/2007	White blood cells		16/02/2007	3.4	4-11x10^9/L
11/05/2007	White blood cells		23/02/2007	3.1	4-11x10^9/L
11/05/2007	White blood cells		02/03/2007	3.5	4-11x10^9/L
11/05/2007	White blood cells		06/03/2007	4.5	4-11x10^9/L

Additional Information:

PHx: Recurrent UTIs, IHD/AMI, laminectomy, CCF, PVD, Narcolepsy, HT, sciatica.

Report Detail

Case Number: 228703 **Seq:** 1 **Gender:** M
Reported: 21/05/2007 **Weight:** 100.00
Hospitalisation: **Age:** 64
Onset Date: 17/04/2007 **DOB:** 27/12/1942
Outcome: Recovered 19/04/2007 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
International normalised ratio inc	Caused or prolonged inpatient hospitalisation	Post extraction of several teeth patient experienced one socket re-bleeding.	Phytomenadione 2mg IV, 2 units of fresh frozen plasma. Further 2 units fresh frozen plasma and phytomenadione 10mg IV. Articane with adrenaline 1:100,000.
Haemorrhage	Caused or prolonged inpatient hospitalisation	haemorrhage post dental extraction	

Medicine Details:

WARFARIN SODIUM (Interaction) **Reason:**
5.0 Milligram
Batch: **Started:** **Stopped:**

ENDEP (Interaction) **Reason:**
20.0 Milligram Daily
Batch: **Started:** 13/04/2007 **Stopped:**

TENORMIN (Other drug) **Reason:**
50.0 Milligram
Batch: **Started:** **Stopped:**

COVERSYL (Other drug) **Reason:**
10.0 Milligram
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 228703 **Seq:** 1 **Gender:** M
Reported: 21/05/2007 **Weight:** 100.00
Hospitalisation: **Age:** 64
Onset Date: 17/04/2007 **DOB:** 27/12/1942
Outcome: Recovered 19/04/2007 **Causality:** Causality possible

Reaction Detail

Medicine Details:

PARACETAMOL (Other drug)			**Reason:**	
	1.0 Gram			
Batch:	**Started:** 17/04/2007		**Stopped:**	
AMOXYCILLIN SODIUM (Other drug)			**Reason:**	
	2.0 Gram			
Batch:	**Started:** 17/04/2007		**Stopped:**	17/04/2007
ENDEP (Other drug)			**Reason:**	
	10.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	13/04/2007

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 230026 **Seq:** 1 **Gender:** F
Reported: 21/06/2007 **Weight:** 0.00
Hospitalisation: Hospitalisation prolonged **Age:** 75Y
Onset Date: 03/04/2007 **DOB:**
Outcome: Recovered 05/04/2007 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Cold sweat Dizziness Dry mouth Hypotension Nausea Vomiting		Patient experienced Serotonin Syndrome, dizziness, vomiting, nausea, dry mouth, hypotension, clammy	Ceased Tramadol changed to Oxycontin, IV fluids, metoclopramide

Medicine Details:

TRAMAL SR (Interaction) **Reason:** Other disturbance of sensation
Tablet 200.0 Milligram Monthly Oral
Batch: **Started:** 02/04/2007 **Stopped:** 04/04/2007

ENDEP (Interaction) **Reason:**
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

LIPITOR (Other drug) **Reason:**
40.0 Milligram
Batch: **Started:** **Stopped:**

CLIMARA (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

correlatedwith increase in tramadol dose. possible drug interaction with TCA

Report Detail

Case Number: 230026 **Seq:** 1
Reported: 21/06/2007
Hospitalisation: Hospitalisation prolonged
Onset Date: 03/04/2007
Outcome: Recovered 05/04/2007

Gender: F
Weight: 0.00
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

CALTRATE (Other drug)		**Reason:**
	1.0 Dose Unspecified	
Batch:	**Started:**	**Stopped:**
OSTELIN 1000 (Other drug)		**Reason:**
	1.0 Dose Unspecified	
Batch:	**Started:**	**Stopped:**
METHOTREXATE (Other drug)		**Reason:**
	20.0 Milligram	
Batch:	**Started:**	**Stopped:**
FOLIC ACID (Other drug)		**Reason:**
Batch:	**Started:**	**Stopped:**

Laboratory Investigations:

Additional Information:

correlatedwith increase in tramadol dose. possible drug interaction with TCA

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 230026 **Seq:** 1 **Gender:** F
Reported: 21/06/2007 **Weight:** 0.00
Hospitalisation: Hospitalisation prolonged **Age:** 75Y
Onset Date: 03/04/2007 **DOB:**
Outcome: Recovered 05/04/2007 **Causality:** Causality possible

Reaction Detail

Medicine Details:

OMEPRAZOLE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
FOSAMAX (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PREDNISOLONE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Additional Information:

correlatedwith increase in tramadol dose. possible drug interaction with TCA

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 231273 **Seq:** 1
Reported: 27/07/2007
Hospitalisation: Admitted to hospital
Onset Date: 19/01/2007
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 63
DOB: 21/07/1943
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion	Caused or prolonged inpatient hospitalisation	Seizures for investigation. Tonic clonic	Zyban ceased. Commenced sodium valproate 200mg/bd.

Medicine Details:

TRIMETHOPRIM-SULFAMETHOXAZOLE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

MORPHINE SULPHATE (Other drug) **Reason:** Pain
Batch: **Started:** **Stopped:**

OXYCODONE HYDROCHLORIDE (Other drug) **Reason:** Pain
Batch: **Started:** **Stopped:**

CREON (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of pancreatitis and long term ETOH usage.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: AMITRIP,AMITRIPTYLINE HYDROCHLORIDE,ELAVIL,ENDEP,LAROXYL,SAROTEN,TRYPTANOL,TRYPTINE

Report Detail

Case Number: 231273 **Seq:** 1
Reported: 27/07/2007
Hospitalisation: Admitted to hospital
Onset Date: 19/01/2007
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 63
DOB: 21/07/1943
Causality: Causality possible

Reaction Detail

Medicine Details:

ALPRAZOLAM (Other drug)			**Reason:**	Anxiety neurosis	
Batch:	**Started:**			**Stopped:**	
AMITRIPTYLINE HYDROCHLORIDE (Suspected)			**Reason:**	Depression	
Tablet	75.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
Zyban (Suspected)			**Reason:**	Smoker	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	19/01/2007

Laboratory Investigations:

Additional Information:
History of pancreatitis and long term ETOH usage.