THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 74730 **Seq:** 1
Reported: 17/12/1991
Hospitalisation:
Onset Date: 11/12/1991
Outcome: Not yet recovered

Gender: F
Weight: 99.00
Age: 46
DOB: 17/11/1945
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Insomnia			Normison 1-2 df daily.
Nausea			Maxolon 3 df daily.

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:**
Oral application 1.0 Gram Daily Oral
Batch: **Started:** 01/01/1983 **Stopped:**

OROXINE (Other drug) **Reason:**
150.0 Microgram Daily
Batch: **Started:** 01/01/1984 **Stopped:**

RENITEC (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** 01/01/1988 **Stopped:**

AURORIX (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 10/12/1991 **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.

Report Detail

Case Number: 74836 **Seq:** 1
Reported: 24/12/1991
Hospitalisation:
Onset Date: 17/12/1991
Outcome: Not yet recovered

Gender: M
Weight: 90.00
Age: 62Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation Speech disorder		Abnormal behaviour. Rambling disorganised speech	

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
2.5 Gram As necessary
Batch: **Started:** 01/01/1983 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 09/12/1991 **Stopped:** 24/12/1991

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: 75051 **Seq:** 1 **Gender:** F
Reported: 13/01/1992 **Weight:** 37.00
Hospitalisation: **Age:** 71Y
Onset Date: 20/12/1991 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flatulence Asthenia Dysarthria Headache Nausea Paraesthesia		Fullness in stomach	

Medicine Details:

XANAX (Other drug)			**Reason:**	Anxiety neurosis	
Tablet	750.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
LANOXIN PG (Other drug)			**Reason:**	Congestive heart failure	
Oral application	125.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
LASIX (Other drug)			**Reason:**	Congestive heart failure	
Oral application	40.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
SLOW-K (Other drug)			**Reason:**		
Tablet	1.2	Gram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 75051 **Seq:** 1 **Gender:** F
Reported: 13/01/1992 **Weight:** 37.00
Hospitalisation: **Age:** 71Y
Onset Date: 20/12/1991 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ASPIRIN (Other drug)				**Reason:**	Acute,ill-def cerebrovascular disease w/o hyperte	
Oral application	150.0	Milligram	Daily		Oral	
Batch:	**Started:**				**Stopped:**	
AURORIX (Suspected)				**Reason:**	Depression	
Oral application	450.0	Milligram	Daily		Oral	
Batch:	**Started:**	18/12/1991			**Stopped:**	24/12/1991

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 75666 **Seq:** 1 **Gender:** F
Reported: 20/02/1992 **Weight:** 85.00
Hospitalisation: **Age:** 71Y
Onset Date: 10/02/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Prothrombin level decreased			

Medicine Details:

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

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

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

SLOW-K (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	International				14/2/92 - abnormal response warfarin inr 2.5 - 6.5.

Additional Information:

Report Detail

Case Number: 75666 **Seq:** 1 **Gender:** F
Reported: 20/02/1992 **Weight:** 85.00
Hospitalisation: **Age:** 71Y
Onset Date: 10/02/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

WARFARIN SODIUM (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**		**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**	Depression
Oral application	450.0 Milligram	Daily		Oral
Batch:	**Started:** 23/01/1992		**Stopped:**	17/02/1992

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 76498 **Seq:** 1 **Gender:** F
Reported: 08/04/1992 **Weight:** 65.00
Hospitalisation: **Age:** 30Y
Onset Date: 06/04/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation	Severe		
Insomnia	Severe		

Medicine Details:

PLENDIL ER (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 18/12/1991 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram 1 time
Batch: **Started:** 06/04/1992 **Stopped:** 06/04/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 76660 **Seq:** 1 **Gender:** F
Reported: 21/04/1992 **Weight:** 67.00
Hospitalisation: **Age:** 19Y
Onset Date: 16/04/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vertigo			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram 1 time Oral
Batch: **Started:** 16/04/1992 **Stopped:** 16/04/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 76748 **Seq:** 1 **Gender:** F
Reported: 22/04/1992 **Weight:** 60.00
Hospitalisation: **Age:** 24Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Delusion			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 10/04/1992 **Stopped:** 10/04/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 76817 **Seq:** 1 **Gender:** M
Reported: 29/04/1992 **Weight:** 65.00
Hospitalisation: **Age:** 71Y
Onset Date: 29/03/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema		Ankle and foot	

Medicine Details:

BETHANECHOL CHLORIDE (Other drug) **Reason:**
30.0 Milligram Daily
Batch: **Started:** 28/02/1992 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 13/03/1992 **Stopped:** 05/04/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 76982 **Seq:** 1 **Gender:** F
Reported: 14/05/1992 **Weight:** 60.00
Hospitalisation: **Age:** 62Y
Onset Date: 08/05/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 08/05/1992 **Stopped:** 10/05/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 77031 **Seq:** 1 **Gender:** M
Reported: 14/05/1992 **Weight:** 73.00
Hospitalisation: **Age:** 48Y
Onset Date: 26/04/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression		Became violently aggressive	

Medicine Details:

TACRINE HYDROCHLORIDE (Other drug) **Reason:**
Oral application 30.0 Milligram Daily Oral
Batch: **Started:** 01/10/1991 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:**
0.0
Batch: **Started:** 23/04/1992 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 77054 **Seq:** 1
Reported: 13/05/1992
Hospitalisation:
Onset Date: 01/05/1992
Outcome: Death, maybe drug

Gender: M
Weight: 87.00
Age: 66Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suicide attempt Drug ineffective		Patient committed suicide by gunshot	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 11/04/1992 **Stopped:** 01/05/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 77431 **Seq:** 1 **Gender:** F
Reported: 03/06/1992 **Weight:** 63.00
Hospitalisation: **Age:** 53Y
Onset Date: 23/05/1992 **DOB:**
Outcome: Recovered with sequelae **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Dysphagia Micturition disorder Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 20/05/1992 **Stopped:** 25/05/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: distress while waiting for replacement medications to take effect.

Report Detail

Case Number: 77476 **Seq:** 1 **Gender:** F
Reported: 06/06/1992 **Weight:** 65.00
Hospitalisation: **Age:** 46Y
Onset Date: 29/05/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mania Vomiting	 Severe	Hypomania	

Medicine Details:

FRISIUM (Other drug) **Reason:** Depression
10.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 77663 **Seq:** 1 **Gender:** U
Reported: 17/06/1992 **Weight:**
Hospitalisation: **Age:**
Onset Date: 13/05/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension Syncope			

Medicine Details:

ADALAT (Other drug)			**Reason:**		
Tablet	20.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
LANOXIN PG (Other drug)			**Reason:**		
Tablet	2.0	Dose Unspecified	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
ISORDIL (Other drug)			**Reason:**	Chron isch heart dis no hyper	
Tablet	20.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN
BETALOC (Other drug)			**Reason:**	Chron isch heart dis no hyper	
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: 77663 **Seq:** 1 **Gender:** U
Reported: 17/06/1992 **Weight:**
Hospitalisation: **Age:**
Onset Date: 13/05/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

ZANTAC (Other drug)			**Reason:**		
	150.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	
ZOCOR (Other drug)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
Oral application	150.0	Milligram	Daily		Oral
Batch:	**Started:**	16/04/1992		**Stopped:**	14/05/1992
MIANSERIN HYDROCHLORIDE (Suspected)			**Reason:**	Depression	
Tablet	20.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 77905 **Seq:** 1 **Gender:** F
Reported: 25/06/1992 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 24/06/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 03/06/1992 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 77906 **Seq:** 1 **Gender:** F
Reported: 25/06/1992 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 11/04/1992 **DOB:**
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence Euphoric mood Headache	 Severe	Deep sleep	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 10/04/1992 **Stopped:** 16/04/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 77939 **Seq:** 1 **Gender:** M
Reported: 29/06/1992 **Weight:** 91.00
Hospitalisation: **Age:** 35Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Dizziness Headache			

Medicine Details:

ATIVAN (Other drug) **Reason:**
Tablet 500.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Anxiety neurosis
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 20/06/1992 **Stopped:** 22/06/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 77982 **Seq:** 1 **Gender:** F
Reported: 01/07/1992 **Weight:** 64.00
Hospitalisation: **Age:** 46Y
Onset Date: 03/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			Motilium

Medicine Details:

PREMARIN (Other drug) **Reason:** Menopausal symptoms
Tablet 625.0 Microgram Daily Oral
Batch: **Started:** 01/10/1991 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 29/04/1991 **Stopped:** 20/06/1992

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: 78040 **Seq:** 1 **Gender:** F
Reported: 08/07/1992 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: 18/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia Insomnia			Normison without effect

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 18/06/1992 **Stopped:** 20/06/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 78052 **Seq:** 1 **Gender:** F
Reported: 04/07/1992 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 08/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Nightmare Psychotic disorder			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 06/06/1992 **Stopped:** 13/06/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 78121 **Seq:** 1 **Gender:** M
Reported: 09/07/1992 **Weight:**
Hospitalisation: **Age:** 46Y
Onset Date: 01/07/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 25/05/1992 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 78309 **Seq:** 1 **Gender:** F
Reported: 16/07/1992 **Weight:** 86.00
Hospitalisation: **Age:** 42Y
Onset Date: 20/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mania			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 20/06/1992 **Stopped:** 23/06/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 78317 **Seq:** 1 **Gender:** F
Reported: 16/07/1992 **Weight:** 90.00
Hospitalisation: **Age:** 36Y
Onset Date: 29/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Headache Nausea Pruritus		Sleep disturbance	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 22/06/1992 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

All effects (nausea, headache, itch) except sleep disturbance ceased to reduction in dosage.

Report Detail

Case Number: 78439 **Seq:** 1 **Gender:** F
Reported: 24/07/1992 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 07/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension			

Medicine Details:

PANADOL (Other drug)		**Reason:** Pain	
	500.0 Milligram	As necessary	
Batch:	**Started:** 01/05/1992	**Stopped:**	CONTIN
PRAMIN (Other drug)		**Reason:** Nausea and vomiting	
	10.0 Milligram	As necessary	
Batch:	**Started:** 01/05/1992	**Stopped:**	CONTIN
COLOXYL (Other drug)		**Reason:** Constipation	
	120.0 Milligram	As necessary	
Batch:	**Started:** 01/05/1992	**Stopped:**	CONTIN
SENOKOT (Other drug)		**Reason:** Constipation	
	75.0 Milligram	As necessary	
Batch:	**Started:** 01/05/1992	**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

Report Detail

Case Number: 78439 **Seq:** 1 **Gender:** F
Reported: 24/07/1992 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 07/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

TEMAZEPAM (Other drug)		**Reason:**		
	10.0 Milligram	As necessary		
Batch:	**Started:** 06/07/1992		**Stopped:**	CONTIN
SERENACE (Other drug)		**Reason:**		
Tablet	5.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/04/1992		**Stopped:**	CONTIN
BETAMIN (Other drug)		**Reason:**		
	100.0 Milligram	Daily		
Batch:	**Started:** 01/05/1992		**Stopped:**	CONTIN
NILSTAT (Other drug)		**Reason:** Moniliasis		
Solution	4.0 Millilitre	Daily	Oral	
Batch:	**Started:** 01/05/1992		**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

Report Detail

Case Number: 78439 **Seq:** 1 **Gender:** F
Reported: 24/07/1992 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 07/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

MAGMIN (Other drug)			**Reason:**		
	1.5 Gram		Daily		
Batch:	**Started:** 06/07/1992			**Stopped:**	CONTIN
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	600.0 Milligram		Total		Oral
Batch:	**Started:** 01/04/1992			**Stopped:**	09/07/1992

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

Report Detail

Case Number: 78592 **Seq:** 1 **Gender:** M
Reported: 29/07/1992 **Weight:** 75.00
Hospitalisation: **Age:** 47Y
Onset Date: 28/07/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination			
Headache			
Vertigo			
Visual disturbance			Stemetil

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 28/07/1992 **Stopped:** 28/07/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred within half an hour of taking aurorix

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 78620 **Seq:** 1 **Gender:** M
Reported: 30/07/1992 **Weight:** 70.00
Hospitalisation: **Age:** 24Y
Onset Date: 23/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chills		Chills	
Somnolence		Hypersomnolence	
Drug ineffective		Increased depressed mood	
Fatigue		Reduced energy	
Anorexia			
Apathy			
Diarrhoea			
Dizziness			

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:** Unspecified schizophrenia

Tablet 1.0 Gram Daily Oral

Batch: **Started:** 01/04/1992 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 23/06/1992 **Stopped:** 20/07/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Onset of symptoms with start of moclobemide and increased in severity when dose doubled the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 78861 **Seq:** 1 **Gender:** M
Reported: 26/08/1992 **Weight:** 60.00
Hospitalisation: **Age:** 13Y
Onset Date: 20/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 19/08/1992 **Stopped:** 20/08/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient rechallenged with reaction recurring.

Report Detail

Case Number: 78999 **Seq:** 1 **Gender:** F
Reported: 26/08/1992 **Weight:** 49.00
Hospitalisation: **Age:** 40Y
Onset Date: 13/08/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Anorexia Diarrhoea Nausea		Abdominal cramps	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 12/08/1992 **Stopped:** 26/08/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 79011 **Seq:** 1 **Gender:** F
Reported: 25/08/1992 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 25/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Dizziness		Collapsed	

Medicine Details:

THYROXINE SODIUM (Other drug) **Reason:**
Tablet 150.0 Microgram Daily Oral
Batch: **Started:** **Stopped:**

RIVOTRIL (Other drug) **Reason:**
Oral application 4.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

PROTHIADEN (Other drug) **Reason:**
Oral application 75.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 21/08/1992

FOLIC ACID (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 79011 **Seq:** 1 **Gender:** F
Reported: 25/08/1992 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 25/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 22/08/1992 **Stopped:** 25/08/1992

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 79190 **Seq:** 1 **Gender:** F
Reported: 05/09/1992 **Weight:** 54.00
Hospitalisation: **Age:** 43Y
Onset Date: 31/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 09/08/1992 **Stopped:** 17/08/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 79490 **Seq:** 1 **Gender:** M
Reported: 03/09/1992 **Weight:** 92.00
Hospitalisation: **Age:** 35Y
Onset Date: 14/08/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flatulence			

Medicine Details:

LITHICARB (Other drug) **Reason:** Depression
1.2 Gram Daily
Batch: **Started:** 01/01/1992 **Stopped:** CONTIN

TEGRETOL (Other drug) **Reason:** Depression
1.4 Gram Daily
Batch: **Started:** 01/02/1992 **Stopped:** CONTIN

CODRAL FORTE (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
20.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 14/08/1992 **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 accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 79545 **Seq:** 1 **Gender:** F
Reported: 16/09/1992 **Weight:** 59.00
Hospitalisation: **Age:** 30Y
Onset Date: 07/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertonia Agitation	 Severe	Tense muscles	

Medicine Details:

HALDOL DECANOATE (Other drug) **Reason:**
Injection 100.0 Milligram Fortnightly Intramuscular
Batch: **Started:** L TERM **Stopped:** CONTIN

HALOPERIDOL (Other drug) **Reason:**
60.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

DISIPAL (Other drug) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 17/08/1992 **Stopped:** 07/09/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 79646 **Seq:** 1 **Gender:** M
Reported: 20/09/1992 **Weight:** 109.00
Hospitalisation: **Age:** 34Y
Onset Date: 31/08/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia Dry mouth			

Medicine Details:

XANAX (Suspected) **Reason:** Anxiety neurosis
500.0 Microgram Daily
Batch: **Started:** 01/02/1992 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 26/08/1992 **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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 80097 **Seq:** 1 **Gender:** M
Reported: 29/09/1992 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 16/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dermatitis bullous		Blistery eruption on face and neck	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: 2 **Started:** 14/09/1992 **Stopped:** 18/09/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction recurred on rechallenge

Report Detail

Case Number: 80622 **Seq:** 1
Reported: 27/10/1992
Hospitalisation: Admitted to hospital
Onset Date: 31/10/1992
Outcome: Unknown

Gender: M
Weight:
Age: 53
DOB: 16/10/1939
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Hypertension Paraesthesia			

Medicine Details:

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

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

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

AURORIX (Suspected) **Reason:**
0.0
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 80686 **Seq:** 1 **Gender:** F
Reported: 04/11/1992 **Weight:** 51.00
Hospitalisation: **Age:** 66Y
Onset Date: 24/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Face oedema Gravitational oedema Rash			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 23/10/1992 **Stopped:** 27/10/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 80770 **Seq:** 1 **Gender:** U
Reported: 26/11/1992 **Weight:**
Hospitalisation: **Age:**
Onset Date: 04/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:**

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 02/10/1992 **Stopped:** 05/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 80789 **Seq:** 1 **Gender:** F
Reported: 07/10/1992 **Weight:** 54.00
Hospitalisation: **Age:** 18Y
Onset Date: 22/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 80834 **Seq:** 1 **Gender:** M
Reported: 09/11/1992 **Weight:** 93.00
Hospitalisation: **Age:** 29Y
Onset Date: 14/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness	Severe		
Headache	Severe		
Nausea	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Phobic neurosis
Oral application 150.0 Milligram Daily Oral
Batch: **Started:** 12/10/1992 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms resolved when dose decreased, when dose gradually increased to 150 mg again, symptoms recurred.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 80864 **Seq:** 1 **Gender:** F
Reported: 11/11/1992 **Weight:**
Hospitalisation: **Age:** AD
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 81001 **Seq:** 1 **Gender:** M
Reported: 14/11/1992 **Weight:** 60.00
Hospitalisation: **Age:** 41Y
Onset Date: 25/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Insomnia Nightmare	Severe		

Medicine Details:

VITAMIN NOS (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 21/08/1992 **Stopped:** 20/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 81014 **Seq:** 1 **Gender:** F
Reported: 16/11/1992 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: 09/11/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Nausea		Visual hallucinations	

Medicine Details:

ORUDIS SR (Other drug)			**Reason:** Osteoarthritis	
Capsule	150.0 Milligram		Daily	Oral
Batch:	**Started:** 01/05/1991		**Stopped:**	
AURORIX (Suspected)			**Reason:** Depression	
Tablet	150.0 Milligram		Daily	Oral
Batch:	**Started:** 29/10/1992		**Stopped:** 09/11/1992	

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81113 **Seq:** 1 **Gender:** F
Reported: 18/11/1992 **Weight:**
Hospitalisation: **Age:** 52Y
Onset Date: 31/08/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gait disturbance Tinnitus		Heaviness in legs.	

Medicine Details:

ALDAZINE (Other drug)		**Reason:**		
Tablet	75.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1988		**Stopped:**	CONTIN
ALODORM (Other drug)		**Reason:**		
Tablet	10.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1980		**Stopped:**	CONTIN
ANTENEX (Other drug)		**Reason:**		
Tablet	20.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1980		**Stopped:**	CONTIN
AURORIX (Suspected)		**Reason:** Depression		
Tablet	300.0 Milligram	Daily	Oral	
Batch:	**Started:** 10/08/1992		**Stopped:**	17/08/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Moclobemide ceased by patient and symptoms resolved within a few days. - 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.

Report Detail

Case Number: 81170 **Seq:** 1
Reported: 26/11/1992
Hospitalisation: Admitted to hospital
Onset Date: 06/10/1992
Outcome: Recovered

Gender: F
Weight: 50.00
Age: 33Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination Agitation		Visual hallucinations	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

0.0

Batch: **Started:** 05/10/1992 **Stopped:** 06/10/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81379 **Seq:** 1 **Gender:** F
Reported: 08/12/1992 **Weight:**
Hospitalisation: **Age:** 26Y
Onset Date: 02/12/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Coordination abnormal Hypotonia			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** 30/11/1992 **Stopped:** 03/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms for 5 days after ceasing moclobemide, patient is chinese and slight build.

Report Detail

Case Number: 81381 **Seq:** 1 **Gender:** F
Reported: 09/12/1992 **Weight:** 54.00
Hospitalisation: **Age:** 48Y
Onset Date: 21/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea			Dose reduced to 150mg bd

Medicine Details:

LEXOTAN (Other drug) **Reason:** Depression
Tablet 3.0 Milligram As necessary Oral
Batch: **Started:** 01/01/1991 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 21/09/1992 **Stopped:** 05/12/1992

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81400 **Seq:** 1 **Gender:** F
Reported: 06/12/1992 **Weight:** 65.00
Hospitalisation: **Age:** 30Y
Onset Date: 14/10/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 08/10/1992 **Stopped:** 26/10/1992

NORDETTE NOS (Suspected) **Reason:** Contraception
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 25/09/1992 **Stopped:** 14/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		14/11/1991	-	
	ALT = SGPT		13/05/1992	-	
	ALT = SGPT		14/10/1992	59	
	ALT = SGPT		19/10/1992	86	
	ALT = SGPT		25/11/1992	130	
	ALT = SGPT		02/11/1992	135	
	ALT = SGPT		28/10/1992	84	

Additional Information:

History of cholestatic jaundice of pregnancy (1989, 1992) requiring cholestyramine 4 gm t.d.s. - resolved after delivery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81400 **Seq:** 1
Reported: 06/12/1992
Hospitalisation:
Onset Date: 14/10/1992
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 30Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression
	150.0 Milligram	Daily		
Batch:	**Started:** 08/10/1992		**Stopped:**	26/10/1992
NORDETTE NOS (Suspected)			**Reason:**	Contraception
Tablet	1.0 Dose Unspecified	Daily		Oral
Batch:	**Started:** 25/09/1992		**Stopped:**	14/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		26/10/1992	104	
	AST = SGOT		14/11/1991	40	
	AST = SGOT		19/10/1992	38	
	AST = SGOT		26/10/1992	48	
	AST = SGOT		28/10/1992	38	
	AST = SGOT		25/11/1992	130	
	AST = SGOT		02/11/1992	60	

Additional Information:

History of cholestatic jaundice of pregnancy (1989, 1992) requiring cholestyramine 4 gm t.d.s. - resolved after delivery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81400 **Seq:** 1 **Gender:** F
Reported: 06/12/1992 **Weight:** 65.00
Hospitalisation: **Age:** 30Y
Onset Date: 14/10/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detai

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression
	150.0 Milligram	Daily		
Batch:	**Started:** 08/10/1992		**Stopped:**	26/10/1992
NORDETTE NOS (Suspected)			**Reason:**	Contraception
Tablet	1.0 Dose Unspecified	Daily		Oral
Batch:	**Started:** 25/09/1992		**Stopped:**	14/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	AST = SGOT		14/10/1992	27	
	AST = SGOT		13/05/1992	111	
	Biochemistry	BIL	14/11/1991	9	
	Biochemistry	BIL	19/10/1992	8	
	Biochemistry	BIL	26/10/1992	9	
	Biochemistry	BIL	28/10/1992	15	
	Biochemistry	BIL	25/11/1992	14	

Additional Information:

History of cholestatic jaundice of pregnancy (1989, 1992) requiring cholestyramine 4 gm t.d.s. - resolved after delivery.

Report Detail

Case Number: 81400 **Seq:** 1
Reported: 06/12/1992
Hospitalisation:
Onset Date: 14/10/1992
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 30Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
	150.0	Milligram	Daily		
Batch:	**Started:**	08/10/1992		**Stopped:**	26/10/1992
NORDETTE NOS (Suspected)			**Reason:**	Contraception	
Tablet	1.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**	25/09/1992		**Stopped:**	14/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry	BIL	02/11/1992	17	
	Biochemistry	BIL	14/10/1992	7	
	Biochemistry	BIL	13/05/1992	27	
	Erythrocyte		14/10/1992	78	
	Erythrocyte		26/10/1992	53	
	Liver biopsy				Liver biopsy 5/11/92 - granulomatous hepatitis consistent with
	SAP = ALP		14/11/1991	130	

Additional Information:

History of cholestatic jaundice of pregnancy (1989, 1992) requiring cholestyramine 4 gm t.d.s. - resolved after delivery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81400 **Seq:** 1
Reported: 06/12/1992
Hospitalisation:
Onset Date: 14/10/1992
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 30Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
	150.0	Milligram	Daily		
Batch:	**Started:**	08/10/1992		**Stopped:**	26/10/1992
NORDETTE NOS (Suspected)			**Reason:**	Contraception	
Tablet	1.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**	25/09/1992		**Stopped:**	14/10/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP		13/05/1992	356	
	SAP = ALP		14/10/1992	313	
	SAP = ALP		19/10/1992	410	
	SAP = ALP		26/10/1992	455	
	SAP = ALP		28/10/1992	387	
	SAP = ALP		02/11/1992	435	
	SAP = ALP		25/11/1992	334	

Additional Information:

History of cholestatic jaundice of pregnancy (1989, 1992) requiring cholestyramine 4 gm t.d.s. - resolved after delivery.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81536 **Seq:** 1 **Gender:** F
Reported: 15/12/1992 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 05/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension	Severe		

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81595 **Seq:** 1 **Gender:** F
Reported: 16/12/1992 **Weight:**
Hospitalisation: **Age:** 18Y
Onset Date: 15/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache			Changed to tofranil

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 06/11/1992 **Stopped:** 03/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 81598 **Seq:** 1 **Gender:** M
Reported: 14/12/1992 **Weight:** 90.00
Hospitalisation: **Age:** 50Y
Onset Date: 23/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Vertigo			

Medicine Details:

PRINIVIL (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 21/09/1992 **Stopped:**

ADALAT 20 (Other drug) **Reason:** Essential benign hypertension
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

TEGRETOL (Other drug) **Reason:**
400.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

LOSEC (Other drug) **Reason:** Other diseases of esophagus
40.0 Milligram Alternate days
Batch: **Started:** **Stopped:** 28/09/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Significant cheese (mature type) intake 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81598 **Seq:** 1 **Gender:** M
Reported: 14/12/1992 **Weight:** 90.00
Hospitalisation: **Age:** 50Y
Onset Date: 23/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 21/11/1992 **Stopped:** 05/12/1992

Laboratory Investigations:

Additional Information:

Significant cheese (mature type) intake the dosage stop date is not necessarily accurate but indicates that stoppage occurred sometime during the month.

Report Detail

Case Number: 81618 **Seq:** 1 **Gender:** F
Reported: 21/12/1992 **Weight:** 49.00
Hospitalisation: **Age:** 52Y
Onset Date: 21/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Supraventricular tachycardia Ventricular tachycardia			

Medicine Details:

DIAZEPAM (Other drug) **Reason:** Depression
Oral application 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

TEMAZEPAM (Other drug) **Reason:** Depression
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 04/11/1992 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81749 **Seq:** 1 **Gender:** F
Reported: 23/12/1992 **Weight:** 73.00
Hospitalisation: **Age:** 65Y
Onset Date: 13/12/1992 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura Vasculitis			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 08/12/1992 **Stopped:** 12/12/1992

MELLERIL (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 09/12/1992 **Stopped:** 12/12/1992

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

DESIPRAMINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Tablet 750.0 Milligram Daily Oral
Batch: **Started:** 24/11/1992 **Stopped:** 27/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haematology				Skin biospy: leukocytoclastic vasculitis in keeping with drug

Additional Information:

Report Detail

Case Number: 81859 **Seq:** 1 **Gender:** F
Reported: 08/01/1993 **Weight:** 78.00
Hospitalisation: **Age:** 43Y
Onset Date: 29/12/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state		Foggy in the head	
Visual disturbance		Staring attacks	
Asthenia			
Dizziness			

Medicine Details:

PRINIVIL (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 01/08/1992 **Stopped:** CONTIN

MURELAX (Suspected) **Reason:** Anxiety neurosis
15.0 Milligram As necessary
Batch: **Started:** **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 29/12/1992 **Stopped:** 04/01/1993

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: 81888 **Seq:** 1 **Gender:** M
Reported: 07/01/1993 **Weight:** 64.00
Hospitalisation: **Age:** 56Y
Onset Date: 25/11/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Severe		

Medicine Details:

CLONAZEPAM (Other drug) **Reason:** Anxiety neurosis

Tablet 2.0 Milligram Daily Oral

Batch: **Started:** 09/07/1992 **Stopped:** 17/12/1992

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 22/10/1992 **Stopped:** 01/12/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 81896 **Seq:** 1 **Gender:** M
Reported: 08/01/1993 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: 10/12/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erectile dysfunction			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 10/11/1992 **Stopped:** 08/01/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 81898 **Seq:** 1 **Gender:** M
Reported: 05/01/1993 **Weight:** 70.00
Hospitalisation: **Age:** 25Y
Onset Date: 12/12/1992 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety			

Medicine Details:

HALOPERIDOL (Other drug) **Reason:** Unspecified psychosis
Oral application 5.0 Milligram Daily Oral
Batch: **Started:** 21/10/1992 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 11/12/1992 **Stopped:** 17/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient rechallenged on moclobemide with same reaction.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82002 **Seq:** 1 **Gender:** F
Reported: 13/01/1993 **Weight:** 63.00
Hospitalisation: **Age:** 56Y
Onset Date: 21/12/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence	Severe		

Medicine Details:

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

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 14/12/1992 **Stopped:** 29/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 82198 **Seq:** 1
Reported: 26/01/1993
Hospitalisation:
Onset Date: 09/11/1992
Outcome: Recovered with sequelae

Gender: M
Weight: 56.00
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle twitching Chills Hyperhidrosis		Spasms of facial muscles and arms.	

Medicine Details:

MADOPAR M (Other drug) **Reason:** Paralysis agitans
Oral application 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/07/1991 **Stopped:**

MADOPAR Q (Other drug) **Reason:** Paralysis agitans
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/04/1990 **Stopped:**

SINEMET CR (Other drug) **Reason:** Paralysis agitans
Tablet 250.0 Milligram Daily Oral
Batch: **Started:** 01/09/1992 **Stopped:**

NORMISON (Other drug) **Reason:** Specific disorders of sleep
10.0 Milligram Daily
Batch: **Started:** L TERM **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. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 82198 **Seq:** 1
Reported: 26/01/1993
Hospitalisation:
Onset Date: 09/11/1992
Outcome: Recovered with sequelae

Gender: M
Weight: 56.00
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** 05/11/1992 **Stopped:**

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.

Report Detail

Case Number: 82201 **Seq:** 1 **Gender:** M
Reported: 22/01/1993 **Weight:** 73.00
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension			

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected) **Reason:** Pain

Injection 75.0 Milligram As necessary Intramuscular

Batch: **Started:** 20/01/1993 **Stopped:** 22/01/1993

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82377 **Seq:** 1 **Gender:** M
Reported: 04/02/1993 **Weight:** 86.00
Hospitalisation: **Age:** 29Y
Onset Date: 23/01/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Constipation Hyperhidrosis Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 21/01/1993 **Stopped:** 03/02/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 82595 **Seq:** 1 **Gender:** M
Reported: 15/02/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 09/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Conjunctivitis	Severe		
Headache	Severe		
Tongue oedema	Severe		

Medicine Details:

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

RIVOTRIL (Other drug) **Reason:**
1.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

EPILIM (Other drug) **Reason:**
1.2 Gram Daily
Batch: **Started:** L TERM **Stopped:**

DILANTIN (Other drug) **Reason:**
300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82595 **Seq:** 1 **Gender:** M
Reported: 15/02/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 09/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ALCOHOL (Suspected)		**Reason:**	
	0.0	1 time	
Batch:	**Started:**		**Stopped:**
AURORIX (Suspected)		**Reason:**	Depression
	300.0 Milligram	Daily	
Batch:	**Started:** 05/02/1993		**Stopped:** 09/02/1993

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 82744 **Seq:** 1 **Gender:** F
Reported: 07/01/1993 **Weight:** 45.00
Hospitalisation: **Age:** 68Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Therapeutic response decreased			

Medicine Details:

EUHYPNOS (Other drug) **Reason:** Anxiety neurosis
10.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

NAPROSYN (Other drug) **Reason:** Osteoarthritis
Tablet 1.0 Gram Daily Oral
Batch: **Started:** L TERM **Stopped:**

SERENACE (Other drug) **Reason:** Otr spec symp psychopathol nec
Tablet 1.5 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

SEREPAX (Other drug) **Reason:** Anxiety neurosis
Tablet 30.0 Milligram Daily Oral
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82744 **Seq:** 1 **Gender:** F
Reported: 07/01/1993 **Weight:** 45.00
Hospitalisation: **Age:** 68Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 21/12/1992 **Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 82785 **Seq:** 1 **Gender:** F
Reported: 23/02/1993 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 18/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Nausea		Sudden pulses in the brain.	

Medicine Details:

PANADEINE FORTE (Suspected) **Reason:** Pain
2.0 Dose Unspecified As necessary
Batch: **Started:** 18/02/1993 **Stopped:** 18/02/1993

PETHIDINE HYDROCHLORIDE (Suspected) **Reason:** Pain
Injection 100.0 Milligram As necessary Intramuscular
Batch: **Started:** 18/02/1993 **Stopped:** 19/02/1993

BUSCOPAN (Suspected) **Reason:** Abdominal pain
Tablet 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** 18/02/1993 **Stopped:** 19/02/1993

FLAGYL (Suspected) **Reason:** Pain
1.2 Gram Daily
Batch: **Started:** 18/02/1993 **Stopped:** 23/02/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergics sulphur, maxolon and panadeine.

Report Detail

Case Number: 82785 **Seq:** 1 **Gender:** F
Reported: 23/02/1993 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 18/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected)		**Reason:** Depression		
	450.0 Milligram	Daily		
Batch:	**Started:** 18/02/1993		**Stopped:** 23/02/1993	
GENTAMICIN SULPHATE (Suspected)		**Reason:** Other cellulitis and abscess		
Injection	160.0 Milligram	Daily	Intramuscular	
Batch:	**Started:** 18/02/1993		**Stopped:** 25/02/1993	
AMPICILLIN (Suspected)		**Reason:** Other cellulitis and abscess		
Injection	4.0 Gram	Daily	Intravenous	
Batch:	**Started:** 18/02/1993		**Stopped:** 22/02/1993	
SEREPAX (Suspected)		**Reason:** Depression		
	60.0 Milligram	Daily		
Batch:	**Started:** 18/02/1993		**Stopped:** 23/03/1993	

Laboratory Investigations:

Additional Information:

Allergics sulphur, maxolon and panadeine.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82833 **Seq:** 1 **Gender:** F
Reported: 25/02/1993 **Weight:** 100.00
Hospitalisation: **Age:** 36Y
Onset Date: 25/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 25/02/1993 **Stopped:** 26/02/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 82840 **Seq:** 1 **Gender:** F
Reported: 24/02/1993 **Weight:** 67.00
Hospitalisation: **Age:** 35Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Insomnia Nausea Vertigo Vomiting			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 08/02/1993 **Stopped:** 22/02/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 82846 **Seq:** 1 **Gender:** F
Reported: 22/02/1993 **Weight:** 66.00
Hospitalisation: **Age:** 51Y
Onset Date: 18/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Paraesthesia			

Medicine Details:

INDERAL (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 18/02/1993 **Stopped:** 20/02/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 82911 **Seq:** 1 **Gender:** F
Reported: 10/02/1993 **Weight:**
Hospitalisation: **Age:** 54Y
Onset Date: 22/01/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Vomiting			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 22/01/1993 **Stopped:** 24/01/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83109 **Seq:** 1 **Gender:** F
Reported: 11/03/1993 **Weight:** 61.00
Hospitalisation: **Age:** 40Y
Onset Date: 25/02/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia Nausea			Moclobemide dose reduced to 300 mg daily.

Medicine Details:

VENTOLIN (Other drug) **Reason:** Asthma
Inhalation 0.0 Inhalation
Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 25/02/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83199 **Seq:** 1 **Gender:** F
Reported: 15/03/1993 **Weight:** 55.00
Hospitalisation: **Age:** 35Y
Onset Date: 10/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Increased appetite			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 05/03/1993 **Stopped:** 13/03/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83221 **Seq:** 1 **Gender:** F
Reported: 15/03/1993 **Weight:** 66.00
Hospitalisation: **Age:** 64Y
Onset Date: 28/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 28/02/1993 **Stopped:** 05/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had nightmares every time after taking moclobemide 150mg.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83430 **Seq:** 1 **Gender:** M
Reported: 23/03/1993 **Weight:** 76.00
Hospitalisation: **Age:** 51Y
Onset Date: 10/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pollakiuria Urinary retention			

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 05/03/1993 **Stopped:** 19/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83458 **Seq:** 1 **Gender:** F
Reported: 25/03/1993 **Weight:**
Hospitalisation: **Age:** 73Y
Onset Date: 15/03/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dry mouth			

Medicine Details:

VENTOLIN (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 1.0 Dose Unspecified As necessary Inhalation
Batch: **Started:** **Stopped:**

ANGININE (Other drug) **Reason:** Angina pectoris w/o hyperten
1.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

PANADOL (Other drug) **Reason:**
2.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

DISPRIN (Other drug) **Reason:** Chron isch heart dis no hyper
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83458 **Seq:** 1
Reported: 25/03/1993
Hospitalisation:
Onset Date: 15/03/1993
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

LANOXIN (Other drug)			**Reason:**	Chron isch heart dis no hyper	
Tablet	250.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
CARDIZEM (Other drug)			**Reason:**	Chron isch heart dis no hyper	
Tablet	240.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
TRANSIDERM-NITRO (Other drug)			**Reason:**	Chron isch heart dis no hyper	
Cream	50.0	Milligram	Daily		Topical
Batch:	**Started:**			**Stopped:**	
ATROVENT (Other drug)			**Reason:**	Otr respiratory systm diseases	
Inhalation	2.0	Millilitre	Daily		Inhalation
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 83458 **Seq:** 1
Reported: 25/03/1993
Hospitalisation:
Onset Date: 15/03/1993
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

PULMICORT (Other drug)			**Reason:**	Otr respiratory systm diseases	
Inhalation	3.2	Milligram	Daily		Inhalation
Batch:	**Started:**			**Stopped:**	
MOGADON (Other drug)			**Reason:**		
Tablet	10.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**	09/03/1993		**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 83597 **Seq:** 1 **Gender:** M
Reported: 25/03/1993 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 21/03/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash morbilliform			Prednisolone dose increased

Medicine Details:

DIPRIVAN (Other drug) **Reason:**
3.0 Milligram Daily
Batch: **Started:** 01/01/1993 **Stopped:** CONTIN

ASPIRIN (Other drug) **Reason:** Other rheumatoid arthritis
Oral application 150.0 Milligram Daily Oral
Batch: **Started:** 01/01/1993 **Stopped:** CONTIN

PLAQUENIL (Suspected) **Reason:** Other rheumatoid arthritis
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** 07/03/1993 **Stopped:** 22/03/1993

PREDNISOLONE (Suspected) **Reason:** Other rheumatoid arthritis
Oral application 5.0 Milligram Daily Oral
Batch: **Started:** 07/03/1993 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83597 **Seq:** 1
Reported: 25/03/1993
Hospitalisation:
Onset Date: 21/03/1993
Outcome: Not yet recovered

Gender: M
Weight:
Age: 60Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected)			**Reason:**		
Tablet	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	10/03/1993	**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.

Report Detail

Case Number: 83653 **Seq:** 1 **Gender:** M
Reported: 27/03/1993 **Weight:** 65.00
Hospitalisation: **Age:** 45Y
Onset Date: 24/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia Depersonalisation Depression Insomnia Nervousness		Over stimulated	

Medicine Details:

LIPEX (Other drug) **Reason:** Othr&unspec metabolic diseases
20.0 Milligram Daily
Batch: **Started:** 29/01/1993 **Stopped:**

DEMAZIN REPETABS (Suspected) **Reason:**
Tablet 1.0 Dose Unspecified 1 time Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 04/03/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83722 **Seq:** 1 **Gender:** F
Reported: 02/04/1993 **Weight:** 55.00
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache	Severe		

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 17/11/1992 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83791 **Seq:** 1 **Gender:** M
Reported: 23/06/1992 **Weight:**
Hospitalisation: **Age:**
Onset Date: 16/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Dizziness			

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 16/06/1992 **Stopped:** 23/06/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83792 **Seq:** 1 **Gender:** F
Reported: 03/08/1992 **Weight:** 73.00
Hospitalisation: **Age:** 72Y
Onset Date: 25/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Insomnia Vomiting			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 24/06/1992 **Stopped:** 12/07/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 83925 **Seq:** 1 **Gender:** F
Reported: 15/04/1993 **Weight:** 48.00
Hospitalisation: **Age:** 53Y
Onset Date: 07/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Syncope			Fluids, rest.

Medicine Details:

SUDAFED (Suspected) **Reason:**
2.0 Dose Unspecified Total
Batch: **Started:** 07/04/1993 **Stopped:**

MELLERIL (Suspected) **Reason:**
50.0 Milligram As necessary
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has mitral valve disease (?rheumatic origin) difficult antidepressant choice.

Report Detail

Case Number: 83970 **Seq:** 1 **Gender:** M
Reported: 07/04/1993 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: 02/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			

Medicine Details:

NAPROSYN SR (Other drug) **Reason:**
750.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 02/04/1993 **Stopped:** 02/04/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 83976 **Seq:** 1 **Gender:** M
Reported: 16/04/1993 **Weight:** 82.00
Hospitalisation: **Age:** 72Y
Onset Date: 02/04/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction			

Medicine Details:

ISOPTIN (Other drug) **Reason:** Essential benign hypertension
240.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

DYAZIDE (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1990 **Stopped:**

CAPOTEN (Other drug) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 19/03/1993 **Stopped:** 16/04/1993

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84101 **Seq:** 1 **Gender:** F
Reported: 23/04/1993 **Weight:** 67.00
Hospitalisation: **Age:** 47Y
Onset Date: 17/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myalgia Drug ineffective		Aching,insommnia, headache & tight head	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 15/04/1993 **Stopped:** 21/04/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84106 **Seq:** 1 **Gender:** F
Reported: 23/04/1993 **Weight:** 69.00
Hospitalisation: **Age:** 45Y
Onset Date: 19/04/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hot flush Diarrhoea		Hot all over body.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 17/04/1993 **Stopped:** 21/04/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 84127 **Seq:** 1 **Gender:** F
Reported: 02/04/1993 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 20/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Constipation Hallucination			

Medicine Details:

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

VALIUM (Other drug) **Reason:**
7.5 Milligram Daily
Batch: **Started:** **Stopped:**

NORMISON (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 13/03/1993 **Stopped:** 29/03/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84194 **Seq:** 1 **Gender:** M
Reported: 26/04/1993 **Weight:** 79.00
Hospitalisation: **Age:** 66Y
Onset Date: 25/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension	Severe		

Medicine Details:

ENALAPRIL MALEATE (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** 01/03/1993 **Stopped:**

ATENOLOL (Other drug) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

WARFARIN SODIUM (Other drug) **Reason:** Phleb&thrombophleb lowr extrem
2.5 Milligram Daily
Batch: **Started:** 01/01/1993 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
150.0 Milligram 3 times
Batch: **Started:** 25/04/1993 **Stopped:** 26/04/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient previously hypertensive-bp 180/120, with three doses of moclobemide 150mg-bp 90/70. when moclobemide ceased, rapid increase to 150/90 in 8 hours. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84194 **Seq:** 1 **Gender:** M
Reported: 26/04/1993 **Weight:** 79.00
Hospitalisation: **Age:** 66Y
Onset Date: 25/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

MELLERIL (Suspected) **Reason:** Otr spec symp psychopathol nec
75.0 Milligram Daily
Batch: **Started:** 24/04/1993 **Stopped:** 26/04/1993

Laboratory Investigations:

Additional Information:

Patient previously hypertensive-bp 180/120, with three doses of moclobemide 150mg-bp 90/70. when moclobemide ceased, rapid increase to 150/90 in 8 hours. 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: 84239 **Seq:** 1 **Gender:** M
Reported: 30/04/1993 **Weight:** 70.00
Hospitalisation: **Age:** 47Y
Onset Date: 01/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia Pruritus			

Medicine Details:

FELDENE (Other drug) **Reason:** Unspecified arthritis
20.0 Milligram As necessary
Batch: **Started:** **Stopped:**

ZANTAC (Other drug) **Reason:** Ulcer of duodenum w/o ment perf
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 27/01/1993 **Stopped:** 03/02/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84274 **Seq:** 1 **Gender:** M
Reported: 03/05/1993 **Weight:**
Hospitalisation: **Age:** 73Y
Onset Date: 25/04/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Prothrombin level decreased			Vitamin k

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 17/04/1993 **Stopped:**

WARFARIN SODIUM (Suspected) **Reason:** Otr&nos disord of heart rhythm
2.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

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

SLOW-K (Suspected) **Reason:** Congestive heart failure
4.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	International		22/04/1993	2.3	
	International		25/04/1993	4.7	
	International		26/04/1993	6.2	
	International		27/04/1993	8.0	

Additional Information:

Report Detail

Case Number: 84274 **Seq:** 1
Reported: 03/05/1993
Hospitalisation:
Onset Date: 25/04/1993
Outcome: Not yet recovered

Gender: M
Weight:
Age: 73Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

LANOXIN (Suspected)			**Reason:**	Otr&nos disord of heart rhythm
	125.0	Microgram	Daily	
Batch:	**Started:**			**Stopped:**
CAPOTEN (Suspected)			**Reason:**	Congestive heart failure
	75.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:** 25/04/1993
SERENACE (Suspected)			**Reason:**	Depression
	1.5	Milligram	Daily	
Batch:	**Started:** 17/04/1993			**Stopped:** 26/04/1993

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84597 **Seq:** 1 **Gender:** F
Reported: 18/05/1993 **Weight:** 60.00
Hospitalisation: **Age:** 33Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety Face oedema			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/04/1993 **Stopped:** 28/04/1993 5D

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84777 **Seq:** 1 **Gender:** F
Reported: 31/05/1993 **Weight:** 93.00
Hospitalisation: **Age:** 60Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension			

Medicine Details:

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

NATRILIX (Other drug) **Reason:**
2.5 Milligram Daily
Batch: **Started:** **Stopped:**

VOLTAREN (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

ISOPTIN SR (Other drug) **Reason:**
240.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Noticeable increase in blood pressure while on aurorix which came down after ceasing drug.

Report Detail

Case Number: 84777 **Seq:** 1 **Gender:** F
Reported: 31/05/1993 **Weight:** 93.00
Hospitalisation: **Age:** 60Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

NOLVADEX (Other drug)			**Reason:**	
	20.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
TEGRETOL (Other drug)			**Reason:**	
	400.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
CALTRATE (Other drug)			**Reason:**	
	600.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
AURORIX (Suspected)			**Reason:**	Depression
	450.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Noticeable increase in blood pressure while on aurorix which came down after ceasing drug.

Report Detail

Case Number: 84780 **Seq:** 1 **Gender:** F
Reported: 31/05/1993 **Weight:** 74.00
Hospitalisation: **Age:** 59Y
Onset Date: 18/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation Dizziness			

Medicine Details:

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

ESTRADERM (Other drug) **Reason:**
Cream 0.0 Transdermal
Batch: **Started:** **Stopped:**

TRITACE (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

PROVERA (Other drug) **Reason:**
1.0 Dose Unspecified 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84780 **Seq:** 1 **Gender:** F
Reported: 31/05/1993 **Weight:** 74.00
Hospitalisation: **Age:** 59Y
Onset Date: 18/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 17/05/1993 **Stopped:** 18/05/1993

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 84794 **Seq:** 1 **Gender:** M
Reported: 31/05/1993 **Weight:**
Hospitalisation: **Age:** 26Y
Onset Date: 17/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Rectal haemorrhage	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 01/10/1992 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85056 **Seq:** 1 **Gender:** F
Reported: 01/06/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 24/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Agitation Malaise		Impaired sleep.	

Medicine Details:

DIAZEPAM (Other drug) **Reason:** Anxiety neurosis

Tablet 5.0 Milligram Daily Oral

Batch: **Started:** 20/04/1993 **Stopped:** 25/04/1993

MOCLOBEMIDE (Suspected) **Reason:** Otr spec symp psychopathol nec

Tablet 150.0 Milligram 1 time Oral

Batch: **Started:** 20/04/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 85108 **Seq:** 1 **Gender:** F
Reported: 03/06/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 27/05/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		Disturbed vision	

Medicine Details:

AURORIX (Suspected) **Reason:** Specific disorders of sleep
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 27/05/1993 **Stopped:** 29/05/1993

DUCENE (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

LIPEX (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

QUINATE (Suspected) **Reason:**
1.0 Dose Unspecified 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85108 **Seq:** 1
Reported: 03/06/1993
Hospitalisation:
Onset Date: 27/05/1993
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

VOLTAREN (Suspected)			**Reason:**	
	50.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
AMPRACE (Suspected)			**Reason:**	
	20.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 85154 **Seq:** 1 **Gender:** M
Reported: 08/06/1993 **Weight:** 66.00
Hospitalisation: **Age:** 68Y
Onset Date: 22/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle contractions involuntary			

Medicine Details:

LANOXIN (Other drug) **Reason:** Otr&nos disord of heart rhythm
375.0 Microgram Daily
Batch: **Started:** 01/01/1985 **Stopped:**

BETALOC (Other drug) **Reason:** Otr&nos disord of heart rhythm
50.0 Microgram Daily
Batch: **Started:** 01/01/1985 **Stopped:**

XANAX (Other drug) **Reason:**
250.0 Microgram Daily
Batch: **Started:** 01/01/1986 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 09/09/1992 **Stopped:** 20/03/1993

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85154 **Seq:** 1 **Gender:** M
Reported: 08/06/1993 **Weight:** 66.00
Hospitalisation: **Age:** 68Y
Onset Date: 22/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TOLVON (Suspected) **Reason:** Depression
30.0 Milligram Daily
Batch: **Started:** 21/03/1993 **Stopped:** 23/03/1993

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.

Report Detail

Case Number: 85267 **Seq:** 1 **Gender:** F
Reported: 10/06/1993 **Weight:** 74.00
Hospitalisation: **Age:** AD
Onset Date: 02/06/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura Pruritus		Purpuric like rash on arms & upper abd.	

Medicine Details:

ATIVAN (Other drug) **Reason:** Anxiety neurosis
1.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 31/05/1993 **Stopped:** 04/06/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85312 **Seq:** 1 **Gender:** M
Reported: 15/06/1993 **Weight:**
Hospitalisation: **Age:** 11
Onset Date: **DOB:** 06/01/1982
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myopathy			

Medicine Details:

RIVOTRIL (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 01/02/1993 **Stopped:**

SINEQUAN (Other drug) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 01/02/1993 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
900.0 Milligram Daily
Batch: **Started:** 01/02/1993 **Stopped:** 28/05/1993

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85339 **Seq:** 1 **Gender:** F
Reported: 15/06/1993 **Weight:** 58.00
Hospitalisation: **Age:** 25Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Headache			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 225.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85348 **Seq:** 1 **Gender:** F
Reported: 15/06/1993 **Weight:** 41.00
Hospitalisation: **Age:** 56Y
Onset Date: 13/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Syncope			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 12/06/1993 **Stopped:** 14/06/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Within 10-15 minutes of taking second dose of aurorix. patient felt dizzy she lay down and felt ok but fainted after standing. symptoms recurred after third dose.

Report Detail

Case Number: 85355 **Seq:** 1 **Gender:** F
Reported: 17/06/1993 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension Insomnia			

Medicine Details:

PROGYNOVA (Other drug) **Reason:** Menopausal symptoms
Tablet 2.0 Milligram Daily Oral
Batch: **Started:** 03/08/1992 **Stopped:** CONTIN

SANDOMIGRAN (Other drug) **Reason:** Migraine
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/01/1989 **Stopped:** CONTIN

TEMAZEPAM (Other drug) **Reason:**
Capsule 10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 08/04/1993 **Stopped:** 01/06/1993

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: 85481 **Seq:** 1 **Gender:** F
Reported: 24/06/1993 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia		Memory lapses.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 29/03/1993 **Stopped:** 02/06/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85837 **Seq:** 1 **Gender:** F
Reported: 02/07/1993 **Weight:**
Hospitalisation: **Age:** 39Y
Onset Date: 03/02/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myalgia	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction resolved spontaneously after increasing dose to 900mg/d.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85863 **Seq:** 1 **Gender:** F
Reported: 06/07/1993 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: 27/05/1992 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Psychotic disorder Thinking abnormal			

Medicine Details:

NORMISON (Other drug) **Reason:**
3.0 Gram Daily
Batch: **Started:** **Stopped:**

NAPROSYN SR (Other drug) **Reason:**
1.0 Gram Daily
Batch: **Started:** **Stopped:**

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

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 22/05/1992 **Stopped:** 27/05/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85945 **Seq:** 1 **Gender:** F
Reported: 09/07/1993 **Weight:** 63.00
Hospitalisation: **Age:** 68Y
Onset Date: 21/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido increased		Nymphomania.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
1.0 Dose Unspecified Daily
Batch: **Started:** 19/06/1993 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85993 **Seq:** 1 **Gender:** F
Reported: 09/07/1993 **Weight:**
Hospitalisation: **Age:** 53Y
Onset Date: 31/08/1992 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Amnesia			

Medicine Details:

AURORIX (Suspected) **Reason:**
450.0 Milligram Daily
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 85994 **Seq:** 1 **Gender:** M
Reported: 09/07/1993 **Weight:**
Hospitalisation: **Age:** 54Y
Onset Date: 17/09/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suicide attempt Agitation Anxiety		Suicidal ideation.	

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 20/08/1992 **Stopped:** 08/09/1992

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86116 **Seq:** 1 **Gender:** M
Reported: 19/07/1993 **Weight:** 100.00
Hospitalisation: **Age:** 44Y
Onset Date: 07/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression			

Medicine Details:

AMFAMOX (Other drug) **Reason:** Other diseases of esophagus
20.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient stopped moclobemide then 1 1/2 days later recommenced with same reaction occurring. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86256 **Seq:** 1 **Gender:** F
Reported: 23/07/1993 **Weight:** 64.00
Hospitalisation: **Age:** 56Y
Onset Date: 16/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence Thrombocytopenia			

Medicine Details:

SUMATRIPTAN (Suspected) **Reason:** Migraine
0.0
Batch: **Started:** **Stopped:** 18/06/1993

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 20/06/1993

INDERAL (Suspected) **Reason:** Headache
Tablet 40.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86286 **Seq:** 1 **Gender:** F
Reported: 27/07/1993 **Weight:** 82.00
Hospitalisation: **Age:** 65Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation		Felling of unreality.	

Medicine Details:

PRINIVIL (Other drug) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** 01/01/1993 **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
200.0 Microgram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

APRINOX (Other drug) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

SEREPAX (Other drug) **Reason:** Anxiety neurosis
15.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on auroxix the next day with same reaction occurring. 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 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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86286 **Seq:** 1 **Gender:** F
Reported: 27/07/1993 **Weight:** 82.00
Hospitalisation: **Age:** 65Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 13/07/1993 **Stopped:** 14/07/1993

Laboratory Investigations:

Additional Information:

Patient was rechallenged on auroxix the next day with same reaction occurring. 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 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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86336 **Seq:** 1 **Gender:** F
Reported: 30/07/1993 **Weight:**
Hospitalisation: **Age:**
Onset Date: 13/07/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression			
Agitation			
Asthenia			
Depersonalisation			
Depression			
Insomnia			
Nausea			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 02/06/1993 **Stopped:** 20/07/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 86584 **Seq:** 1 **Gender:** F
Reported: 04/08/1993 **Weight:** 42.00
Hospitalisation: **Age:** 77Y
Onset Date: 25/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety Hyperkinesia	Severe		

Medicine Details:

NIFEDIPINE (Other drug) **Reason:** Essential benign hypertension
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

SEREPAX (Other drug) **Reason:** Specific disorders of sleep
30.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

TIMOPTOL (Other drug) **Reason:** Unspecified glaucoma
Eye Drops 2.0 Dose Unspecified Daily Ophthalmic
Batch: **Started:** 01/09/1992 **Stopped:**

PILOCARPINE (Other drug) **Reason:** Unspecified glaucoma
Eye Drops 2.0 Dose Unspecified Daily Ophthalmic
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. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 86584 **Seq:** 1 **Gender:** F
Reported: 04/08/1993 **Weight:** 42.00
Hospitalisation: **Age:** 77Y
Onset Date: 25/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected)		**Reason:**	Depression
	300.0 Milligram	Daily	
Batch:	**Started:** 04/05/1993	**Stopped:**	18/06/1993
AURORIX (Suspected)		**Reason:**	Depression
	450.0 Milligram	Daily	
Batch:	**Started:** 18/06/1993	**Stopped:**	15/07/1993

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86595 **Seq:** 1 **Gender:** F
Reported: 04/08/1993 **Weight:** 48.00
Hospitalisation: **Age:** 79Y
Onset Date: 31/03/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Myalgia Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 08/03/1993 **Stopped:** 01/04/1993

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: 86617 **Seq:** 1 **Gender:** M
Reported: 04/08/1993 **Weight:** 80.00
Hospitalisation: **Age:** 46Y
Onset Date: 25/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tardive dyskinesia		Dyskinesia tardive mouth and tongue.	

Medicine Details:

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

FLUPHENAZINE DECANOATE (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** 20/06/1993

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/05/1993 **Stopped:** 20/06/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 86656 **Seq:** 1 **Gender:** M
Reported: 09/08/1993 **Weight:** 63.00
Hospitalisation: **Age:** 76Y
Onset Date: 24/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression			
Dizziness			
Dysgeusia			
Dyspnoea			
Headache			
Oliguria			

Medicine Details:

NARDIL (Other drug) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1987 **Stopped:** CONTIN

VENTOLIN (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 4.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/01/1986 **Stopped:**

INTAL (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 4.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/01/1986 **Stopped:**

BECOTIDE (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 2.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/01/1986 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Discontinue aurorix - taking nardil. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86656 **Seq:** 1 **Gender:** M
Reported: 09/08/1993 **Weight:** 63.00
Hospitalisation: **Age:** 76Y
Onset Date: 24/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 24/04/1993 **Stopped:** 26/04/1993

Laboratory Investigations:

Additional Information:

Discontinue aurorix - taking nardil. 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.

Report Detail

Case Number: 86818 **Seq:** 1 **Gender:** F
Reported: 16/08/1993 **Weight:** 51.00
Hospitalisation: **Age:** 44Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal			

Medicine Details:

PANADEINE FORTE (Other drug) **Reason:** Pain
Oral application 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

MERSYNDOL FORTE (Other drug) **Reason:** Pain
Tablet 2.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

PROZAC (Other drug) **Reason:** Depression
Capsule 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

LARGACTIL (Other drug) **Reason:** Depression
Oral application 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergy to cod po4 inj, indocid, di-gesic and thyroxine. see original report for rest of other drugs.

Report Detail

Case Number: 86818 **Seq:** 1
Reported: 16/08/1993
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 51.00
Age: 44Y
DOB:
Causality: Causality probable

Reaction Detail

Medicine Details:

AMOXIL (Other drug)			**Reason:**	Otr&nos infec&parasit diseases
Oral application	1.0	Gram	Daily	Oral
Batch:	**Started:**	07/08/1993	**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression
Tablet	300.0	Milligram	Daily	Oral
Batch:	**Started:**		**Stopped:**	10/08/1993

Laboratory Investigations:

Additional Information:

Allergy to cod po4 inj, indocid, di-gesic and thyroxine. see original report for rest of other drugs.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 86921 **Seq:** 1 **Gender:** F
Reported: 19/08/1993 **Weight:** 48.00
Hospitalisation: **Age:** 71Y
Onset Date: 28/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema Pain Paraesthesia			

Medicine Details:

VENTOLIN (Other drug) **Reason:** Asthma
Inhalation 8.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/01/1990 **Stopped:** CONTIN

PULMICORT (Other drug) **Reason:** Asthma
Inhalation 4.0 Dose Unspecified Daily Inhalation
Batch: **Started:** 01/01/1992 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/07/1993 **Stopped:** 03/08/1993

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: 87002 **Seq:** 1 **Gender:** F
Reported: 23/08/1993 **Weight:** 70.00
Hospitalisation: **Age:** 36Y
Onset Date: 31/01/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperhidrosis Hypertension Mydriasis Tachycardia Tremor			

Medicine Details:

NORDETTE 28 (Other drug)			**Reason:**	Contraception	
Pill	1.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**	01/01/1980		**Stopped:**	
BECLOFORTE (Other drug)			**Reason:**	Asthma	
Inhalation	4.0	Dose Unspecified	Daily		Inhalation
Batch:	**Started:**	01/01/1992		**Stopped:**	
VENTOLIN (Other drug)			**Reason:**	Asthma	
Inhalation	4.0	Dose Unspecified	Daily		Inhalation
Batch:	**Started:**	01/01/1992		**Stopped:**	
AURORIX (Suspected)			**Reason:**		
Tablet	525.0	Milligram	Daily		Oral
Batch:	**Started:**	12/01/1993		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient continues on aurorix at a reduced level of 500mg daily with no reaction occurring. - 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.

Report Detail

Case Number: 87034 **Seq:** 1 **Gender:** M
Reported: 24/08/1993 **Weight:** 69.00
Hospitalisation: **Age:** 32Y
Onset Date: 06/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hallucination			

Medicine Details:

ROHYPNOL (Other drug) **Reason:** Specific disorders of sleep
Tablet 4.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 05/08/1993 **Stopped:** 07/08/1993

PARNATE (Suspected) **Reason:** Depression
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 29/05/1990 **Stopped:** 04/08/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87238 **Seq:** 1 **Gender:** M
Reported: 23/08/1993 **Weight:** 85.00
Hospitalisation: **Age:** 48Y
Onset Date: 06/04/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Hepatitis Nausea Vomiting			Prepulsid, pepcidine.

Medicine Details:

COLOFAC (Other drug) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

XANAX (Other drug) **Reason:** Anxiety neurosis
0.5 Milligram As necessary
Batch: **Started:** 31/03/1993 **Stopped:** 06/04/1993

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 31/03/1993 **Stopped:** 30/04/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	5 - 40	02/05/1993	81	Minor elevation of lfts.

Additional Information:

Symptoms recurred when recommenced on aurorix.

Report Detail

Case Number: 87568 **Seq:** 1 **Gender:** F
Reported: 10/09/1993 **Weight:** 58.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 22/03/1993 **Stopped:** 21/06/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87597 **Seq:** 1 **Gender:** M
Reported: 10/09/1993 **Weight:** 105.00
Hospitalisation: **Age:** 26Y
Onset Date: 27/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Disturbance in attention			

Medicine Details:

STELAZINE (Suspected) **Reason:** Unspecified schizophrenia
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** 01/01/1992 **Stopped:** 20/08/1993

AURORIX (Suspected) **Reason:** Unspecified schizophrenia
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 27/07/1993 **Stopped:** 20/08/1993

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: 87635 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:** 36.00
Hospitalisation: **Age:** 60Y
Onset Date: 28/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Jerking movements of arms, legs & head.	

Medicine Details:

VENTOLIN (Other drug) **Reason:** Asthma
Inhalation 0.0 As necessary Inhalation
Batch: **Started:** L TERM **Stopped:**

BECOTIDE (Other drug) **Reason:** Asthma
Inhalation 2.0 Dose Unspecified Daily Inhalation
Batch: **Started:** L TERM **Stopped:**

PRIMOLUT-N (Other drug) **Reason:** Osteoporosis
Tablet 0.5 Dose Unspecified Daily Oral
Batch: **Started:** 01/01/1991 **Stopped:** CONTIN

CALCITRIOL (Other drug) **Reason:** Osteoporosis
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1991 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87635 **Seq:** 1 **Gender:** F
Reported: 13/09/1993 **Weight:** 36.00
Hospitalisation: **Age:** 60Y
Onset Date: 28/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 26/08/1993 **Stopped:** 28/08/1993

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87804 **Seq:** 1 **Gender:** F
Reported: 22/09/1993 **Weight:** 90.00
Hospitalisation: **Age:** 51Y
Onset Date: 15/05/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thinking abnormal			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 15/04/1993 **Stopped:** 09/09/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87839 **Seq:** 1
Reported: 21/09/1993
Hospitalisation: Admitted to hospital
Onset Date: 03/09/1993
Outcome: Not yet recovered

Gender: M
Weight: 53.00
Age: 76Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Deafness			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 18/05/1993 **Stopped:**

TEMAZEPAM (Suspected) **Reason:** Specific disorders of sleep
Capsule 10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

PIROXICAM (Suspected) **Reason:** Osteoarthritis
Capsule 20.0 Milligram Daily Oral
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87871 **Seq:** 1 **Gender:** F
Reported: 23/09/1993 **Weight:** 62.00
Hospitalisation: **Age:** 31Y
Onset Date: 22/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Palpitations	Severe		

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 75.0 Milligram Daily Oral

Batch: **Started:** 27/07/1993 **Stopped:** 26/08/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87897 **Seq:** 1 **Gender:** M
Reported: 23/09/1993 **Weight:** 60.00
Hospitalisation: **Age:** 41Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erectile dysfunction			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 17/08/1993 **Stopped:** 31/08/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 87956 **Seq:** 1 **Gender:** F
Reported: 28/09/1993 **Weight:** 53.00
Hospitalisation: **Age:** 78Y
Onset Date: 07/09/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Agitation Anorexia Confusional state Flatulence			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 07/09/1993 **Stopped:** 07/09/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had no known previous allergies.

Report Detail

Case Number: 88293 **Seq:** 1 **Gender:** F
Reported: 05/10/1993 **Weight:**
Hospitalisation: **Age:** 40Y
Onset Date: 27/07/1993 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrasystoles Palpitations		Ventricular ectopics.	

Medicine Details:

MORPHINE TARTRATE (Suspected) **Reason:** Premedication
Injection 2.5 Milligram 1 time Intravenous
Batch: **Started:** 27/07/1993 **Stopped:**

LOMOTIL (Suspected) **Reason:** Diarrhea
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

RIVOTRIL (Suspected) **Reason:** Unspecified psychosis
Oral application 500.0 Microgram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 88293 **Seq:** 1 **Gender:** F
Reported: 05/10/1993 **Weight:**
Hospitalisation: **Age:** 40Y
Onset Date: 27/07/1993 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

FLAGYL (Suspected)			**Reason:**		
Suppository	1.5 Gram		Daily		Rectal
Batch:	**Started:**	26/07/1993		**Stopped:**	27/07/1993
PANADOL (Suspected)			**Reason:**	Pain	
Suppository	6.0 Gram		Daily		Rectal
Batch:	**Started:**	26/07/1993		**Stopped:**	29/07/1993

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 88333 **Seq:** 1 **Gender:** M
Reported: 06/10/1993 **Weight:**
Hospitalisation: **Age:** 22Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Nausea	 Severe	Brief loss of conciousness.	

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CANNABIS PREPARATION (Interaction) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 88535 **Seq:** 1 **Gender:** F
Reported: 18/10/1993 **Weight:** 55.00
Hospitalisation: **Age:** 27Y
Onset Date: 25/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nocturia			

Medicine Details:

TRIFEME 28 (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** 23/08/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on aurorix with same reaction occurring. patient has had no known previous allergies.

Report Detail

Case Number: 88730 **Seq:** 1 **Gender:** M
Reported: 28/10/1993 **Weight:**
Hospitalisation: **Age:** 44Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

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

PROSTIN VR (Suspected) **Reason:**
Injection 4.5 Microgram Daily Intravenous
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 88978 **Seq:** 1 **Gender:** M
Reported: 02/11/1993 **Weight:** 60.00
Hospitalisation: **Age:** 59Y
Onset Date: 20/10/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction Skin exfoliation			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 05/10/1993 **Stopped:** 20/10/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 88979 **Seq:** 1 **Gender:** F
Reported: 02/11/1993 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: 14/06/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia			Benztropine prn and

Medicine Details:

ALDAZINE (Other drug) **Reason:** Specific disorders of sleep
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 24/05/1993 **Stopped:**

RIVOTRIL (Other drug) **Reason:** Otr spec symp psychopathol nec
Oral application 2.0 Milligram Daily Oral
Batch: **Started:** 08/06/1993 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 750.0 Milligram Daily Oral
Batch: **Started:** 23/05/1993 **Stopped:** 19/06/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 89058 **Seq:** 1 **Gender:** M
Reported: 04/11/1993 **Weight:** 70.00
Hospitalisation: **Age:** 29Y
Onset Date: 19/07/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension Bradycardia		100/50 lying, 70/50 standing.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 08/07/1993 **Stopped:** 27/07/1993

INDERAL (Suspected) **Reason:** Abnormal involuntary movement
40.0 Milligram Daily
Batch: **Started:** 15/07/1993 **Stopped:** 19/07/1993

PRIADEL (Suspected) **Reason:** Depression
800.0 Milligram Daily
Batch: **Started:** 22/06/1993 **Stopped:** 22/07/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89070 **Seq:** 1 **Gender:** U
Reported: 05/11/1993 **Weight:** 56.00
Hospitalisation: **Age:** 55Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Dry mouth Hyperhidrosis Tremor Visual disturbance			

Medicine Details:

AURORIX (Suspected) **Reason:**
150.0 Milligram 1 time
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had no known previous allergies.

Report Detail

Case Number: 89259 **Seq:** 1 **Gender:** F
Reported: 15/11/1993 **Weight:**
Hospitalisation: **Age:** 47Y
Onset Date: 31/10/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Migraine			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 02/07/1993 **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.

Report Detail

Case Number: 89261 **Seq:** 1 **Gender:** F
Reported: 15/11/1993 **Weight:** 76.00
Hospitalisation: **Age:** 58Y
Onset Date: 01/11/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Dry mouth Nausea Paraesthesia		Head booming.	

Medicine Details:

AURORIX (Suspected) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** 01/11/1993 **Stopped:** 05/11/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89359 **Seq:** 1 **Gender:** F
Reported: 23/11/1993 **Weight:** 60.00
Hospitalisation: **Age:**
Onset Date: 13/11/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Diarrhoea Vomiting	Severe Severe		Gastrolyte 1df 6 times a day

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
1.0 Dose Unspecified Daily
Batch: **Started:** 12/11/1993 **Stopped:** 13/11/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89419 **Seq:** 1 **Gender:** F
Reported: 26/11/1993 **Weight:** 58.00
Hospitalisation: **Age:** 65Y
Onset Date: 15/11/1993 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 27/09/1993 **Stopped:** 15/10/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occured on administering 300mg and again on 150mg_daily

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89446 **Seq:** 1 **Gender:** F
Reported: 29/11/1993 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 17/11/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 16/11/1993 **Stopped:** 20/11/1993

TELDANE (Suspected) **Reason:** Chronic nasopharyngitis
Tablet 120.0 Milligram Daily Oral
Batch: **Started:** 17/11/1993 **Stopped:** 19/11/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 89457 **Seq:** 1 **Gender:** F
Reported: 29/11/1993 **Weight:**
Hospitalisation: **Age:** 39
Onset Date: 06/11/1993 **DOB:** 06/08/1954
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			
Hyperhidrosis			
Hypertonia			
Myalgia			No response to cogentin, diazepam, and phenytoin so given
Mydriasis			
Pupillary reflex impaired			
Pyrexia			
Respiratory disorder			
Salivary hypersecretion			

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 01/11/1993 **Stopped:**

IMIPRAMINE HYDROCHLORIDE (Interaction) **Reason:**
200.0 Milligram Daily
Batch: **Started:** 01/11/1993 **Stopped:**

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine [illegible] Radiololgy		07/11/1993	3,200 U/L	Serum tricyclics - 1940 mcg/l.

Additional Information:

Alcoholism. 1992 bipolar affective disorder. paroxysmal supraventricular tachycardia controlled on verapamil and digoxin. still some residual calf and thigh discomfort, flushing, shivering, amnesia.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89457 **Seq:** 1 **Gender:** F
Reported: 29/11/1993 **Weight:**
Hospitalisation: **Age:** 39
Onset Date: 06/11/1993 **DOB:** 06/08/1954
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Speech disorder Stupor			Chlorpromazine, penicillin and ceftriaxone with some improvement

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
7.5 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Alcoholism. 1992 bipolar affective disorder. paroxysmal supraventricular tachycardia controlled on verapamil and digoxin. still some residual calf and thigh discomfort, flushing, shivering, amnesia.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89580 **Seq:** 1 **Gender:** F
Reported: 01/12/1993 **Weight:** 47.00
Hospitalisation: **Age:** 88Y
Onset Date: 20/10/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Hyperkinesia			

Medicine Details:

OXAZEPAM (Other drug) **Reason:** Anxiety neurosis
15.0 Milligram Daily
Batch: **Started:** 01/10/1993 **Stopped:**

ASPIRIN (Other drug) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 15/10/1993 **Stopped:** 16/11/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89818 **Seq:** 1 **Gender:** M
Reported: 13/12/1993 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 26/11/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Palpitations			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 25/02/1993 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 89835 **Seq:** 1 **Gender:** F
Reported: 13/12/1993 **Weight:** 60.00
Hospitalisation: **Age:** 32Y
Onset Date: 27/11/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:**

300.0 Milligram Daily

Batch: **Started:** **Stopped:** 2D

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89844 **Seq:** 1
Reported: 14/12/1993
Hospitalisation:
Onset Date: 14/10/1993
Outcome: Recovered with sequelae

Gender: F
Weight: 43.00
Age: 24Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuroleptic malignant syndrome			Bromocriptine

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 08/10/1993 **Stopped:** 15/10/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

"(patient exhibited) muscle rigidity, tremulousness of upper extremities, and was mute. (also characterised by) elevated body temperature, elevated cpk levels, elevated urea/creatinine ratio, elevated white cell count. also a transient elevation of the asot."

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 89898 **Seq:** 1
Reported: 02/12/1993
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 64.00
Age: 82Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

Batch: **Started:** **Stopped:** 01M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 90031 **Seq:** 1 **Gender:** F
Reported: 24/12/1993 **Weight:** 68.00
Hospitalisation: **Age:** 80Y
Onset Date: 07/12/1993 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

VERACAPS SR (Other drug) **Reason:**
Capsule 160.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

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

UREX (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 04/12/1993 **Stopped:** 08/12/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90545 **Seq:** 1
Reported: 18/01/1994
Hospitalisation:
Onset Date: 14/01/1994
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 52Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Flushing Palpitations	Severe	Rash on chest,back,face,and arms.	

Medicine Details:

ATIVAN (Suspected) **Reason:**
1.0 Milligram As necessary
Batch: **Started:** 11/01/1994 **Stopped:** 13/01/1994

AURORIX (Suspected) **Reason:** Depression
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 14/01/1994 **Stopped:** 14/01/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90561 **Seq:** 1 **Gender:** F
Reported: 19/01/1994 **Weight:** 60.00
Hospitalisation: **Age:** 62Y
Onset Date: 01/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Conjunctivitis		Eyes swollen and burned	
Salivary hypersecretion		Mucus in throat	
Dysphonia			Antihistamine, adrenaline.
Pruritus			
Rash			
Rhinitis			

Medicine Details:

PREMARIN (Other drug) **Reason:** Menopausal symptoms
0.0
Batch: **Started:** 01/12/1993 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:**
300.0 Milligram 1 time
Batch: **Started:** 01/01/1994 **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.

Report Detail

Case Number: 90563 **Seq:** 1 **Gender:** F
Reported: 20/01/1994 **Weight:**
Hospitalisation: **Age:** 79Y
Onset Date: 13/08/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vasculitis Red blood cell sedimentation rate		Vasculitic skin lesions	

Medicine Details:

ENALAPRIL MALEATE (Other drug) **Reason:** Congestive heart failure
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 24/08/1993

RANITIDINE (Suspected) **Reason:** Peptic ulcer nos w/o ment perf
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 14/08/1993

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 07/08/1993 **Stopped:** 14/08/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90770 **Seq:** 1
Reported: 31/01/1994
Hospitalisation:
Onset Date: 22/01/1994
Outcome: Not yet recovered

Gender: M
Weight: 102.00
Age: 57Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash maculo-papular			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 4.0 Dose Unspecified Daily Oral

Batch: **Started:** 17/01/1994 **Stopped:** 22/01/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90791 **Seq:** 1 **Gender:** F
Reported: 31/01/1994 **Weight:** 87.00
Hospitalisation: **Age:** 39Y
Onset Date: 13/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myalgia	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 12/01/1994 **Stopped:** 19/01/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90859 **Seq:** 1 **Gender:** M
Reported: 02/02/1994 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia			Surgical excision.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 27/04/1993 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90930 **Seq:** 1 **Gender:** M
Reported: 07/02/1994 **Weight:** 83.00
Hospitalisation: **Age:** 48Y
Onset Date: 11/11/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Ejaculation failure			

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 18/09/1986 **Stopped:**

ZYLOPRIM (Other drug) **Reason:** Gout
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/09/1985 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 12/03/1993 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90941 **Seq:** 1 **Gender:** M
Reported: 07/02/1994 **Weight:**
Hospitalisation: **Age:** 44
Onset Date: 24/01/1994 **DOB:** 24/01/1950
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Prothrombin level decreased			

Medicine Details:

AURORIX (Interaction) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 21/01/1994 **Stopped:**

WARFARIN SODIUM (Interaction) **Reason:**
5.0 Milligram Daily
Batch: **Started:** 01/07/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prothrombin time		24/01/1994	2.4	Usual warfarin-prothrombin time ratio=2.0 (2.0-2.3).
	Prothrombin time		02/02/1994	3.7	Usual warfarin-prothrombin time ratio=2.0 (2.0-2.3).

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90953 **Seq:** 1 **Gender:** M
Reported: 09/02/1994 **Weight:** 85.00
Hospitalisation: **Age:** 60Y
Onset Date: 06/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence	Severe		

Medicine Details:

ISOPTIN (Other drug)			**Reason:** Essential benign hypertension		
Tablet	40.0	Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1990			**Stopped:**	CONTIN
ZANTAC (Other drug)			**Reason:**		
Tablet	600.0	Milligram	Daily	Oral	
Batch:	**Started:** 23/12/1993			**Stopped:**	11/01/1994
AURORIX (Suspected)			**Reason:** Depression		
Tablet	150.0	Milligram	1 time	Oral	
Batch:	**Started:** 06/01/1994			**Stopped:**	07/01/1994
FLAGYL (Suspected)			**Reason:**		
Oral application	600.0	Milligram	Daily	Oral	
Batch:	**Started:** 04/01/1994			**Stopped:**	11/01/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 90984 **Seq:** 1 **Gender:** F
Reported: 09/02/1994 **Weight:** 70.00
Hospitalisation: **Age:** 56Y
Onset Date: 01/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia			

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

OXAZEPAM (Suspected) **Reason:**
360.0 Milligram Daily
Batch: **Started:** **Stopped:** 31/01/1994

LITHIUM CARBONATE (Suspected) **Reason:** Depression
500.0 Milligram Daily
Batch: **Started:** **Stopped:** 01/02/1994

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 09/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Neutrophils	1.1-7.8	01/02/1994	1600	
	Neutrophils	1.1-7.8	04/02/1994	2100	
	Neutrophils	1.1-7.8	07/02/1994	1400	
	Neutrophils	1.1-7.8	11/02/1994	2700	
	Neutrophils	1.1-7.8	14/02/1994	3900	
	Neutrophils	1.1-7.8	09/02/1994	1700	
	White blood cells	4000 - 11000			Wcc 4,4000/ul,on cessation serepax(detox with diazepam) wcc

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91192 **Seq:** 1 **Gender:** F
Reported: 21/02/1994 **Weight:** 55.00
Hospitalisation: **Age:** 37Y
Onset Date: 25/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 24/01/1994 **Stopped:** 28/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Itchy rash on face occured 2 days after commencing aurorix.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91285 **Seq:** 1 **Gender:** F
Reported: 25/02/1994 **Weight:** 52.00
Hospitalisation: **Age:** 76Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anaemia Bone marrow failure	Severe		

Medicine Details:

LASIX (Other drug) **Reason:** Congestive heart failure
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

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

LOPID (Suspected) **Reason:** Othr&unspec metabolic diseases
1.2 Gram Daily
Batch: **Started:** **Stopped:** 20/01/1994

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 13/01/1994 **Stopped:** 20/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haemoglobin				Hb 8.1

Additional Information:

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

Report Detail

Case Number: 91285 **Seq:** 1 **Gender:** F
Reported: 25/02/1994 **Weight:** 52.00
Hospitalisation: **Age:** 76Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TOLVON (Suspected)		**Reason:**	Depression
	40.0 Milligram	Daily	
Batch:	**Started:** 01/12/1993	**Stopped:**	20/01/1994
RENITEC (Suspected)		**Reason:**	Congestive heart failure
	20.0 Milligram	Daily	
Batch:	**Started:**	**Stopped:**	20/01/1994

Laboratory Investigations:

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91397 **Seq:** 1 **Gender:** F
Reported: 03/03/1994 **Weight:** 53.00
Hospitalisation: **Age:** 38Y
Onset Date: 26/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal			
Diplopia			
Hyperacusis			
Nausea			
Tetany			
Visual disturbance			

Medicine Details:

CODRAL TENSION HEADACHE (Other drug) **Reason:** Headache
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Total Oral
Batch: **Started:** 26/02/1994 **Stopped:** 26/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No previous reaction to drugs. past history of asthma and depression

Report Detail

Case Number: 91399 **Seq:** 1 **Gender:** F
Reported: 04/03/1994 **Weight:** 47.00
Hospitalisation: **Age:** 32Y
Onset Date: 31/12/1993 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea	Severe		
Vomiting	Severe		

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:** Nausea and vomiting
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/12/1992 **Stopped:** 28/02/1994

METHADONE HYDROCHLORIDE (Interaction) **Reason:** Pain
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** 01/12/1993 **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 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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91412 **Seq:** 1 **Gender:** F
Reported: 07/03/1994 **Weight:** 80.00
Hospitalisation: **Age:** 29Y
Onset Date: 31/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 01/12/1993 **Stopped:** 21/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Not warned to avoid tyramine containing food - apparently this is not in literature in australia (it is in th uk). - 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91436 **Seq:** 1 **Gender:** M
Reported: 08/03/1994 **Weight:** 58.00
Hospitalisation: **Age:** 90Y
Onset Date: 02/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia	Severe		

Medicine Details:

FLUOXETINE HYDROCHLORIDE (Suspected)			**Reason:** Depression	
Capsule	20.0	Milligram	Daily	Oral
Batch:	**Started:** 17/12/1993			**Stopped:** 02/02/1994
DILANTIN (Suspected)			**Reason:** Convulsions	
Capsule	200.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:** 07/12/1993
MOCLOBEMIDE (Suspected)			**Reason:** Depression	
Tablet	600.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:** 07/12/1993
LITHIUM CARBONATE (Suspected)			**Reason:** Depression	
Tablet	500.0	Milligram	Daily	Oral
Batch:	**Started:** 01/12/1993			**Stopped:** 07/12/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium				116

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91494 **Seq:** 1 **Gender:** M
Reported: 09/03/1994 **Weight:** 84.00
Hospitalisation: **Age:** 65Y
Onset Date: 01/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia Headache		Tingling hands and legs	

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** 08/07/1993 **Stopped:**

ZANTAC (Other drug) **Reason:** Otr disorders stomach function
300.0 Milligram Daily
Batch: **Started:** 10/11/1993 **Stopped:**

LIPEX (Other drug) **Reason:** Othr&unspec metabolic diseases
10.0 Milligram Daily
Batch: **Started:** 01/01/1991 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 10/11/1993 **Stopped:** 18/01/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91540 **Seq:** 1 **Gender:** F
Reported: 11/03/1994 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression Abdominal pain Anxiety Malaise Nausea Syncope		Tearfulness	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Anxiety neurosis
300.0 Milligram Daily
Batch: **Started:** 17/02/1994 **Stopped:** 23/02/1994

TRIMETHOPRIM (Suspected) **Reason:** Othr diseases of urinary tract
300.0 Milligram Daily
Batch: **Started:** 18/02/1994 **Stopped:** 23/02/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 91559 **Seq:** 1 **Gender:** M
Reported: 14/03/1994 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: 22/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle twitching		Tourette syndrome	Small doses of alprazolam

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 15/03/1993 **Stopped:** 22/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 92273 **Seq:** 1 **Gender:** F
Reported: 18/04/1994 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: 22/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Nausea			

Medicine Details:

TEMAZEPAM (Other drug) **Reason:** Specific disorders of sleep
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 22/02/1994 **Stopped:** 28/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Soon after taking each dose of aurorix same reaction occurred. symptoms stopped immediately after stopping treatment.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 92351 **Seq:** 1 **Gender:** F
Reported: 18/04/1994 **Weight:** 75.00
Hospitalisation: **Age:** 67Y
Onset Date: 03/03/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 03/03/1994 **Stopped:** 06/03/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 92404 **Seq:** 1 **Gender:** M
Reported: 21/04/1994 **Weight:** 70.00
Hospitalisation: **Age:** 41Y
Onset Date: 10/02/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertonia Confusional state Coordination abnormal Depersonalisation Dry mouth Nausea		Tightness in legs.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
3.0 Dose Unspecified Daily
Batch: **Started:** 09/02/1994 **Stopped:** 15/03/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 92675 **Seq:** 1 **Gender:** F
Reported: 02/05/1994 **Weight:** 55.00
Hospitalisation: **Age:** 37Y
Onset Date: 24/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Face oedema Hepatic function abnormal Malaise		Hand swelling	

Medicine Details:

INSULIN NOS (Other drug) **Reason:** Diabetes mellitus
0.0
Batch: **Started:** L TERM **Stopped:**

VERACAPS SR (Other drug) **Reason:** Essential benign hypertension
240.0 Milligram Daily
Batch: **Started:** 01/01/1993 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 15/01/1994 **Stopped:** 17/02/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	(N. 0-60)	17/02/1994	137	
	GGT = SGGT = SGTP	(N 0-80)	17/02/1994	177	
	SAP = ALP	(N 28-126)	17/02/1994	178	

Additional Information:

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

Report Detail

Case Number: 92804 **Seq:** 1 **Gender:** F
Reported: 09/05/1994 **Weight:**
Hospitalisation: **Age:** 66
Onset Date: 17/03/1994 **DOB:** 18/09/1927
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state	Severe		
Dizziness	Severe		
Eosinophilia	Severe		
Nausea	Severe		
Orthostatic hypotension	Severe		
Pyrexia	Severe		
Red blood cell sedimentation rate	Severe		
Somnolence	Severe		Antipyretics
Vomiting	Severe		

Medicine Details:

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

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

RIVOTRIL (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** **Stopped:** 10/04/1994

CLOZAPINE (Suspected) **Reason:** Unspecified schizophrenia
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 16/03/1994 **Stopped:** 04/04/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 92804 **Seq:** 1 **Gender:** F
Reported: 09/05/1994 **Weight:**
Hospitalisation: **Age:** 66
Onset Date: 17/03/1994 **DOB:** 18/09/1927
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression
Tablet	600.0	Milligram	Daily	Oral
Batch:	**Started:**	16/03/1994	**Stopped:**	23/03/1994
MOCLOBEMIDE (Suspected)			**Reason:**	Depression
Tablet	300.0	Milligram	Daily	Oral
Batch:	**Started:**	16/03/1994	**Stopped:**	27/03/1994
MOCLOBEMIDE (Suspected)			**Reason:**	Depression
Tablet	150.0	Milligram	Daily	Oral
Batch:	**Started:**	16/03/1994	**Stopped:**	27/03/1994
PROPRANOLOL HYDROCHLORIDE (Suspected)			**Reason:**	Disturbance of coordination
Tablet	30.0	Milligram	Daily	Oral
Batch:	**Started:**	17/03/1994	**Stopped:**	18/03/1994

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 92857 **Seq:** 1 **Gender:** M
Reported: 10/05/1994 **Weight:** 93.00
Hospitalisation: **Age:** 44Y
Onset Date: 26/04/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Prothrombin level decreased			

Medicine Details:

AURORIX (Interaction) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

MAREVAN (Interaction) **Reason:**
4.0 Milligram Daily
Batch: **Started:** 26/04/1994 **Stopped:**

PARACETAMOL (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prothrombin ratio				Prothrombin time ratio increased from ~ 2.0 to 3.7 after beginning to take aurorix then stablising warfarin dose, aurorix was withdrawn and prothrombin time ratio decreased from ~ 2.3 to 1.8

Additional Information:

Report Detail

Case Number: 92906 **Seq:** 1 **Gender:** F
Reported: 12/05/1994 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 08/03/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Hypertension Visual disturbance			

Medicine Details:

DOXEPIN HYDROCHLORIDE (Other drug) **Reason:** Depression

Capsule 50.0 Milligram Daily Oral

Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 11/11/1993 **Stopped:** 08/03/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 93124 **Seq:** 1 **Gender:** F
Reported: 17/05/1994 **Weight:**
Hospitalisation: **Age:** 61
Onset Date: 15/11/1993 **DOB:** 02/10/1932
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 05/11/1992 **Stopped:** 08/11/1992

MOCLOBEMIDE (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 09/11/1993 **Stopped:** 15/11/1993

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 93145 **Seq:** 1 **Gender:** F
Reported: 25/05/1994 **Weight:** 60.00
Hospitalisation: **Age:** 38Y
Onset Date: 12/05/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Euphoric mood Hyperkinesia Insomnia Mania		Elevated mood Over activity.	

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:** Unspecified schizophrenia
Tablet 1.2 Gram Daily Oral
Batch: **Started:** 01/11/1991 **Stopped:**

AURORIX (Suspected) **Reason:**
450.0 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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 93163 **Seq:** 1 **Gender:** F
Reported: 23/05/1994 **Weight:** 64.00
Hospitalisation: **Age:** 36Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mouth ulceration			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/11/1993 **Stopped:** 18/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Aurorix has been tried on two occasions - ulcers rear when it is started, worsen if dose is increased to 3/day.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 93190 **Seq:** 1 **Gender:** F
Reported: 26/05/1994 **Weight:** 74.00
Hospitalisation: **Age:** 37Y
Onset Date: 23/05/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 20/05/1994 **Stopped:** 22/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergic to bactrim and buscopan.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 93619 **Seq:** 1 **Gender:** F
Reported: 10/06/1994 **Weight:** 65.00
Hospitalisation: **Age:** 28Y
Onset Date: 03/03/1994 **DOB:**
Outcome: Recovered without treatment **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 03/03/1994 **Stopped:** 14/03/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93660 **Seq:** 1 **Gender:** F
Reported: 06/06/1994 **Weight:**
Hospitalisation: **Age:** 45
Onset Date: 22/05/1994 **DOB:** 06/03/1949
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium Pallor Tremor			

Medicine Details:

PAROXETINE HYDROCHLORIDE (Suspected) **Reason:** Depression
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** 06/05/1994 **Stopped:** 20/05/1994

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 22/05/1994 **Stopped:** 22/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93771 **Seq:** 1 **Gender:** F
Reported: 17/06/1994 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Eosinophilia Pruritus Rash morbilliform			

Medicine Details:

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

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93791 **Seq:** 1 **Gender:** M
Reported: 20/06/1994 **Weight:** 60.00
Hospitalisation: **Age:** 83Y
Onset Date: 16/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing Pruritus Rash erythematous			Phomethazine 25mg. im.

Medicine Details:

MORPHINE NOS (Other drug) **Reason:** Pain
Injection 0.0 Intramuscular
Batch: **Started:** 13/06/1994 **Stopped:**

MERSYNDOL FORTE (Other drug) **Reason:** Pain
Tablet 6.0 Dose Unspecified Daily Oral
Batch: **Started:** 08/06/1994 **Stopped:**

LANOXIN (Other drug) **Reason:**
Tablet 250.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

MURELAX (Other drug) **Reason:**
Tablet 15.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93791 **Seq:** 1 **Gender:** M
Reported: 20/06/1994 **Weight:** 60.00
Hospitalisation: **Age:** 83Y
Onset Date: 16/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TETREX (Other drug)			**Reason:**		
Capsule	2.5	Dose Unspecified	Daily		Oral
Batch:	**Started:**	L TERM		**Stopped:**	
THEO-DUR (Other drug)			**Reason:**		
Tablet	600.0	Milligram	Daily		Oral
Batch:	**Started:**	L TERM		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	2.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**	14/06/1994		**Stopped:**	16/06/1994

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 93902 **Seq:** 1 **Gender:** M
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: 26/10/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia			

Medicine Details:

PREDNISOLONE (Other drug) **Reason:** Otr respiratory systm diseases
Oral application 0.0 Reducing Oral
Batch: **Started:** 20/10/1993 **Stopped:**

VENTOLIN (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 5.0 Milligram Daily Inhalation
Batch: **Started:** 20/10/1993 **Stopped:**

ATROVENT (Other drug) **Reason:** Otr respiratory systm diseases
Inhalation 3.0 Milligram Daily Inhalation
Batch: **Started:** 20/10/1993 **Stopped:**

ZANTAC (Other drug) **Reason:** Peptic ulcer nos w/o ment perf
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 20/10/1993 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 93902 **Seq:** 1 **Gender:** M
Reported: 24/06/1994 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: 26/10/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

THEO-DUR (Other drug)			**Reason:**	Otr respiratory systm diseases
Tablet	600.0	Milligram	Daily	Oral
Batch:	**Started:**	20/10/1993		**Stopped:**
AUGMENTIN FORTE (Other drug)			**Reason:**	Otr respiratory systm diseases
Tablet	1.5	Gram	Daily	Oral
Batch:	**Started:**	20/10/1993		**Stopped:**
AURORIX (Suspected)			**Reason:**	Depression
	0.0			
Batch:	**Started:**	20/10/1993		**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 94013 **Seq:** 1 **Gender:** F
Reported: 29/06/1994 **Weight:** 94.00
Hospitalisation: **Age:** 46Y
Onset Date: 20/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous Pruritus	Severe Severe	Blotchy rash over torso	

Medicine Details:

PROVERA (Other drug) **Reason:**
Tablet 2.5 Milligram Daily Oral
Batch: **Started:** 01/01/1992 **Stopped:**

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

SALAZOPYRIN EN (Other drug) **Reason:**
6.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1993 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 16/06/1994 **Stopped:** 21/06/1994

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.

Report Detail

Case Number: 94387 **Seq:** 1 **Gender:** F
Reported: 14/07/1994 **Weight:** 70.00
Hospitalisation: **Age:** 33Y
Onset Date: 06/07/1994 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Nausea Vomiting			

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:** Depression
Tablet 750.0 Milligram Daily Oral
Batch: **Started:** 01/04/1994 **Stopped:**

STELAZINE (Other drug) **Reason:** Depression
Oral application 1.0 Milligram Daily Oral
Batch: **Started:** 08/06/1994 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** 21/06/1994 **Stopped:** 07/07/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient rechallenged on aurorix with same reaction. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 94483 **Seq:** 1 **Gender:** M
Reported: 21/07/1994 **Weight:** 83.00
Hospitalisation: **Age:** 73Y
Onset Date: 25/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Diarrhoea Nausea			

Medicine Details:

SULINDAC (Other drug) **Reason:** Unspecified arthritis
400.0 Milligram Daily
Batch: **Started:** 01/01/1993 **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
100.0 Microgram Daily
Batch: **Started:** 01/08/1993 **Stopped:**

IMDUR (Other drug) **Reason:** Chron isch heart dis no hyper
60.0 Milligram Daily
Batch: **Started:** 01/09/1993 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 24/06/1994 **Stopped:** 30/06/1994

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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 94538 **Seq:** 1 **Gender:** F
Reported: 26/07/1994 **Weight:** 69.00
Hospitalisation: **Age:** 46Y
Onset Date: 20/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension Paraesthesia			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 750.0 Milligram Daily Oral

Batch: **Started:** 06/05/1994 **Stopped:** 06/07/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 94660 **Seq:** 1 **Gender:** F
Reported: 28/07/1994 **Weight:** 51.00
Hospitalisation: **Age:** 41Y
Onset Date: 14/07/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia			
Dry skin			
Headache			
Hot flush			
Hypotension			
Insomnia			
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 13/07/1994 **Stopped:** 17/07/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 94751 **Seq:** 1 **Gender:** F
Reported: 01/08/1994 **Weight:** 74.00
Hospitalisation: **Age:** 30Y
Onset Date: 23/06/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Nightmare			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/06/1994 **Stopped:** 30/06/1994

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 Detai

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: 95230 **Seq:** 1 **Gender:** M
Reported: 22/08/1994 **Weight:** 180.00
Hospitalisation: **Age:** 36Y
Onset Date: 20/07/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus			

Medicine Details:

AURORIX (Suspected) **Reason:**
4.0 Dose Unspecified Daily
Batch: **Started:** 06/07/1994 **Stopped:** 08/08/1994

CISAPRIDE (Suspected) **Reason:**
6.0 Dose Unspecified Daily
Batch: **Started:** 06/07/1994 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95317 **Seq:** 1 **Gender:** M
Reported: 24/08/1994 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: 09/07/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 750.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95400 **Seq:** 1 **Gender:** F
Reported: 31/08/1994 **Weight:** 78.00
Hospitalisation: **Age:** 37Y
Onset Date: 20/08/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Depersonalisation Gait disturbance			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 17/08/1992 **Stopped:** 22/08/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 95418 **Seq:** 1 **Gender:** F
Reported: 31/08/1994 **Weight:** 70.00
Hospitalisation: **Age:** 70Y
Onset Date: 12/07/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Dizziness Nausea			

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

PEPCIDINE (Interaction) **Reason:** Ulcer of stomach w/o ment perf
40.0 Milligram Daily
Batch: **Started:** 12/07/1994 **Stopped:**

PANAMAX (Other drug) **Reason:** Depression
1.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

PROTHIADEN (Other drug) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 95643 **Seq:** 1 **Gender:** F
Reported: 02/09/1994 **Weight:** 55.00
Hospitalisation: **Age:** 29Y
Onset Date: 08/08/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mania			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 19/07/1994 **Stopped:** 08/08/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95687 **Seq:** 1 **Gender:** F
Reported: 07/09/1994 **Weight:**
Hospitalisation: **Age:** 84Y
Onset Date: 17/05/1994 **DOB:**
Outcome: Recovered with sequelae **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cerebrovascular disorder			

Medicine Details:

VALIUM (Suspected) **Reason:** Anxiety neurosis
5.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:** 17/05/1994

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 14/05/1994 **Stopped:** 17/05/1994

ASPIRIN (Suspected) **Reason:** Otr peripheral vasculr disease
150.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 17/05/1994

CAPOTEN (Suspected) **Reason:** Essential benign hypertension
50.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 17/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: short term memory deficits and nominal dysphasia mild.

Report Detail

Case Number: 95840 **Seq:** 1 **Gender:** M
Reported: 12/09/1994 **Weight:** 70.00
Hospitalisation: **Age:** 52Y
Onset Date: 27/08/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nightmare Paraesthesia			

Medicine Details:

DOLOBID (Other drug) **Reason:**
500.0 Milligram Daily
Batch: **Started:** 26/08/1994 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 26/08/1994 **Stopped:** 02/09/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Similar reaction to tofranil which he had about 10 years ago.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95923 **Seq:** 1 **Gender:** M
Reported: 19/09/1994 **Weight:** 75.00
Hospitalisation: **Age:** 45Y
Onset Date: 22/08/1994 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing Influenza like illness			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 17/08/1994 **Stopped:** 22/08/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on aurorix with same reaction occurring.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95968 **Seq:** 1 **Gender:** F
Reported: 19/09/1994 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Coordination abnormal		Jerking movements of limbs	

Medicine Details:

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

SLOW-K (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

CARDIPRIN 100 (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

LANOXIN (Other drug) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 95968 **Seq:** 1 **Gender:** F
Reported: 19/09/1994 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

LASIX (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
AURORIX (Suspected)		**Reason:**	Depression
	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 96192 **Seq:** 1 **Gender:** F
Reported: 30/09/1994 **Weight:** 62.00
Hospitalisation: **Age:** 58Y
Onset Date: 20/09/1994 **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain			

Medicine Details:

LOSEC (Other drug) **Reason:**
Capsule 0.0 Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

PANAMAX (Other drug) **Reason:**
Oral application 0.0 Oral
Batch: **Started:** **Stopped:** CONTIN

ESTRADERM (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:** CONTIN

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred with every dose of aurorix.

Report Detail

Case Number: 96460 **Seq:** 1 **Gender:** M
Reported: 11/10/1994 **Weight:** 98.00
Hospitalisation: **Age:** 27Y
Onset Date: 28/09/1994 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Ejaculation failure			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** 23/09/1994 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 96653 **Seq:** 1 **Gender:** F
Reported: 24/10/1994 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 07/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

OESTROGENS NOS (Suspected) **Reason:** Menopausal symptoms
Cream 0.0 Topical
Batch: **Started:** **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 01/05/1994 **Stopped:** 14/07/1994

CALCITRIOL (Suspected) **Reason:** Osteoporosis
1.0 Milligram Daily
Batch: **Started:** 16/05/1994 **Stopped:** CONTIN

TEMAZEPAM (Suspected) **Reason:**
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		07/06/1994	48	
	ALT = SGPT		03/07/1994	47	
	AST = SGOT		07/06/1994	52	
	Bilirubin		07/06/1994	9	
	Bilirubin		03/07/1994	8	
	GGT = SGGT =		07/06/1994	153	

Additional Information:

Abuse of alcohol, sedatives, aperients. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 96653 **Seq:** 1 **Gender:** F
Reported: 24/10/1994 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 07/06/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

OESTROGENS NOS (Suspected)			**Reason:**	Menopausal symptoms	
Cream	0.0			Topical	
Batch:	**Started:**			**Stopped:**	CONTIN
MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
Tablet	450.0 Milligram		Daily	Oral	
Batch:	**Started:** 01/05/1994			**Stopped:** 14/07/1994	
CALCITRIOL (Suspected)			**Reason:**	Osteoporosis	
	1.0 Milligram		Daily		
Batch:	**Started:** 16/05/1994			**Stopped:**	CONTIN
TEMAZEPAM (Suspected)			**Reason:**		
Oral application	20.0 Milligram		Daily	Oral	
Batch:	**Started:**			**Stopped:**	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Liver biopsy				Needle biopsy of liver - 12/7/94 - mixed pattern of moderate fatty change and evidence of recent scattered liver cell necrosis. the pattern is consistent with drug-induced reaction and alcohol induced liver damage. the pattern of the liver cell damage is not that usually seen with alcohol but fatty change and megamitochondria are consistent with the latter.
	SAP = ALP		07/06/1994	147	
	SAP = ALP		03/07/1994	126	

Additional Information:

Abuse of alcohol, sedatives, aperients. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 97409 **Seq:** 1 **Gender:** F
Reported: 28/11/1994 **Weight:** 80.00
Hospitalisation: **Age:** 30Y
Onset Date: 19/11/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash vesicular Urticaria		Rash on anterior elbow and upper arm.	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 03/11/1994 **Stopped:** 20/11/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 97622 **Seq:** 1 **Gender:** M
Reported: 09/12/1994 **Weight:** 88.00
Hospitalisation: **Age:** 65Y
Onset Date: 30/11/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea		Up to 5 bowel actions daily.	

Medicine Details:

ZANTAC (Other drug) **Reason:** Otr disorders stomach function
300.0 Milligram Daily
Batch: **Started:** 01/01/1990 **Stopped:**

BURINEX (Other drug) **Reason:** Congestive heart failure
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1990 **Stopped:**

MINIDIAB (Other drug) **Reason:** Diabetes mellitus
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1994 **Stopped:**

WARFARIN SODIUM (Other drug) **Reason:**
5.5 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 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. 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: 97622 **Seq:** 1
Reported: 09/12/1994
Hospitalisation:
Onset Date: 30/11/1994
Outcome: Not yet recovered

Gender: M
Weight: 88.00
Age: 65Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

LANOXIN (Other drug)		**Reason:**	Congestive heart failure
	250.0 Microgram	Daily	
Batch:	**Started:** 01/01/1990	**Stopped:**	
MOCLOBEMIDE (Suspected)		**Reason:**	Depression
	300.0 Milligram	Daily	
Batch:	**Started:** 29/11/1994	**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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 97829 **Seq:** 1 **Gender:** F
Reported: 14/12/1994 **Weight:**
Hospitalisation: **Age:** 72Y
Onset Date: 19/11/1994 **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cardiomyopathy Left ventricular failure			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 12/10/1994 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 97836 **Seq:** 1 **Gender:** M
Reported: 12/12/1994 **Weight:** 105.00
Hospitalisation: **Age:** 41Y
Onset Date: 15/11/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			

Medicine Details:

LITHIUM CARBONATE (Other drug) **Reason:**
1.7 Gram Daily
Batch: **Started:** L TERM **Stopped:**

TEGRETOL (Other drug) **Reason:**
1.0 Gram Daily
Batch: **Started:** L TERM **Stopped:**

NEULACTIL (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/08/1994 **Stopped:** 26/11/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 98024 **Seq:** 1 **Gender:** F
Reported: 29/12/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 20/10/1994 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia	Severe		

Medicine Details:

CALTRATE (Suspected) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

DIGOXIN (Suspected) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

LASIX (Suspected) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 03/10/1994 **Stopped:** 14/12/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 98024 **Seq:** 1 **Gender:** F
Reported: 29/12/1994 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 20/10/1994 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

ZANTAC (Suspected)			**Reason:**	
	300.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
CAPTOPRIL (Suspected)			**Reason:**	
	75.0	Milligram	Daily	
Batch:	**Started:**		L TERM	**Stopped:**

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 98054 **Seq:** 1 **Gender:** F
Reported: 29/12/1994 **Weight:** 70.00
Hospitalisation: **Age:** 65Y
Onset Date: 14/12/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal		Cerebellar ataxia.	

Medicine Details:

SEREPAX (Other drug) **Reason:** Anxiety neurosis
30.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
50.0 Microgram Daily
Batch: **Started:** 07/11/1994 **Stopped:**

HYGROTON (Other drug) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

LOPRESOR (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98054 **Seq:** 1 **Gender:** F
Reported: 29/12/1994 **Weight:** 70.00
Hospitalisation: **Age:** 65Y
Onset Date: 14/12/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 08/12/1994 **Stopped:** 14/12/1994

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98231 **Seq:** 1 **Gender:** M
Reported: 10/01/1995 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia			
Dizziness			
Fatigue			
Nausea			
Tremor			

Medicine Details:

AROPAX (Interaction) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 22/07/1994 **Stopped:** 18/10/1994

AURORIX (Interaction) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 30/12/1993 **Stopped:** 21/07/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98314 **Seq:** 1 **Gender:** F
Reported: 13/01/1995 **Weight:** 66.00
Hospitalisation: **Age:** 71Y
Onset Date: 14/12/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous Pruritus		Rash on shoulders, abdomen and back.	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Alternating-other
Batch: **Started:** 01/12/1994 **Stopped:** 31/12/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 98387 **Seq:** 1 **Gender:** F
Reported: 18/01/1995 **Weight:**
Hospitalisation: Admitted to hospital **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Suicide attempt			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98459 **Seq:** 1 **Gender:** M
Reported: 23/01/1995 **Weight:** 87.00
Hospitalisation: **Age:** 66Y
Onset Date: 06/01/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thinking abnormal Aggression Agitation Dizziness Drug ineffective Dyspnoea Myalgia Tachycardia Tinnitus		Inability to think	

Medicine Details:

DUCENE (Other drug) **Reason:**
Tablet 1.0 Milligram As necessary Oral
Batch: **Started:** 19/12/1994 **Stopped:**

DIAZEPAM (Other drug) **Reason:**
Tablet 5.0 Milligram As necessary Oral
Batch: **Started:** L TERM **Stopped:**

SURMONTIL (Other drug) **Reason:**
Capsule 25.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

BETALOC (Other drug) **Reason:**
Tablet 25.0 Milligram Daily Oral
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98459 **Seq:** 1 **Gender:** M
Reported: 23/01/1995 **Weight:** 87.00
Hospitalisation: **Age:** 66Y
Onset Date: 06/01/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 06/01/1995 **Stopped:** 06/01/1995

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98486 **Seq:** 1 **Gender:** F
Reported: 24/01/1995 **Weight:** 57.00
Hospitalisation: **Age:** 73Y
Onset Date: 31/10/1994 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea			

Medicine Details:

DUCENE (Other drug) **Reason:** Anxiety neurosis
5.0 Milligram As necessary
Batch: **Started:** 01/01/1992 **Stopped:**

ASPIRIN (Other drug) **Reason:** Transient cereb isch no hypert
150.0 Milligram Daily
Batch: **Started:** 01/01/1992 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/09/1994 **Stopped:** 05/01/1995

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98616 **Seq:** 1 **Gender:** M
Reported: 31/01/1995 **Weight:** 78.00
Hospitalisation: **Age:** 84Y
Onset Date: 20/01/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Insomnia			

Medicine Details:

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

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

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

LASIX (Other drug) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98616 **Seq:** 1 **Gender:** M
Reported: 31/01/1995 **Weight:** 78.00
Hospitalisation: **Age:** 84Y
Onset Date: 20/01/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

LANOXIN (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
BETALOC (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
AURORIX (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98646 **Seq:** 1 **Gender:** M
Reported: 01/02/1995 **Weight:**
Hospitalisation: **Age:** 64Y
Onset Date: 13/01/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Therapeutic response unexpecte			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 13/01/1995 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has suffered post-traumatic deafness and tinnitus since 1985, since starting aurorix the tinnitus has been marked (somewhat to his delight).

Report Detail

Case Number: 98760 **Seq:** 1 **Gender:** M
Reported: 07/02/1995 **Weight:** 62.00
Hospitalisation: **Age:** 62Y
Onset Date: 24/07/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain			
Agitation			
Anorexia			
Arthralgia			
Arthritis			
Asthenia			
Deafness			
Depression			
Drug ineffective			
Malaise			
Pruritus			

Medicine Details:

METOPROLOL TARTRATE (Other drug) **Reason:** Essential benign hypertension
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 01/01/1977 **Stopped:** CONTIN

CHLORDIAZEPOXIDE HYDROCHLORIDE (Other drug) **Reason:** Depression
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** 01/01/1984 **Stopped:** 28/01/1995

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/07/1994 **Stopped:** 28/01/1995

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98760 **Seq:** 1
Reported: 07/02/1995
Hospitalisation:
Onset Date: 24/07/1994
Outcome: Not yet recovered

Gender: M
Weight: 62.00
Age: 62Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia	Required Specialist Consultation		

Medicine Details:

METOPROLOL TARTRATE (Other drug)			**Reason:** Essential benign hypertension	
Tablet	100.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1977		**Stopped:**	CONTIN
CHLORDIAZEPOXIDE HYDROCHLORIDE (Other drug)			**Reason:** Depression	
Oral application	10.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/01/1984		**Stopped:** 28/01/1995	
MOCLOBEMIDE (Suspected)			**Reason:** Depression	
Tablet	300.0 Milligram	Daily	Oral	
Batch:	**Started:** 22/07/1994		**Stopped:** 28/01/1995	

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

Report Detail

Case Number: 98798 **Seq:** 1 **Gender:** F
Reported: 07/02/1995 **Weight:** 48.00
Hospitalisation: **Age:** 77Y
Onset Date: 16/11/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased		Sexually disinhibited	Dothiepin

Medicine Details:

PROPRANOLOL HYDROCHLORIDE (Other drug) **Reason:** Otr spec symp psychopathol nec
1.0 Dose Unspecified Daily
Batch: **Started:** 11/10/1994 **Stopped:** 13/10/1994

FAMOTIDINE (Other drug) **Reason:** Ulcer of duodenum w/o ment perf
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

DIAZEPAM (Other drug) **Reason:** Otr spec symp psychopathol nec
6.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 28/10/1994 **Stopped:** 17/11/1994

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 98798 **Seq:** 1
Reported: 07/02/1995
Hospitalisation:
Onset Date: 16/11/1994
Outcome: Not yet recovered

Gender: F
Weight: 48.00
Age: 77Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

RIVOTRIL (Suspected) **Reason:** Otr spec symp psychopathol nec
1.0 Gram Daily
Batch: **Started:** L TERM **Stopped:** 25/10/1994

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 98971 **Seq:** 1 **Gender:** F
Reported: 16/02/1995 **Weight:** 49.00
Hospitalisation: **Age:** 67Y
Onset Date: 29/12/1994 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Palpitations			

Medicine Details:

DYAZIDE (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

NILSTAT (Other drug) **Reason:** Debility&undue fatigue
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** S TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 12/12/1994 **Stopped:** 05/02/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on aurorix with same reaction occurring. patient is also allergic to penicillin, codiene and erythromycin - rash.

Report Detail

Case Number: 99349 **Seq:** 1 **Gender:** F
Reported: 07/03/1995 **Weight:** 70.00
Hospitalisation: **Age:** 38Y
Onset Date: 10/02/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus		Pruritus on trunk and limbs	
Rash		Rash on trunk and limbs	
Insomnia			
Nausea			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 10/02/1995 **Stopped:** 22/02/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 99720 **Seq:** 1 **Gender:** F
Reported: 20/03/1995 **Weight:**
Hospitalisation: **Age:** 34Y
Onset Date: 03/03/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dry mouth Somnolence			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
3.0 Dose Unspecified Daily
Batch: **Started:** 03/03/1995 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Changed to night dose

Report Detail

Case Number: 99933 **Seq:** 1 **Gender:** F
Reported: 30/03/1995 **Weight:** 65.00
Hospitalisation: **Age:** 38Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
0.0 Dose Unspecified
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 99935 **Seq:** 1 **Gender:** F
Reported: 30/03/1995 **Weight:** 55.00
Hospitalisation: **Age:** 30Y
Onset Date: 15/03/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vertigo		Motion sickness.	
Acne			
Diarrhoea			
Insomnia			
Nausea	Severe		
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 09/03/1995 **Stopped:** 27/03/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 100278 **Seq:** 1 **Gender:** M
Reported: 18/04/1995 **Weight:** 70.00
Hospitalisation: **Age:** 52Y
Onset Date: 08/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral	Severe	Non-pitting oedema affecting both hands.	

Medicine Details:

BETAMIN (Other drug)		**Reason:**		
	100.0 Milligram	Daily		
Batch:	**Started:** 01/01/1993		**Stopped:**	CONTIN
ROHYPNOL (Other drug)		**Reason:**	Specific disorders of sleep	
	4.0 Milligram	Daily		
Batch:	**Started:** 01/01/1993		**Stopped:**	CONTIN
LARGACTIL (Other drug)		**Reason:**	Otr spec symp psychopathol nec	
	150.0 Milligram	Daily		
Batch:	**Started:** 09/03/1995		**Stopped:**	CONTIN
TRYPTANOL (Other drug)		**Reason:**	Depression	
	250.0 Milligram	Daily		
Batch:	**Started:** 01/01/1992		**Stopped:**	06/03/1995

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100278 **Seq:** 1 **Gender:** M
Reported: 18/04/1995 **Weight:** 70.00
Hospitalisation: **Age:** 52Y
Onset Date: 08/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 20/03/1995 **Stopped:** 10/04/1995

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100642 **Seq:** 1 **Gender:** M
Reported: 05/05/1995 **Weight:** 105.00
Hospitalisation: **Age:** 38Y
Onset Date: 30/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			Advised no pot and no aurorix on 1.5.95

Medicine Details:

CANNABIS PREPARATION (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 10/04/1995 **Stopped:** 01/05/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100874 **Seq:** 1 **Gender:** F
Reported: 19/05/1995 **Weight:** 65.00
Hospitalisation: **Age:** 80Y
Onset Date: 09/05/1995 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thinking abnormal Psychotic disorder		Felt "funny in the head".	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 11/04/1995 **Stopped:** 09/05/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred every time patient took aurorix. see original report for other drugs.

Report Detail

Case Number: 100963 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:** 40.00
Hospitalisation: **Age:** 23Y
Onset Date: 10/04/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Salivary hypersecretion		Excessive salivation	

Medicine Details:

MOGADON (Other drug) **Reason:**
5.0 Milligram As necessary
Batch: **Started:** 01/01/1994 **Stopped:**

VALIUM (Other drug) **Reason:** Anxiety neurosis
5.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 30/03/1995 **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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100990 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:**
Hospitalisation: **Age:** 47Y
Onset Date: 19/04/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Fatigue Headache Hepatic function abnormal Nausea			

Medicine Details:

XANAX (Suspected) **Reason:** Anxiety neurosis
500.0 Microgram As necessary
Batch: **Started:** 28/12/1994 **Stopped:** 18/01/1995

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 10/10/1994 **Stopped:** 18/04/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	(UP TO 45)	19/04/1995	1880	
	ALT = SGPT	(UP TO 45)	26/04/1995	1650	
	ALT = SGPT	(UP TO 45)	24/05/1995	73	
	AST = SGOT	(UP TO 41)	19/04/1995	1110	
	AST = SGOT	(UP TO 41)	24/05/1995	37	
	AST = SGOT	(UP TO 41)	26/04/1995	730	
	Bilirubin	(2-20)	19/04/1995	62	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100990 **Seq:** 1
Reported: 22/05/1995
Hospitalisation:
Onset Date: 19/04/1995
Outcome: Not yet recovered

Gender: F
Weight:
Age: 47Y
DOB:
Causality: Causality possible

Reaction Detai

Medicine Details:

XANAX (Suspected)		**Reason:**	Anxiety neurosis
	500.0 Microgram		As necessary
Batch:	**Started:** 28/12/1994		**Stopped:** 18/01/1995
AURORIX (Suspected)		**Reason:**	Depression
	600.0 Milligram		Daily
Batch:	**Started:** 10/10/1994		**Stopped:** 18/04/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	(2-20)	26/04/1995	27	
	Bilirubin	(2-20)	24/05/1995	11	
	GGT = SGGT =	(UP TO 45)	19/04/1995	229	
	GGT = SGGT =	(UP TO 45)	26/04/1995	193	
	GGT = SGGT =	(UP TO 45)	24/05/1995	41	
	SAP = ALP	(30-115)	19/04/1995	180	
	SAP = ALP	(30-115)	26/04/1995	174	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 100990 **Seq:** 1
Reported: 22/05/1995
Hospitalisation:
Onset Date: 19/04/1995
Outcome: Not yet recovered

Gender: F
Weight:
Age: 47Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

XANAX (Suspected)		**Reason:** Anxiety neurosis	
	500.0 Microgram	As necessary	
Batch:	**Started:** 28/12/1994		**Stopped:** 18/01/1995
AURORIX (Suspected)		**Reason:** Depression	
	600.0 Milligram	Daily	
Batch:	**Started:** 10/10/1994		**Stopped:** 18/04/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP	(30-115)	24/05/1995	74	

Additional Information:

Report Detail

Case Number: 101007 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:** 75.00
Hospitalisation: **Age:** 53Y
Onset Date: 13/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety			
Chest pain			
Confusional state			
Dyspnoea			
Headache			
Nervousness			
Thinking abnormal			

Medicine Details:

ATIVAN (Other drug) **Reason:** Anxiety neurosis
3.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

TRISEQUENS (Other drug) **Reason:** Menopausal symptoms
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1995 **Stopped:**

PREPULSID (Other drug) **Reason:** Other diseases of esophagus
40.0 Milligram Daily
Batch: **Started:** 06/04/1995 **Stopped:**

INDERAL (Other drug) **Reason:** Essential benign hypertension
80.0 Milligram Daily
Batch: **Started:** L TERM **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101007 **Seq:** 1 **Gender:** F
Reported: 22/05/1995 **Weight:** 75.00
Hospitalisation: **Age:** 53Y
Onset Date: 13/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 12/04/1995 **Stopped:** 15/04/1995

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: 101063 **Seq:** 1 **Gender:** F
Reported: 24/05/1995 **Weight:** 95.00
Hospitalisation: **Age:** 68Y
Onset Date: 15/05/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Headache Palpitations	Severe		

Medicine Details:

ALDOMET (Other drug) **Reason:** Essential benign hypertension
500.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 14/05/1995 **Stopped:** 17/05/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 101069 **Seq:** 1 **Gender:** F
Reported: 25/05/1995 **Weight:** 64.00
Hospitalisation: **Age:** 50Y
Onset Date: 09/05/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Nausea Somnolence			

Medicine Details:

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

VOLTAREN (Other drug) **Reason:** Osteoarthritis
0.0 As necessary
Batch: **Started:** **Stopped:**

LOSEC (Other drug) **Reason:** Other diseases of esophagus
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 08/05/1995 **Stopped:** 11/05/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 101149 **Seq:** 1 **Gender:** F
Reported: 26/05/1995 **Weight:** 84.00
Hospitalisation: **Age:** 40Y
Onset Date: 17/05/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alcohol intolerance Dizziness Syncope Tinnitus			In bed for following three days.

Medicine Details:

ALCOHOL (Interaction) **Reason:**
2.0 Dose Unspecified Total
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:** Depression
3.0 Dose Unspecified Daily
Batch: **Started:** 03/05/1995 **Stopped:** 17/05/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101154 **Seq:** 1 **Gender:** F
Reported: 26/05/1995 **Weight:** 50.00
Hospitalisation: **Age:** 78Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety Delirium			

Medicine Details:

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

AROPAX (Suspected) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
Tablet 300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101514 **Seq:** 1 **Gender:** F
Reported: 15/06/1995 **Weight:** 74.00
Hospitalisation: **Age:** 35Y
Onset Date: 08/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 1.0 Dose Unspecified 1 time Oral
Batch: **Started:** 08/06/1995 **Stopped:** 09/06/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101517 **Seq:** 1 **Gender:** M
Reported: 15/06/1995 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 31/01/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 10/03/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	<55	31/01/1995	98	
	Bilirubin		31/01/1995	11	
	GGT = SGGT = SGTP	7-43	31/01/1995	111	
	SAP = ALP	30-110	31/01/1995	176	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101611 **Seq:** 1
Reported: 19/06/1995
Hospitalisation:
Onset Date: 01/06/1995
Outcome: Not yet recovered

Gender: F
Weight: 63.00
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Chest pain Rash maculo-papular	Severe	Itch of palms and ears and eyelids.	Avomine

Medicine Details:

FLUVAX (Other drug) **Reason:** Prophylactic inoculation and vaccination
Injection 1.0 Dose Unspecified 1 time Intramuscular
Batch: **Started:** 04/05/1995 **Stopped:**

LASIX (Other drug) **Reason:** Edema and dropsy
Tablet 80.0 Milligram Weekly Oral
Batch: **Started:** L TERM **Stopped:**

INDERAL (Other drug) **Reason:** Headache
Tablet 80.0 Milligram Weekly Oral
Batch: **Started:** L TERM **Stopped:**

ATENOLOL (Other drug) **Reason:** Essential benign hypertension
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 101611 **Seq:** 1
Reported: 19/06/1995
Hospitalisation:
Onset Date: 01/06/1995
Outcome: Not yet recovered

Gender: F
Weight: 63.00
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ZANTAC (Other drug)			**Reason:**		
Tablet	150.0	Milligram	Daily		Oral
Batch:	**Started:** 23/03/1995			**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	2.0	Dose Unspecified	Daily		Oral
Batch:	**Started:** 23/03/1995			**Stopped:**	03/06/1995
ORUDIS SR (Suspected)			**Reason:**	Unspecified arthritis	
Capsule	100.0	Milligram	Daily		Oral
Batch:	**Started:** 12/01/1995			**Stopped:**	03/06/1995

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 101706 **Seq:** 1 **Gender:** F
Reported: 23/06/1995 **Weight:** 79.00
Hospitalisation: **Age:** 45Y
Onset Date: 12/06/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuralgia		Right facial neuralgic pain and numbness	

Medicine Details:

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** L TERM **Stopped:**

PROVERA (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 12/06/1995 **Stopped:** 17/06/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on moclobemide with same reaction occurring.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101745 **Seq:** 1 **Gender:** M
Reported: 26/06/1995 **Weight:**
Hospitalisation: **Age:** 60
Onset Date: 04/04/1995 **DOB:** 23/01/1935
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Confusional state			Haloperidol

Medicine Details:

TEMAZEPAM (Other drug) **Reason:**
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

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

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 14/03/1995 **Stopped:** 04/04/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101746 **Seq:** 1 **Gender:** M
Reported: 26/06/1995 **Weight:**
Hospitalisation: **Age:** 60
Onset Date: 04/04/1995 **DOB:** 23/01/1935
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine.

Medicine Details:

HALOPERIDOL (Suspected) **Reason:**
Injection 10.0 Milligram 1 time Intramuscular
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 14/03/1995 **Stopped:** 04/04/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101813 **Seq:** 1 **Gender:** F
Reported: 28/06/1995 **Weight:**
Hospitalisation: **Age:** 70Y
Onset Date: 18/06/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise Abdominal pain Headache Hypertension Insomnia Nausea		Feeling awful	

Medicine Details:

VALIUM (Other drug) **Reason:** Otr spec symp psychopathol nec
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

ADALAT (Other drug) **Reason:** Angina pectoris w/o hyperten
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

TAGAMET (Other drug) **Reason:** Otr disorders stomach function
400.0 Milligram Daily
Batch: **Started:** **Stopped:**

PROVERA (Other drug) **Reason:**
5.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101813 **Seq:** 1
Reported: 28/06/1995
Hospitalisation:
Onset Date: 18/06/1995
Outcome: Not yet recovered

Gender: F
Weight:
Age: 70Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 16/06/1995 **Stopped:** 23/06/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101819 **Seq:** 1 **Gender:** M
Reported: 29/06/1995 **Weight:** 57.00
Hospitalisation: **Age:** 72
Onset Date: 30/05/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia Inappropriate antidiuretic hormon		Suspected SIADH	

Medicine Details:

RANITIDINE (Other drug) **Reason:** Peptic ulcer nos w/o ment perf
300.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

NOROXIN (Other drug) **Reason:** Othr diseases of urinary tract
800.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:**
0.0
Batch: **Started:** 05/05/1995 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 101895 **Seq:** 1 **Gender:** U
Reported: 03/07/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension Visual disturbance			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/08/1994 **Stopped:** 28/06/1995

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.

Report Detail

Case Number: 101908 **Seq:** 1 **Gender:** M
Reported: 03/07/1995 **Weight:** 70.00
Hospitalisation: **Age:** 30Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 22/06/1991 **Stopped:** 25/06/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Hypertensive reaction to cheese while on aurorix on two occasions.

Report Detail

Case Number: 101914 **Seq:** 1
Reported: 03/07/1995
Hospitalisation: Admitted to hospital
Onset Date: 23/05/1995
Outcome: Recovered

Gender: F
Weight: 57.00
Age: 74Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia		Had not slept for days.	
Psychotic disorder		Writing notes to the devil.	
Aggression			
Encephalopathy			Serence
Mania			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

300.0 Milligram Daily

Batch: **Started:** 10/05/1995 **Stopped:** 23/05/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 101948 **Seq:** 1 **Gender:** F
Reported: 05/07/1995 **Weight:**
Hospitalisation: **Age:** 70Y
Onset Date: 27/09/1994 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Atrial fibrillation Cerebrovascular disorder Tachycardia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 20/09/1994 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102040 **Seq:** 1 **Gender:** F
Reported: 05/07/1995 **Weight:** 66.00
Hospitalisation: **Age:** 25Y
Onset Date: 24/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Migraine Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 22/06/1995 **Stopped:** 03/07/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102388 **Seq:** 1 **Gender:** F
Reported: 26/07/1995 **Weight:** 68.00
Hospitalisation: **Age:** 39Y
Onset Date: 20/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

3.0 Dose Unspecified Daily

Batch: **Started:** 14/06/1995 **Stopped:** 26/06/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102613 **Seq:** 1 **Gender:** F
Reported: 04/08/1995 **Weight:** 63.00
Hospitalisation: **Age:** 58Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Hearing voices in her head	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/10/1994 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurrs every time aurorix dose is increased from 150mg bd to 300mg bd. see original report for other drugs. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102617 **Seq:** 1 **Gender:** F
Reported: 04/08/1995 **Weight:** 65.00
Hospitalisation: **Age:** 64Y
Onset Date: 26/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Glossitis	Severe	Burning in tongue	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 18/07/1995 **Stopped:** 28/07/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on aurorix with same reaction occurring.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102640 **Seq:** 1 **Gender:** F
Reported: 07/08/1995 **Weight:** 81.00
Hospitalisation: **Age:** 76Y
Onset Date: 29/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Disturbance in attention			
Hypertension			
Pallor			
Pyrexia			
Tachycardia			
Tremor			Valium

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 28/07/1995 **Stopped:** 29/07/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102652 **Seq:** 1 **Gender:** F
Reported: 02/08/1995 **Weight:**
Hospitalisation: **Age:** 27Y
Onset Date: 27/04/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache			

Medicine Details:

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

LOGYNON ED (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 27/04/1995 **Stopped:** 14/07/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102710 **Seq:** 1 **Gender:** F
Reported: 09/08/1995 **Weight:** 60.00
Hospitalisation: **Age:** 65Y
Onset Date: 10/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia			

Medicine Details:

MOGADON (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

ZANTAC (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** **Stopped:**

PROVERA (Other drug) **Reason:**
5.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102710 **Seq:** 1 **Gender:** F
Reported: 09/08/1995 **Weight:** 60.00
Hospitalisation: **Age:** 65Y
Onset Date: 10/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 3.0 Dose Unspecified Total Oral
Batch: **Started:** 10/06/1995 **Stopped:** 11/06/1995

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102788 **Seq:** 1 **Gender:** M
Reported: 11/08/1995 **Weight:**
Hospitalisation: **Age:** 84Y
Onset Date: 01/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression			
Delirium			
Delusion			
Dizziness			
Hypertensive encephalopathy			Haloperidol
Paranoia			
Vomiting			

Medicine Details:

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

LASIX (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

CAPOTEN (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

ISOPTIN (Other drug) **Reason:**
80.0 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: 102788 **Seq:** 1 **Gender:** M
Reported: 11/08/1995 **Weight:**
Hospitalisation: **Age:** 84Y
Onset Date: 01/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

VOLTAREN (Other drug)			**Reason:**	
	100.0 Milligram	Daily		
Batch:	**Started:** 01/05/1995		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Other&unspecified epilepsy
Tablet	2.0 Dose Unspecified	Daily		Oral
Batch:	**Started:** 19/06/1995		**Stopped:**	01/07/1995

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: 102848 **Seq:** 1 **Gender:** F
Reported: 14/08/1995 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 15/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/07/1995 **Stopped:** 28/07/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102867 **Seq:** 1 **Gender:** U
Reported: 14/08/1995 **Weight:**
Hospitalisation: **Age:**
Onset Date: 09/08/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Face oedema Headache Insomnia Nervousness			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 09/08/1995 **Stopped:** 10/08/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 102947 **Seq:** 1 **Gender:** F
Reported: 21/08/1995 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 06/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue			

Medicine Details:

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

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

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

PRESOLOL (Other drug) **Reason:**
2.0 Dose Unspecified 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102947 **Seq:** 1 **Gender:** F
Reported: 21/08/1995 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 06/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

APRESOLINE (Other drug)		**Reason:**	
	2.0 Dose Unspecified Daily		
Batch:	**Started:**	**Stopped:**	
AURORIX (Suspected)		**Reason:**	Depression
	2.0 Dose Unspecified Daily		
Batch:	**Started:** 06/08/1995	**Stopped:**	09/08/1995

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 102950 **Seq:** 1 **Gender:** F
Reported: 21/08/1995 **Weight:** 73.00
Hospitalisation: **Age:** 36Y
Onset Date: 24/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 30/06/1995 **Stopped:** 08/08/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergic to erythromycin

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103180 **Seq:** 1 **Gender:** F
Reported: 30/08/1995 **Weight:**
Hospitalisation: **Age:** 34Y
Onset Date: 29/06/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Galactorrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 01/06/1994 **Stopped:** 14/07/1995

PREPULSID (Suspected) **Reason:** Hematemesis
3.0 Dose Unspecified Daily
Batch: **Started:** 01/06/1995 **Stopped:** 29/06/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103362 **Seq:** 1 **Gender:** F
Reported: 07/09/1995 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 25/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Euphoric mood Hyperkinesia Mania			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/08/1995 **Stopped:** 20/08/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103562 **Seq:** 1 **Gender:** F
Reported: 14/09/1995 **Weight:** 75.00
Hospitalisation: **Age:** 47Y
Onset Date: 02/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypoaesthesia Insomnia Palpitations Paraesthesia			

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 01/08/1995 **Stopped:** 03/08/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103569 **Seq:** 1 **Gender:** F
Reported: 18/09/1995 **Weight:** 116.00
Hospitalisation: **Age:** 43Y
Onset Date: 05/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Thinking abnormal			

Medicine Details:

ZANTAC (Other drug) **Reason:** Diaph hern of abd cav w/o obst
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1992 **Stopped:** CONTIN

COLOFAC (Other drug) **Reason:** Irritable colon
2.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1993 **Stopped:** CONTIN

INDERAL (Other drug) **Reason:** Essential benign hypertension
160.0 Milligram Daily
Batch: **Started:** 01/01/1989 **Stopped:** CONTIN

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
2.0 Milligram Daily
Batch: **Started:** 01/01/1989 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103569 **Seq:** 1 **Gender:** F
Reported: 18/09/1995 **Weight:** 116.00
Hospitalisation: **Age:** 43Y
Onset Date: 05/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 30/08/1995 **Stopped:** 06/09/1995

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.

Report Detail

Case Number: 103596 **Seq:** 1 **Gender:** F
Reported: 18/09/1995 **Weight:** 63.00
Hospitalisation: **Age:** 66Y
Onset Date: 30/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

LASIX (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

SLOW-K (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

PREMARIN (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

SOLPRIN (Other drug) **Reason:**
0.0
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103596 **Seq:** 1 **Gender:** F
Reported: 18/09/1995 **Weight:** 63.00
Hospitalisation: **Age:** 66Y
Onset Date: 30/08/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detai

Medicine Details:

CARDIZEM (Other drug) **Reason:**

0.0

Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 17/08/1995 **Stopped:** 02/09/1995

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 103617 **Seq:** 1 **Gender:** F
Reported: 18/09/1995 **Weight:** 54.00
Hospitalisation: **Age:** 39Y
Onset Date: 08/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Nausea			

Medicine Details:

ERYTHROMYCIN (Suspected) **Reason:** Acute apical periodontitis

Oral application 1.5 Gram Daily Oral

Batch: **Started:** 31/08/1995 **Stopped:** 07/09/1995

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 04/09/1995 **Stopped:** 07/09/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		08/09/1995	318	
	AST = SGOT		08/09/1995	209	
	Albumin		08/09/1995	41	
	Bilirubin		08/09/1995	14	
	GGT = SGGT =		08/09/1995	142	
	Prolactin		08/09/1995	65	
	SAP = ALP		08/09/1995	220	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103697 **Seq:** 1 **Gender:** F
Reported: 25/09/1995 **Weight:**
Hospitalisation: **Age:** 30Y
Onset Date: 12/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dystonia Extrapyramidal disorder Hyperkinesia			

Medicine Details:

RANITIDINE (Other drug) **Reason:** Peptic ulcer nos w/o ment perf
300.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 28/07/1995 **Stopped:** 15/09/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103731 **Seq:** 1
Reported: 27/09/1995
Hospitalisation:
Onset Date: 13/09/1995
Outcome: Not yet recovered

Gender: M
Weight: 115.00
Age: 40Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Severe		
Headache	Severe		
Vertigo			
Visual disturbance			

Medicine Details:

MERSYNDOL (Other drug) **Reason:** Headache
0.0 As necessary
Batch: **Started:** 01/01/1994 **Stopped:**

PANADEINE FORTE (Other drug) **Reason:** Headache
4.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 12/09/1995 **Stopped:** 19/09/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103756 **Seq:** 1
Reported: 29/09/1995
Hospitalisation:
Onset Date: 06/09/1995
Outcome: Not yet recovered

Gender: M
Weight: 68.00
Age: 30Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Euphoric mood Mania			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

300.0 Milligram Daily

Batch: **Started:** 06/09/1995 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient experienced reaction to aurorix after each dose.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 103794 **Seq:** 1 **Gender:** F
Reported: 03/10/1995 **Weight:**
Hospitalisation: **Age:** 78Y
Onset Date: 03/09/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			

Medicine Details:

AURORIX (Suspected) **Reason:** Pain

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 31/08/1995 **Stopped:** 04/09/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104089 **Seq:** 1 **Gender:** F
Reported: 17/10/1995 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: 04/10/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			Calamine lotion

Medicine Details:

SIMVASTATIN (Other drug) **Reason:** Othr&unspec metabolic diseases
10.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 30/09/1995 **Stopped:** 07/10/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104098 **Seq:** 1 **Gender:** F
Reported: 18/10/1995 **Weight:** 62.00
Hospitalisation: **Age:** 41Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104116 **Seq:** 1 **Gender:** F
Reported: 19/10/1995 **Weight:**
Hospitalisation: **Age:** 79Y
Onset Date: 03/10/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Pruritus		Rash on trunk and limbs.	

Medicine Details:

PLENDIL ER (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

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

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

BECOTIDE (Other drug) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104116 **Seq:** 1
Reported: 19/10/1995
Hospitalisation:
Onset Date: 03/10/1995
Outcome: Not yet recovered

Gender: F
Weight:
Age: 79Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 3.0 Dose Unspecified Daily Oral

Batch: **Started:** 26/09/1995 **Stopped:** 10/10/1995

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 104209 **Seq:** 1 **Gender:** F
Reported: 25/10/1995 **Weight:** 52.00
Hospitalisation: **Age:** 69Y
Onset Date: 03/09/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression Nausea Palpitations Thinking abnormal			

Medicine Details:

VENTOLIN (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** L TERM **Stopped:**

OROXINE (Other drug) **Reason:**
200.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

CARTIA (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1992 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 26/08/1995 **Stopped:** 01/10/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104400 **Seq:** 1 **Gender:** F
Reported: 02/11/1995 **Weight:** 84.00
Hospitalisation: **Age:** 59Y
Onset Date: 26/10/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Rash inner thighs upper	

Medicine Details:

LOSEC (Other drug) **Reason:** Other diseases of esophagus
0.0
Batch: **Started:** 01/01/1992 **Stopped:**

MAXOLON (Other drug) **Reason:** Nausea and vomiting
0.0
Batch: **Started:** 01/01/1992 **Stopped:**

THYROXINE SODIUM (Other drug) **Reason:** Myxedema
Tablet 100.0 Microgram Daily Oral
Batch: **Started:** 01/01/1980 **Stopped:**

RIVOTRIL (Other drug) **Reason:**
Oral application 1.0 Milligram Daily Oral
Batch: **Started:** 01/01/1990 **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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104400 **Seq:** 1
Reported: 02/11/1995
Hospitalisation:
Onset Date: 26/10/1995
Outcome: Not yet recovered

Gender: F
Weight: 84.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AMINOGLUTETHIMIDE (Other drug)			**Reason:**	Malignant neoplasm of breast
Tablet	250.0 Milligram	Daily		Oral
Batch:	**Started:** 01/01/1992		**Stopped:**	
ZANTAC (Other drug)			**Reason:**	Other diseases of esophagus
Tablet	300.0 Milligram	Daily		Oral
Batch:	**Started:** 01/01/1992		**Stopped:**	
AURORIX (Suspected)			**Reason:**	
Tablet	450.0 Milligram	Daily		Oral
Batch:	**Started:** 11/10/1995		**Stopped:**	27/10/1995

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104454 **Seq:** 1 **Gender:** F
Reported: 07/11/1995 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: 30/10/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperprolactinaemia		Prolactin increased.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/01/1995 **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: 104462 **Seq:** 1 **Gender:** F
Reported: 07/11/1995 **Weight:** 80.00
Hospitalisation: **Age:** A3
Onset Date: 01/11/1995 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			
Dry mouth			
Headache			
Hypoaesthesia			
Insomnia			
Paralysis			
Thinking abnormal			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 01/11/1995 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 104523 **Seq:** 1 **Gender:** M
Reported: 09/11/1995 **Weight:** 84.00
Hospitalisation: **Age:** 28Y
Onset Date: 12/10/1995 **DOB:**
Outcome: Recovered without treatment **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence Thinking abnormal			

Medicine Details:

ERGODRYL (Suspected) **Reason:**
1.0 Dose Unspecified 1 time
Batch: **Started:** 12/10/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 12/10/1995 **Stopped:** 12/10/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 104541 **Seq:** 1
Reported: 10/11/1995
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered with sequelae

Gender: F
Weight: 62.00
Age: 45Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation Headache Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

Batch: **Started:** 25/10/1995 **Stopped:** 30/10/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104544 **Seq:** 1 **Gender:** F
Reported: 10/11/1995 **Weight:** 77.00
Hospitalisation: **Age:** 63Y
Onset Date: 01/11/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 27/10/1995 **Stopped:** 02/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 104615 **Seq:** 1 **Gender:** F
Reported: 15/11/1995 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia Fatigue			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 04/05/1995 **Stopped:** 20/10/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 104639 **Seq:** 1 **Gender:** F
Reported: 15/11/1995 **Weight:** 68.00
Hospitalisation: **Age:** 45Y
Onset Date: 19/10/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Nausea Vomiting			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 19/10/1995 **Stopped:** 02/11/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104701 **Seq:** 1 **Gender:** F
Reported: 20/11/1995 **Weight:**
Hospitalisation: **Age:** A7
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Agitation Hyperkinesia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/01/1993 **Stopped:**

ZOLOFT (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 1D

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104723 **Seq:** 1 **Gender:** F
Reported: 21/11/1995 **Weight:** 52.00
Hospitalisation: **Age:** 90Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuropathy			

Medicine Details:

DAPSONE (Suspected) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 01/01/1993 **Stopped:** 08/08/1995

CODEINE (Suspected) **Reason:** Cough
Oral Liquid 15.0 Millilitre Daily Oral
Batch: **Started:** **Stopped:**

POLARAMINE (Suspected) **Reason:** Chronic sinusitis
4.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Falls and unilateral foot drop. see original report for other drugs. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104723 **Seq:** 1
Reported: 21/11/1995
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 52.00
Age: 90Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

CAPOTEN (Suspected) **Reason:** Essential benign hypertension
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

Falls and unilateral foot drop. see original report for other drugs. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104790 **Seq:** 1
Reported: 21/11/1995
Hospitalisation:
Onset Date: 02/10/1995
Outcome: Death, maybe drug

Gender: F
Weight: 38.00
Age: 32Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sudden death			

Medicine Details:

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

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

MOCLOBEMIDE (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 104964 **Seq:** 1
Reported: 29/11/1995
Hospitalisation: Admitted to hospital
Onset Date: 05/08/1995
Outcome: Recovered

Gender: F
Weight:
Age: 52Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia			
Dysarthria			
Dyskinesia			
Dyspnoea			
Headache			
Hypertension			
Pyrexia			

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
2.5 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

CELERIAC (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
450.0 Milligram Daily
Batch: **Started:** 01/06/1995 **Stopped:**

ROBITUSSIN DM (Suspected) **Reason:**
10.0 Millilitre 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: 105005 **Seq:** 1 **Gender:** F
Reported: 30/11/1995 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 24/11/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Confusional state Drug withdrawal syndrome			

Medicine Details:

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
2.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

NATRILIX (Other drug) **Reason:** Essential benign hypertension
2.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:**
4.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1993 **Stopped:** 21/10/1995

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: 105010 **Seq:** 1 **Gender:** F
Reported: 30/11/1995 **Weight:** 45.00
Hospitalisation: **Age:** 17Y
Onset Date: 20/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haematemesis			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 06/11/1995 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105036 **Seq:** 1 **Gender:** F
Reported: 04/12/1995 **Weight:** 62.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia Convulsion		Left fingers With absenses	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 17/10/1995 **Stopped:** 06/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 105076 **Seq:** 1 **Gender:** M
Reported: 04/12/1995 **Weight:**
Hospitalisation: **Age:** 67Y
Onset Date: 31/10/1995 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Diarrhoea Dizziness Visual disturbance			

Medicine Details:

VERAPAMIL HYDROCHLORIDE (Other drug) **Reason:**
400.0 Milligram Daily
Batch: **Started:** **Stopped:**

RANITIDINE (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged with moclobemide and same thing happened. - 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105113 **Seq:** 1 **Gender:** M
Reported: 05/12/1995 **Weight:** 94.00
Hospitalisation: **Age:** 52Y
Onset Date: 02/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Influenza like illness Arthralgia Myalgia		Flu-like state	

Medicine Details:

COVERSYL (Other drug) **Reason:** Essential benign hypertension

Tablet 4.0 Milligram Daily Oral

Batch: **Started:** 01/10/1993 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 05/10/1995 **Stopped:** 29/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Medicine ceased and restarted on three occasions with same result on each. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 105149 **Seq:** 1 **Gender:** F
Reported: 07/12/1995 **Weight:**
Hospitalisation: **Age:** 67Y
Onset Date: 20/10/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Albuminuria			
Confusional state			
Headache			
Hepatic function abnormal			
Hyperreflexia			
Hypertension			
Hypertensive encephalopathy			Nifedipine, enalapril, atenolol
Hypokalaemia			
Insomnia			
Pyrexia			

Medicine Details:

CLONAZEPAM (Other drug) **Reason:** Otr spec symp psychopathol nec

Oral application 0.5 Milligram Daily Oral

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

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 18/10/1995 **Stopped:** 20/10/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		21/10/1995	197	
	Bilirubin		21/10/1995	20	
	Computerised [illegible]				Ct scan showed only aged related atrophy.
	GGT = SGGT = [illegible]		21/10/1995	114	All abnormalities resolved within seventy two hours.
	LDH		21/10/1995	481	
	Other data				Lumbar punctures - xanthochromia. 25/11/1995 bp was 200/100 in the right arm and about 170/90 in the left arm.

Additional Information:

Patient unable to dress herself, one small flame haemorrhage in each eye, arteriolar spasm. patient recovered within 10 days.

Report Detail

Case Number: 105208 **Seq:** 1 **Gender:** F
Reported: 08/12/1995 **Weight:** 57.00
Hospitalisation: **Age:** 48Y
Onset Date: 09/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria	Severe		Antihistamine

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 08/11/1995 **Stopped:** 09/11/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105265 **Seq:** 1 **Gender:** F
Reported: 11/12/1995 **Weight:** 52.00
Hospitalisation: **Age:** 32Y
Onset Date: 05/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			Phenergan

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 1.0 Dose Unspecified 1 time Oral
Batch: **Started:** 05/12/1995 **Stopped:** 05/12/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 105444 **Seq:** 1 **Gender:** M
Reported: 18/12/1995 **Weight:** 83.00
Hospitalisation: **Age:** 84Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness			

Medicine Details:

MODURETIC (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 26/10/1995 **Stopped:** 03/11/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105457 **Seq:** 1 **Gender:** F
Reported: 19/12/1995 **Weight:**
Hospitalisation: **Age:** 64Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Confusional state Coordination abnormal Diarrhoea Malaise			

Medicine Details:

IMIGRAN (Interaction) **Reason:** Migraine

Tablet	100.0	Milligram	1 time	Oral

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

AURORIX (Interaction) **Reason:** Depression

Tablet	200.0	Milligram	Daily	Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Chronic fatigue syndrome from 1985..

Report Detail

Case Number: 105586 **Seq:** 1 **Gender:** F
Reported: 02/01/1996 **Weight:** 63.00
Hospitalisation: **Age:** 36Y
Onset Date: 15/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Agitation Diarrhoea Dry mouth Dysphagia Hypoaesthesia Nausea		Extreme panic	

Medicine Details:

ALCOHOL (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 09/12/1995 **Stopped:**

TRIQUILAR (Other drug) **Reason:** Contraception
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1989 **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: 105616 **Seq:** 1 **Gender:** F
Reported: 02/01/1996 **Weight:**
Hospitalisation: **Age:** 82Y
Onset Date: 10/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Orthostatic hypotension			

Medicine Details:

CARDIPRIN 100 (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:**

QUINATE (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:**

ROHYPNOL (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/11/1995 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105630 **Seq:** 1 **Gender:** F
Reported: 02/01/1996 **Weight:** 65.00
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Dysgeusia Pruritus		Off food +++	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 05/09/1995 **Stopped:** 28/10/1995

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: 105641 **Seq:** 1 **Gender:** F
Reported: 02/01/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: 07/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura			

Medicine Details:

PREMARIN (Other drug) **Reason:** Menopausal symptoms
2.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

DUPHASTON (Other drug) **Reason:** Menopausal symptoms
10.0 Milligram Cyclical
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 04/12/1995 **Stopped:** 11/12/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 105691 **Seq:** 1 **Gender:** F
Reported: 04/01/1996 **Weight:**
Hospitalisation: **Age:** 75
Onset Date: 01/12/1995 **DOB:** 23/03/1920
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular			

Medicine Details:

RAMIPRIL (Other drug) **Reason:** Congestive heart failure
Capsule 2.5 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

FRUSEMIDE (Other drug) **Reason:** Congestive heart failure
Oral application 40.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 30/11/1995 **Stopped:** 01/12/1995

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 105733 **Seq:** 1
Reported: 05/01/1996
Hospitalisation:
Onset Date: 19/12/1995
Outcome: Not yet recovered

Gender: M
Weight: 70.00
Age: 36Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Folliculitis Pruritus Vasculitis			

Medicine Details:

DI-GESIC (Suspected) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 27/10/1995 **Stopped:** 24/12/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Skin biopsy - vasculitis

Additional Information:

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: 105951 **Seq:** 1 **Gender:** M
Reported: 18/01/1996 **Weight:** 89.00
Hospitalisation: **Age:** 38Y
Onset Date: 09/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Flushing Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 08/01/1996 **Stopped:** 11/01/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106042 **Seq:** 1
Reported: 24/01/1996
Hospitalisation:
Onset Date: 06/01/1996
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 22Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia Abdominal pain Conjunctivitis Diarrhoea Headache Hepatic function abnormal Leukopenia Pruritus Rash erythematous Rash maculo-papular Thrombocytopenia		Fever to 40 degrees	Iv fluids

Medicine Details:

CEPHALEXIN (Other drug) **Reason:** Pyrexia of unknown origin

1.0 Dose Unspecified 1 time

Batch: **Started:** 05/01/1996 **Stopped:** 05/01/1996

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 27/12/1995 **Stopped:** 10/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				The hepatic enzymes rose during admission, with ast and alt peaking at three times the normal range.
	Platelets		10/01/1996	137	
	White blood cells		10/01/1996	3.1	

Additional Information:

Symptoms settled with cessation of the drug and developed again on rechallenge with moclobemide.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106042 **Seq:** 1 **Gender:** F
Reported: 24/01/1996 **Weight:** 53.00
Hospitalisation: **Age:** 22Y
Onset Date: 06/01/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting			

Medicine Details:

CEPHALEXIN (Other drug) **Reason:** Pyrexia of unknown origin
1.0 Dose Unspecified 1 time
Batch: **Started:** 05/01/1996 **Stopped:** 05/01/1996

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 27/12/1995 **Stopped:** 10/01/1996

Laboratory Investigations:

Additional Information:

Symptoms settled with cessation of the drug and developed again on rechallenge with moclobemide.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106185 **Seq:** 1 **Gender:** F
Reported: 01/02/1996 **Weight:** 57.00
Hospitalisation: **Age:** 56Y
Onset Date: 27/11/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Paraesthesia			 Teldene

Medicine Details:

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

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

ZANTAC (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Weekly Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
0.0
Batch: **Started:** 15/11/1995 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106187 **Seq:** 1 **Gender:** F
Reported: 01/02/1996 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: 26/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia Neurosis	Severe		

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 20/01/1996 **Stopped:** 30/01/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106232 **Seq:** 1 **Gender:** M
Reported: 02/02/1996 **Weight:**
Hospitalisation: **Age:** 43
Onset Date: 27/01/1996 **DOB:** 25/04/1952
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 09/12/1995 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106286 **Seq:** 1 **Gender:** F
Reported: 07/02/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
0.0
Batch: **Started:** **Stopped:**

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 106289 **Seq:** 1 **Gender:** F
Reported: 05/02/1996 **Weight:** 50.00
Hospitalisation: **Age:** 79Y
Onset Date: 22/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise Rash erythematous Urticaria			

Medicine Details:

SERC (Other drug) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** 01/01/1995 **Stopped:**

MODURETIC (Other drug) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** 01/01/1995 **Stopped:**

PLENDIL (Other drug) **Reason:** Essential benign hypertension
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 23/12/1995 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106289 **Seq:** 1 **Gender:** F
Reported: 05/02/1996 **Weight:** 50.00
Hospitalisation: **Age:** 79Y
Onset Date: 22/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
3.0 Dose Unspecified Daily
Batch: **Started:** 14/01/1996 **Stopped:** 23/01/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106340 **Seq:** 1 **Gender:** M
Reported: 07/02/1996 **Weight:** 75.00
Hospitalisation: **Age:** 38Y
Onset Date: 20/12/1995 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Choking Dyspnoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 15/11/1995 **Stopped:** 25/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Trial of re-introduction after 7 days without aurorix (symptoms nearly completely subsided) caused reaction to occur only worse in intensity.

Report Detail

Case Number: 106374 **Seq:** 1 **Gender:** M
Reported: 09/02/1996 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: 18/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dermatitis bullous Urticaria			Antihistamines

Medicine Details:

CAPOTEN (Other drug) **Reason:** Essential benign hypertension

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression

150.0 Milligram 1 time

Batch: **Started:** 18/01/1996 **Stopped:** 18/01/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106403 **Seq:** 1 **Gender:** F
Reported: 12/02/1996 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Anxiety			
Confusional state			
Constipation			
Diarrhoea			
Diplopia			
Disturbance in attention			
Dizziness			
Headache			
Insomnia			
Nausea			

Medicine Details:

THYROXINE SODIUM (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 29/12/1995 **Stopped:** 01/02/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106403 **Seq:** 1 **Gender:** F
Reported: 12/02/1996 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thirst			

Medicine Details:

THYROXINE SODIUM (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 29/12/1995 **Stopped:** 01/02/1996

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 106415 **Seq:** 1 **Gender:** M
Reported: 12/02/1996 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 06/02/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypotension			

Medicine Details:

ECOTRIN (Other drug) **Reason:**
Tablet 1.3 Gram Weekly Oral
Batch: **Started:** L TERM **Stopped:**

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

SIMVASTATIN (Other drug) **Reason:** Othr&unspec metabolic diseases
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 31/07/1995 **Stopped:**

TAGAMET (Other drug) **Reason:** Other diseases of esophagus
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 25/01/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106415 **Seq:** 1 **Gender:** M
Reported: 12/02/1996 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 06/02/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

PULMICORT (Other drug)			**Reason:**	Asthma	
Inhalation	400.0	Microgram	Daily		Inhalation
Batch:	**Started:**	L TERM		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**	30/01/1996		**Stopped:**	06/02/1996

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106854 **Seq:** 1 **Gender:** F
Reported: 04/03/1996 **Weight:** 63.00
Hospitalisation: **Age:** 33Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysuria Micturition disorder			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 13/02/1996 **Stopped:** 15/02/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106907 **Seq:** 1 **Gender:** M
Reported: 06/03/1996 **Weight:** 70.00
Hospitalisation: **Age:** 44Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:** 27/02/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 106910 **Seq:** 1
Reported: 06/03/1996
Hospitalisation:
Onset Date: 29/02/1996
Outcome: Not yet recovered

Gender: F
Weight: 82.00
Age: 59Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Face oedema Pharyngitis			

Medicine Details:

DUCENE (Other drug) **Reason:** Anxiety neurosis
Tablet 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** L TERM **Stopped:**

MODURETIC (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

SLOW-K (Other drug) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** 28/02/1996 **Stopped:**

AURORIX (Suspected) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** 28/02/1996 **Stopped:** 01/03/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has allergy to aspirin-itchy rash, clinoril-rash itchy.

Report Detail

Case Number: 106934 **Seq:** 1 **Gender:** M
Reported: 07/03/1996 **Weight:** 78.00
Hospitalisation: **Age:** 67Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Vomiting			

Medicine Details:

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

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 02/12/1995 **Stopped:** 08/02/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 106943 **Seq:** 1 **Gender:** F
Reported: 08/03/1996 **Weight:** 60.00
Hospitalisation: **Age:** 51Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thirst			

Medicine Details:

EPILIM (Other drug)		**Reason:**	Otr spec symp psychopathol nec	
Tablet	100.0 Milligram	Daily	Oral	
Batch:	**Started:** 24/01/1996		**Stopped:**	CONTIN
NAVANE (Other drug)		**Reason:**	Unspecified psychosis	
Tablet	2.0 Milligram	Daily	Oral	
Batch:	**Started:** 09/02/1995		**Stopped:**	CONTIN
ARTANE (Other drug)		**Reason:**		
Tablet	2.0 Milligram	Daily	Oral	
Batch:	**Started:** 15/02/1996		**Stopped:**	CONTIN
AURORIX (Suspected)		**Reason:**	Depression	
Tablet	150.0 Milligram	Daily	Oral	
Batch:	**Started:** 22/05/1995		**Stopped:**	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Rechallenge: reaction clears after about 2 days when medication discontinued. reaction recurs 2 days after resuming medication.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107007 **Seq:** 1 **Gender:** F
Reported: 12/03/1996 **Weight:** 50.00
Hospitalisation: **Age:** 46Y
Onset Date: 27/02/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia		Perioral paraesthesia	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 20/02/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Rechallenge: reaction begins 1-1.5 hours after medication, lasts for 2 hours.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107126 **Seq:** 1 **Gender:** F
Reported: 20/03/1996 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: 12/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Face oedema Urticaria			S/c adrenaline; im

Medicine Details:

KAPANOL (Other drug) **Reason:** Pain
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** 06/02/1996 **Stopped:**

FRUSEMIDE (Other drug) **Reason:** Noninfec dis of lymph channels
Oral application 500.0 Milligram Daily Oral
Batch: **Started:** 21/02/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 11/03/1996 **Stopped:** 11/03/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107144 **Seq:** 1 **Gender:** F
Reported: 20/03/1996 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 14/01/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nausea Vertigo	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
1.0 Dose Unspecified Daily
Batch: **Started:** 10/01/1996 **Stopped:** 14/01/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107195 **Seq:** 1 **Gender:** F
Reported: 21/03/1996 **Weight:** 55.00
Hospitalisation: **Age:** 29Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Drug withdrawal syndrome	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Severe anxiety symptoms when dose reduced from 450mg daily to 100mg. reduction attempted twice and same reaction occurred. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107202 **Seq:** 1 **Gender:** F
Reported: 21/03/1996 **Weight:** 54.00
Hospitalisation: **Age:** 26Y
Onset Date: 29/02/1996 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vaginal infection		Vaginal odour.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 26/02/1996 **Stopped:** 29/02/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Same reaction occurred on rechallenge with aurorix.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107211 **Seq:** 1 **Gender:** F
Reported: 21/03/1996 **Weight:** 75.00
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nocturia Pollakiuria			

Medicine Details:

VOLTAREN (Other drug) **Reason:** Unspecified arthritis
100.0 Milligram As necessary
Batch: **Started:** 01/12/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 24/11/1995 **Stopped:** 03/03/1996

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107435 **Seq:** 1 **Gender:** F
Reported: 29/03/1996 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 18/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Dry mouth Flushing Pyrexia			

Medicine Details:

DI-GESIC (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
Tablet 0.0 As necessary Oral
Batch: **Started:** **Stopped:**

TEMAZE (Other drug) **Reason:** Specific disorders of sleep
Capsule 1.0 Dose Unspecified As necessary Oral
Batch: **Started:** **Stopped:**

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
Tablet 6.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

BURINEX (Other drug) **Reason:** Essential benign hypertension
Tablet 1.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107435 **Seq:** 1 **Gender:** F
Reported: 29/03/1996 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 18/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 14/03/1996 **Stopped:** 18/03/1996

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 107692 **Seq:** 1 **Gender:** F
Reported: 03/04/1996 **Weight:**
Hospitalisation: **Age:** 78Y
Onset Date: 26/03/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Melaena			

Medicine Details:

MACRODANTIN (Suspected) **Reason:** Othr diseases of urinary tract
50.0 Milligram Daily
Batch: **Started:** 01/12/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 01/07/1995 **Stopped:**

COGNEX (Suspected) **Reason:** Presenile dementia
160.0 Milligram Daily
Batch: **Started:** 01/10/1994 **Stopped:** 26/03/1996

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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107759 **Seq:** 1 **Gender:** F
Reported: 09/04/1996 **Weight:** 70.00
Hospitalisation: **Age:** 62Y
Onset Date: 20/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 20/03/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Same reaction occurred on rechallenge with aurorix three days later.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107772 **Seq:** 1 **Gender:** F
Reported: 10/04/1996 **Weight:** 70.00
Hospitalisation: **Age:** 56Y
Onset Date: 28/03/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 08/03/1996 **Stopped:** 28/03/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	< 56	28/03/1996	202	
	AST = SGOT	< 41	28/03/1996	123	
	Bilirubin	< 21	28/03/1996	35	
	GGT = SGGT =	< 51	28/03/1996	257	

Additional Information:

Lft's were normal on 9/2/96

Report Detail

Case Number: 107804 **Seq:** 1 **Gender:** F
Reported: 10/04/1996 **Weight:**
Hospitalisation: **Age:** 24Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Agitation Convulsion Convulsions local Flushing Serotonin syndrome Tremor Urinary retention Vomiting		Burning feeling in chest.	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 10W

FLUNITRAZEPAM (Suspected) **Reason:**
2.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Serum level (steady state) on nortriptyline 100mg nocte was 73mu g/l, giving a calculated clearance rate of 0.73; those with 2d6 deficiency exhibit values of 1.5 or more.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107943 **Seq:** 1 **Gender:** F
Reported: 17/04/1996 **Weight:** 55.00
Hospitalisation: **Age:** 30Y
Onset Date: 09/02/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope		Collapse	
Chills		Shivering attacks	
Flushing			

Medicine Details:

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

AURORIX (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 107996 **Seq:** 1 **Gender:** F
Reported: 22/04/1996 **Weight:** 50.00
Hospitalisation: **Age:** 50Y
Onset Date: 24/02/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Pruritus Rash erythematous Rash maculo-papular			

Medicine Details:

XANAX (Other drug) **Reason:** Anxiety neurosis
Tablet 1.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 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.

Report Detail

Case Number: 108074 **Seq:** 1 **Gender:** F
Reported: 26/04/1996 **Weight:** 82.00
Hospitalisation: **Age:**
Onset Date: 02/04/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Urticaria	Severe	Severe, painless, urgent diarrhoea	

Medicine Details:

ESTRADERM (Other drug) **Reason:** Menopausal symptoms

Cream 100.0 Microgram Daily Transdermal

Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:**

450.0 Milligram Daily

Batch: **Started:** 25/03/1995 **Stopped:** 19/04/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 108132 **Seq:** 1 **Gender:** F
Reported: 30/04/1996 **Weight:**
Hospitalisation: **Age:** 67Y
Onset Date: 16/04/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis cholestatic			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/03/1996 **Stopped:** 18/04/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 108365 **Seq:** 1 **Gender:** F
Reported: 08/05/1996 **Weight:** 66.00
Hospitalisation: **Age:** 71Y
Onset Date: 10/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema Periorbital oedema			

Medicine Details:

HALOPERIDOL (Other drug) **Reason:** Otr spec symp psychopathol nec
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

VERAPAMIL HYDROCHLORIDE (Other drug) **Reason:** Otr&nos disord of heart rhythm
Oral application 240.0 Milligram Daily Oral
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

DIGOXIN (Other drug) **Reason:** Otr&nos disord of heart rhythm
375.0 Microgram Daily
Batch: **Started:** 01/01/1994 **Stopped:** CONTIN

DOTHIEPIN HYDROCHLORIDE (Other drug) **Reason:** Depression
100.0 Milligram Daily
Batch: **Started:** 01/01/1982 **Stopped:** 07/03/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 108365 **Seq:** 1 **Gender:** F
Reported: 08/05/1996 **Weight:** 66.00
Hospitalisation: **Age:** 71Y
Onset Date: 10/03/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 09/03/1996 **Stopped:** 14/03/1996

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.

Report Detail

Case Number: 108639 **Seq:** 1 **Gender:** M
Reported: 23/05/1996 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photophobia			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
3.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction ceased with withdrawal and recurred with reintroduction.

Report Detail

Case Number: 108949 **Seq:** 1 **Gender:** F
Reported: 03/06/1996 **Weight:** 70.00
Hospitalisation: **Age:** 32Y
Onset Date: 14/04/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Acne		Large cysts on jaw and back	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 09/04/1996 **Stopped:** 09/05/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109013 **Seq:** 1 **Gender:** F
Reported: 06/06/1996 **Weight:** 70.00
Hospitalisation: **Age:** 42Y
Onset Date: 27/05/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria Pruritus		Rash on trunk and limbs.	

Medicine Details:

PROVELLE-14 (Other drug) **Reason:**
0.0
Batch: **Started:** 01/03/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 14/05/1996 **Stopped:** 27/05/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of allergy to penicillin and tetracycline. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 109135 **Seq:** 1
Reported: 14/06/1996
Hospitalisation: Admitted to hospital
Onset Date: 06/06/1996
Outcome: Recovered

Gender: F
Weight: 45.00
Age: 28Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperreflexia		Increased reflexes, increased jaw reflexes	
Dystonia		Stiffness in legs	
Amnesia			
Anxiety			
Hypertonia			
Serotonin syndrome			
Tachycardia			
Tremor			Cogentin 2mg imi, valium 5mg

Medicine Details:

MELLERIL (Other drug) **Reason:**
Oral application 75.0 Milligram Daily Oral
Batch: **Started:** 01/05/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 05/06/1996 **Stopped:** 07/06/1996

AROPAX (Suspected) **Reason:**
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 01/04/1996 **Stopped:** 05/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Stiffness (?dystonic) came on > 3hrs after 1st dose of aurorix and the same reaction occurred after 2nd dose. 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: 109205 **Seq:** 1 **Gender:** F
Reported: 18/06/1996 **Weight:** 58.00
Hospitalisation: **Age:** 52Y
Onset Date: 28/05/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Depression Drug ineffective		Parodoxical reaction	

Medicine Details:

CAPOTEN (Other drug) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

CHLOTRIDE (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

OGEN (Other drug) **Reason:** Menopausal symptoms
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

PROVERA (Other drug) **Reason:** Menopausal symptoms
1.0 Dose Unspecified Daily
Batch: **Started:** L TERM **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109205 **Seq:** 1 **Gender:** F
Reported: 18/06/1996 **Weight:** 58.00
Hospitalisation: **Age:** 52Y
Onset Date: 28/05/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 28/05/1996 **Stopped:** 11/06/1996

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109441 **Seq:** 1 **Gender:** F
Reported: 24/06/1996 **Weight:** 84.00
Hospitalisation: **Age:** 47Y
Onset Date: 13/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous Rash maculo-papular			

Medicine Details:

TESTOSTERONE (Other drug) **Reason:**
0.0
Batch: **Started:** 01/02/1996 **Stopped:**

EVENING PRIMROSE OIL (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ESTRADERM (Other drug) **Reason:** Menopausal symptoms
2.0 Dose Unspecified Weekly
Batch: **Started:** 01/01/1995 **Stopped:**

PROVERA (Other drug) **Reason:** Menopausal symptoms
Tablet 2.5 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **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 necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 109441 **Seq:** 1 **Gender:** F
Reported: 24/06/1996 **Weight:** 84.00
Hospitalisation: **Age:** 47Y
Onset Date: 13/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

SUSTANON 100 (Other drug)			**Reason:** Menopausal symptoms	
	100.0 Milligram	Daily		
Batch:	**Started:** 23/05/1996		**Stopped:**	
AURORIX (Suspected)			**Reason:** Depression	
Tablet	600.0 Milligram	Daily		Oral
Batch:	**Started:** 03/06/1996		**Stopped:** 14/06/1996	

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 necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 109542 **Seq:** 1 **Gender:** F
Reported: 26/06/1996 **Weight:** 44.00
Hospitalisation: **Age:** 39Y
Onset Date: 01/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash erythematous	Required Specialist Consultation		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 17/05/1996 **Stopped:** 04/06/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109552 **Seq:** 1 **Gender:** F
Reported: 28/06/1996 **Weight:** 61.00
Hospitalisation: **Age:** 22Y
Onset Date: 21/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 11/06/1996 **Stopped:** 20/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of allergy to aspirin.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109747 **Seq:** 1 **Gender:** M
Reported: 05/07/1996 **Weight:**
Hospitalisation: **Age:** 56Y
Onset Date: 21/06/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Vasculitis	Severe		

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

PLENDIL ER (Other drug) **Reason:** Essential benign hypertension
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 20/06/1996 **Stopped:** 24/06/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109782 **Seq:** 1 **Gender:** F
Reported: 08/07/1996 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Lymphocytic urticaria	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Oral application 0.0 Oral
Batch: **Started:** 01/11/1995 **Stopped:** 28/05/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Histology				Histology showing a perivascular lymphocytic infiltrate

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109849 **Seq:** 1 **Gender:** M
Reported: 09/07/1996 **Weight:** 60.00
Hospitalisation: **Age:** 58Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 18/06/1996 **Stopped:** 21/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction became disabling when dose increased and quickly settled on stopping medication.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109878 **Seq:** 1 **Gender:** F
Reported: 10/07/1996 **Weight:** 78.00
Hospitalisation: **Age:** 50Y
Onset Date: 24/06/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Rash around upper abdomen, wrists,	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 16/06/1996 **Stopped:** 27/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109887 **Seq:** 1 **Gender:** F
Reported: 11/07/1996 **Weight:** 67.00
Hospitalisation: **Age:** 70Y
Onset Date: 13/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Euphoric mood		'on a high'	
Agitation		On edge/couldnt sit still	
Insomnia		Unable to sleep	

Medicine Details:

MINIPRESS (Other drug) **Reason:** Essential benign hypertension
1.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

SEREPAX (Other drug) **Reason:** Anxiety neurosis
15.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Sudden death (cause unknown)
300.0 Milligram Daily
Batch: **Started:** 13/06/1996 **Stopped:** 14/06/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109931 **Seq:** 1 **Gender:** F
Reported: 11/07/1996 **Weight:** 75.00
Hospitalisation: **Age:** 36Y
Onset Date: 05/07/1996 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Agitation Depression			Serepax 15mg to calm down

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Aurorix tried twice before, resulting in increased anxiety.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109939 **Seq:** 1 **Gender:** F
Reported: 15/07/1996 **Weight:** 77.00
Hospitalisation: **Age:** 75Y
Onset Date: 20/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Faecal incontinence Urinary incontinence			

Medicine Details:

RENITEC (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

PREMARIN (Other drug) **Reason:**
300.0 Microgram Daily
Batch: **Started:** **Stopped:**

PROVERA (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 07/06/1996 **Stopped:** 30/06/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109945 **Seq:** 1 **Gender:** F
Reported: 15/07/1996 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: 28/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia		Burning feeling in feet	

Medicine Details:

OROXINE (Other drug) **Reason:**
Tablet 100.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 28/06/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 109975 **Seq:** 1 **Gender:** F
Reported: 16/07/1996 **Weight:**
Hospitalisation: **Age:** 56Y
Onset Date: 01/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Constipation Diarrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 02/03/1996 **Stopped:** 01/07/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 109985 **Seq:** 1
Reported: 16/07/1996
Hospitalisation:
Onset Date: 10/07/1996
Outcome: Not yet recovered

Gender: M
Weight: 86.00
Age: 69Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Supraventricular tachycardia Tachycardia			

Medicine Details:

VERACAPS SR (Other drug) **Reason:**
400.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 27/06/1996 **Stopped:** 12/07/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110272 **Seq:** 1 **Gender:** F
Reported: 26/07/1996 **Weight:**
Hospitalisation: **Age:** 27Y
Onset Date: 10/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Dizziness Palpitations			

Medicine Details:

ORTHOXICOL DRY COUGH MIXTURE (WITH PARACETAMOL) (Suspe **Reason:** Ac upr resp inf,mult,unsp stes
1.0 Dose Unspecified 1 time
Batch: **Started:** 10/06/1996 **Stopped:** 10/06/1996

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 07/06/1996 **Stopped:** 10/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient took aurorix one year ago.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110292 **Seq:** 1
Reported: 29/07/1996
Hospitalisation:
Onset Date: 09/07/1996
Outcome: Recovered with sequelae

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Confusional state Delirium			Rivotril (clonazepam) 0.5mg im

Medicine Details:

SERENACE (Other drug) **Reason:** Otr spec symp psychopathol nec
Oral application 5.0 Milligram Daily Oral
Batch: **Started:** 14/06/1996 **Stopped:** 17/06/1996

PROZAC (Other drug) **Reason:** Depression
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 12/06/1996

ZANTAC (Other drug) **Reason:** Other diseases of esophagus
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

HEPARIN SODIUM (Other drug) **Reason:** Prophylaxis
Injection 10.0 Thousand Internat Daily Subcutaneous
Batch: **Started:** 27/05/1996 **Stopped:** 04/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had concurrent urinary tract infection. see original report for other drugs.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110292 **Seq:** 1
Reported: 29/07/1996
Hospitalisation:
Onset Date: 09/07/1996
Outcome: Recovered with sequelae

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

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 09/07/1996 **Stopped:** 11/07/1996

Laboratory Investigations:

Additional Information:

Patient had concurrent urinary tract infection. see original report for other drugs.

Report Detail

Case Number: 110353 **Seq:** 1 **Gender:** F
Reported: 31/07/1996 **Weight:** 55.00
Hospitalisation: **Age:** 71Y
Onset Date: 12/07/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle twitching Hyperkinesia Muscle spasms	Severe	Kicking and twitching.	

Medicine Details:

WARFARIN SODIUM (Other drug) **Reason:**
7.0 Milligram Daily
Batch: **Started:** 01/06/1995 **Stopped:**

CAPOTEN (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** 01/06/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
2.0 Dose Unspecified Daily
Batch: **Started:** 12/07/1996 **Stopped:** 18/07/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110452 **Seq:** 1
Reported: 02/08/1996
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: M
Weight: 60.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression Agitation Drug ineffective Insomnia		Exacerbation of illness symptoms.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/06/1996 **Stopped:** 18/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

When dose of aurorix inceased symptoms became worse.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110498 **Seq:** 1 **Gender:** F
Reported: 05/08/1996 **Weight:** 68.00
Hospitalisation: **Age:** 55Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Increase in temporal lobe epilepsy	

Medicine Details:

TEGRETOL (Other drug) **Reason:** Partl,templobe,psychomo epilep
30.0 Milligram As necessary
Batch: **Started:** L TERM **Stopped:**

LANOXIN (Other drug) **Reason:**
250.0 Microgram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

ASPIRIN (Other drug) **Reason:**
0.5 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 28/05/1996 **Stopped:** 20/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had increase in temporal lobe epilepsy. seizures increased from one every 4-6 weeks, to daily, after about 3 weeks on aurorix.

Report Detail

Case Number: 110643 **Seq:** 1 **Gender:** M
Reported: 08/08/1996 **Weight:** 87.00
Hospitalisation: **Age:** 45Y
Onset Date: 22/07/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sensory disturbance Headache		Altered sensation feels cold in face and	

Medicine Details:

PLENDIL (Other drug) **Reason:** Essential benign hypertension
2.5 Milligram Daily
Batch: **Started:** 28/06/1996 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/07/1996 **Stopped:** 06/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 110806 **Seq:** 1 **Gender:** F
Reported: 13/08/1996 **Weight:**
Hospitalisation: **Age:** 70
Onset Date: 25/07/1996 **DOB:** 05/07/1926
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			Phenergan

Medicine Details:

RIFAMPICIN (Other drug) **Reason:** Bacterial disease nec
Oral application 1.8 Gram Daily Oral
Batch: **Started:** **Stopped:** 26/07/1996

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

COVERSYL (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ALDACTONE (Other drug) **Reason:**
25.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drugs

Report Detail

Case Number: 110806 **Seq:** 1
Reported: 13/08/1996
Hospitalisation:
Onset Date: 25/07/1996
Outcome: Not yet recovered

Gender: F
Weight:
Age: 70
DOB: 05/07/1926
Causality: Causality possible

Reaction Detail

Medicine Details:

SLOW-K (Other drug)			**Reason:**		
	2.0	Dose Unspecified	Daily		
Batch:	**Started:**	L TERM		**Stopped:**	
CIPROXIN (Suspected)			**Reason:**	Unspecified osteomyelitis	
Tablet	1.5	Gram	Daily		Oral
Batch:	**Started:**	24/07/1996		**Stopped:**	26/07/1996
AURORIX (Suspected)			**Reason:**		
	300.0	Milligram	Daily		
Batch:	**Started:**	22/07/1996		**Stopped:**	

Laboratory Investigations:

Additional Information:
See original report for other drugs

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110907 **Seq:** 1 **Gender:** F
Reported: 16/08/1996 **Weight:** 60.00
Hospitalisation: **Age:** 33Y
Onset Date: 11/08/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache	Severe	Throbbing headache - very severe	

Medicine Details:

XANAX (Other drug) **Reason:** Phobic neurosis
6.0 Milligram Daily
Batch: **Started:** 05/08/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Phobic neurosis
450.0 Milligram Daily
Batch: **Started:** 05/08/1996 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 110930 **Seq:** 1 **Gender:** F
Reported: 15/08/1996 **Weight:** 70.00
Hospitalisation: **Age:** 57Y
Onset Date: 07/07/1996 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea Palpitations			

Medicine Details:

PROVERA (Other drug) **Reason:** Menopausal symptoms
10.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

PREMARIN (Other drug) **Reason:** Menopausal symptoms
625.0 Microgram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 03/07/1996 **Stopped:** 20/07/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had same symptoms while on lower dose of aurorix in may.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111124 **Seq:** 1 **Gender:** F
Reported: 23/08/1996 **Weight:** 70.00
Hospitalisation: **Age:** 31Y
Onset Date: 07/08/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Galactorrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 31/07/1996 **Stopped:** 07/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prolactin	N < 24.2	07/08/1996	32.4	

Additional Information:

Report Detail

Case Number: 111188 **Seq:** 1 **Gender:** M
Reported: 26/08/1996 **Weight:**
Hospitalisation: **Age:** 53Y
Onset Date: 16/07/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression			

Medicine Details:

PEPCIDINE (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 16/07/1996 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111407 **Seq:** 1 **Gender:** F
Reported: 05/09/1996 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 25/08/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 16/08/1996 **Stopped:** 30/08/1996

HALCION (Suspected) **Reason:** Depression
Tablet 125.0 Milligram Daily Oral
Batch: **Started:** 23/08/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of allergy to sulpha drugs and diazepam.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111500 **Seq:** 1 **Gender:** M
Reported: 09/09/1996 **Weight:** 67.00
Hospitalisation: **Age:** 70Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug level increased		Elevated serum perhexiline concentrations	

Medicine Details:

PEXID (Interaction) **Reason:** Angina pectoris w/o hyperten
200.0 Milligram Daily
Batch: **Started:** 01/06/1995 **Stopped:** CONTIN

AROPAX (Interaction) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** 18/01/1996 **Stopped:** 27/02/1996

AURORIX (Interaction) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/05/1996 **Stopped:** CONTIN

CARDIZEM (Other drug) **Reason:** Angina pectoris w/o hyperten
180.0 Milligram Daily
Batch: **Started:** 01/05/1995 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level		11/07/1995	0.33	Perhexiline serum
	Serum drug level		15/08/1995	0.31	Perhexiline serum
	Serum drug level		07/09/1995	0.28	Perhexiline serum
	Serum drug level		25/01/1996	0.88	Perhexiline serum
	Serum drug level		13/02/1996	1.02	Perhexiline serum
	Serum drug level		20/02/1996	1.54	Perhexiline serum
	Serum drug level		05/03/1996	0.65	Perhexiline serum

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111500 **Seq:** 1 **Gender:** M
Reported: 09/09/1996 **Weight:** 67.00
Hospitalisation: **Age:** 70Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

CARTIA (Other drug) **Reason:** Chron isch heart dis no hyper
100.0 Milligram Daily
Batch: **Started:** 01/09/1995 **Stopped:** CONTIN

ENDONE (Other drug) **Reason:** Pain
15.0 Milligram Daily
Batch: **Started:** 01/09/1995 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level		19/03/1996	0.61	Perhexiline serum
	Serum drug level		22/04/1996	0.43	Perhexiline serum
	Serum drug level		20/06/1996	0.25	Perhexiline serum
	Serum drug level		01/07/1996	0.59	Perhexiline serum

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111545 **Seq:** 1 **Gender:** F
Reported: 12/09/1996 **Weight:** 94.00
Hospitalisation: **Age:** 28Y
Onset Date: 05/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal		Loss of balance - couldn't stand.	
Tremor		Shaking uncontrolable.	
Visual disturbance		Tunnel vision	Diazepam, maxolon
Dizziness			
Vomiting			

Medicine Details:

AROPAX (Suspected) **Reason:** Depression
Tablet 30.0 Milligram Daily Oral
Batch: **Started:** 20/05/1996 **Stopped:** 03/09/1996

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 05/09/1996 **Stopped:** 05/09/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111646 **Seq:** 1 **Gender:** F
Reported: 16/09/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blood creatine phosphokinase in Fatigue Myalgia			

Medicine Details:

LIPEX (Suspected) **Reason:** Othr&unspec metabolic diseases
0.0
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine	(150-160)			Cpk = 150 u/l

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111659 **Seq:** 1 **Gender:** M
Reported: 17/09/1996 **Weight:**
Hospitalisation: **Age:** 62Y
Onset Date: 27/08/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			

Medicine Details:

PANADOL (Suspected) **Reason:** Pain
Oral application 1.0 Gram As necessary Oral
Batch: **Started:** 11/08/1996 **Stopped:** CONTIN

DILANTIN (Suspected) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 11/08/1996 **Stopped:** 06/09/1996

DEXAMETHASONE (Suspected) **Reason:**
Oral application 12.0 Milligram Daily Oral
Batch: **Started:** 11/08/1996 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 11/08/1996 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 111719 **Seq:** 1 **Gender:** F
Reported: 20/09/1996 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: 05/07/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia Tremor			

Medicine Details:

ZOLOFT (Suspected) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 05/07/1996 **Stopped:**

AURORIX (Suspected) **Reason:**
1.2 Gram Daily
Batch: **Started:** S TERM **Stopped:** 05/07/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111724 **Seq:** 1 **Gender:** M
Reported: 20/09/1996 **Weight:**
Hospitalisation: **Age:** 36Y
Onset Date: 31/08/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective		? reversal of antidepressant activity of	Augmentin ceased, depression resolving.

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** 01/07/1996 **Stopped:** CONTIN

AUGMENTIN (Interaction) **Reason:** Chronic sinusitis
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** 22/08/1996 **Stopped:** 30/08/1996

NITRAZEPAM (Other drug) **Reason:**
Tablet 6.2 Milligram Reducing Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

"on aurorix constant dose since july 1996. took augmentin for sinusitis - near the end of 1st course, then by halfway through 2nd course: gradual --> full ? reversal of antidepressant activity of aurorix. felt original symptoms coming back. ? augmentin blocked action of aurorix or greatly enhanced its metabolism." - 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111760 **Seq:** 1 **Gender:** F
Reported: 23/09/1996 **Weight:**
Hospitalisation: **Age:** 47
Onset Date: 18/09/1996 **DOB:** 28/11/1948
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Peripheral ischaemia		Cold feet	
Dyskinesia		Facial contortions	
Chills		Shaking	
Vomiting			

Medicine Details:

MURELAX (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:** 18/09/1996

AURORIX (Interaction) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 18/09/1996

EFEXOR (Interaction) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 18/09/1996 **Stopped:** 18/09/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Aurorix stopped after morning dose 18/9/96. efexor started with 7mg evening dose on the same day - reaction then occurred.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111844 **Seq:** 1 **Gender:** F
Reported: 25/09/1996 **Weight:** 57.00
Hospitalisation: **Age:** 48Y
Onset Date: 31/08/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 18/07/1996 **Stopped:** 28/09/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111902 **Seq:** 1 **Gender:** M
Reported: 30/09/1996 **Weight:** 75.00
Hospitalisation: **Age:** 27Y
Onset Date: 18/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Dizziness Hallucination Headache Nausea			

Medicine Details:

LOGICIN COUGH MIXTURE FOR DRY COUGHS (Interaction) **Reason:** Cough
Solution 40.0 Millilitre Daily Oral
Batch: **Started:** 18/09/1996 **Stopped:** 18/09/1996

AURORIX (Interaction) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 04/07/1996 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 111961 **Seq:** 1
Reported: 02/10/1996
Hospitalisation: Admitted to hospital
Onset Date: 26/09/1996
Outcome: Recovered

Gender: F
Weight: 50.00
Age: 39Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Anaphylactoid reaction Hypotension Vomiting	Severe		Adrenaline, oxygen, hydrocortisone, phenergan.

Medicine Details:

AURORIX (Suspected) **Reason:**

150.0 Milligram 1 time

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had history of 3-4 days of prozac in 1993 no effects.

Report Detail

Case Number: 112214 **Seq:** 1 **Gender:** F
Reported: 11/10/1996 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 25/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Targetoid rash lower chest, upper	

Medicine Details:

HALOPERIDOL (Other drug) **Reason:** Otr spec symp psychopathol nec
Tablet 500.0 Microgram As necessary Oral
Batch: **Started:** 04/09/1996 **Stopped:**

DILTIAZEM HYDROCHLORIDE (Other drug) **Reason:** Essential benign hypertension
Tablet, modified release 240.0 Milligram Daily Oral
Batch: **Started:** 03/09/1996 **Stopped:**

CARBAMAZEPINE (Other drug) **Reason:** Other&unspecified epilepsy
200.0 Milligram Daily
Batch: **Started:** 03/09/1996 **Stopped:**

DIGOXIN (Other drug) **Reason:** Otr&nos disord of heart rhythm
125.0 Microgram Daily
Batch: **Started:** 03/09/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Biopsy: shows parakeratosis and spongiosus with exocytosis. -> spongiotic dermatitis consistent with drug eruption.

Additional Information:

Report Detail

Case Number: 112214 **Seq:** 1 **Gender:** F
Reported: 11/10/1996 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 25/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

WARFARIN SODIUM (Other drug)			**Reason:**	Otr&nos disord of heart rhythm
	4.0 Milligram	Daily		
Batch:	**Started:** 03/09/1996		**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**	Depression
Tablet	300.0 Microgram	Daily		Oral
Batch:	**Started:** 12/09/1996		**Stopped:**	25/09/1996

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 112330 **Seq:** 1 **Gender:** F
Reported: 17/10/1996 **Weight:** 50.00
Hospitalisation: **Age:** 73Y
Onset Date: 02/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal Pancytopenia			

Medicine Details:

SERENACE (Suspected) **Reason:**
1.5 Milligram Daily
Batch: **Started:** 04/07/1996 **Stopped:** 29/08/1996

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 25/06/1996 **Stopped:** 29/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		02/09/1996	63	
	ALT = SGPT		13/09/1996	17	
	Haemoglobin	(120-160)	02/09/1996	95	
	Haemoglobin	(120-160)	13/09/1996	117	

Additional Information:

Report Detail

Case Number: 112430 **Seq:** 1
Reported: 22/10/1996
Hospitalisation:
Onset Date: 10/10/1996
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 32Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suppressed lactation		Milk supply dried up	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 06/09/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 10/10/1996 **Stopped:** 17/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had previously taken aurorix with side effect of headache, but had not been lactating.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 112587 **Seq:** 1 **Gender:** F
Reported: 28/10/1996 **Weight:** 85.00
Hospitalisation: **Age:** 52Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperprolactinaemia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 4.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/03/1996 **Stopped:**

PROGYNOVA (Suspected) **Reason:** Menopausal symptoms
Tablet 2.0 Milligram Daily Oral
Batch: **Started:** 01/01/1991 **Stopped:**

PRIMOLUT-N (Suspected) **Reason:** Menopausal symptoms
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prolactin	64 - 472	08/07/1996	1844	

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 112625 **Seq:** 1 **Gender:** M
Reported: 29/10/1996 **Weight:**
Hospitalisation: **Age:** 66Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 01/05/1996 **Stopped:** 28/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	GGT = SGGT =				Ggt rised from 905 to 1445.

Additional Information:

Primary biliary cirrhosis. 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: 112730 **Seq:** 1 **Gender:** F
Reported: 01/11/1996 **Weight:** 60.00
Hospitalisation: **Age:** 22Y
Onset Date: 26/10/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

VENTOLIN (Other drug) **Reason:** Asthma

Inhalation 2.0 Dose Unspecified As necessary Inhalation

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

AURORIX (Suspected) **Reason:** Depression

300.0 Milligram Daily

Batch: **Started:** 22/10/1996 **Stopped:** 27/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergy to keflex previously.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 112761 **Seq:** 1 **Gender:** M
Reported: 04/11/1996 **Weight:** 70.00
Hospitalisation: **Age:** 54Y
Onset Date: 26/10/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash erythematous			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 17/10/1996 **Stopped:** 29/10/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 112950 **Seq:** 1 **Gender:** F
Reported: 08/11/1996 **Weight:** 63.00
Hospitalisation: **Age:** 68
Onset Date: **DOB:** 25/10/1928
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Fluid restriction
Nausea	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

Batch: **Started:** **Stopped:** 2W

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium	(135-146)	23/10/1996	129	
	Sodium	(135-146)	24/10/1996	130	
	Sodium	(135-146)	29/10/1996	134	

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113013 **Seq:** 1 **Gender:** M
Reported: 11/11/1996 **Weight:** 69.00
Hospitalisation: **Age:** 72Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia Agitation		Constantly moving arms and legs.	 Lorazepam

Medicine Details:

PREDNISOLONE (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 01/03/1996 **Stopped:**

TEMAZEPAM (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/05/1996 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 20/09/1996 **Stopped:** 24/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Has been hyponatraemic-na+ 126 (136-146) for several months. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113095 **Seq:** 1 **Gender:** U
Reported: 12/11/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise		Feels terrible	
Tinnitus		Ringing and buzzing in ears	
Dizziness			

Medicine Details:

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 113125 **Seq:** 1 **Gender:** F
Reported: 13/11/1996 **Weight:** 55.00
Hospitalisation: **Age:** 40Y
Onset Date: 03/11/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hemiparesis Hypoaesthesia		Left sided weakness	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 02/11/1996 **Stopped:** 03/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 113243 **Seq:** 1 **Gender:** F
Reported: 19/11/1996 **Weight:** 70.00
Hospitalisation: **Age:** 75Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia		Hyperarousal	

Medicine Details:

THYROXINE SODIUM (Other drug) **Reason:**
Tablet 150.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

DILTIAZEM HYDROCHLORIDE (Other drug) **Reason:** Essential benign hypertension
Oral application 240.0 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
150.0 Milligram Daily
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113245 **Seq:** 1 **Gender:** F
Reported: 19/11/1996 **Weight:** 62.00
Hospitalisation: **Age:** 44Y
Onset Date: 06/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Hypertension	Severe		

Medicine Details:

MODECATE (Other drug) **Reason:**
0.0
Batch: **Started:** 01/02/1987 **Stopped:**

MONOPRIL (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** 01/04/1993 **Stopped:**

ALLOPURINOL (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

PREMARIN (Other drug) **Reason:** Menopausal symptoms
625.0 Microgram Daily
Batch: **Started:** 01/02/1995 **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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113245 **Seq:** 1 **Gender:** F
Reported: 19/11/1996 **Weight:** 62.00
Hospitalisation: **Age:** 44Y
Onset Date: 06/06/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/07/1995 **Stopped:**

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113272 **Seq:** 1 **Gender:** M
Reported: 22/11/1996 **Weight:** 71.00
Hospitalisation: **Age:** 71Y
Onset Date: 19/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension	Required Visit to Doctor	Mild to moderate increase in blood	Cease aurorix and change to

Medicine Details:

MONOPRIL (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

NORVASC (Other drug) **Reason:**
7.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ISOPTIN SR (Other drug) **Reason:**
240.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

HYDRALAZINE HYDROCHLORIDE (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Average bp 16-18/9/96(n=8) 19-21/9/96(n=8) 26-28/9/96(n=8) lying sbp 173.5 177.3 165.8 lying dbp 68.3 70.5 67.8 standing sbp 149.0 152.5 138.0 standing dbp 63.0 64.0 62.8

Additional Information:

Report Detail

Case Number: 113272 **Seq:** 1 **Gender:** M
Reported: 22/11/1996 **Weight:** 71.00
Hospitalisation: **Age:** 71Y
Onset Date: 19/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

MADOPAR (Other drug)			**Reason:**		
	3.0	Dose Unspecified	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
SALAZOPYRIN (Other drug)			**Reason:**		
Injection	0.0		Daily	Intravenous	
Batch:	**Started:**		L TERM	**Stopped:**	
ORUDIS SR (Other drug)			**Reason:**		
	200.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	
ZYLOPRIM (Other drug)			**Reason:**		
	100.0	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113272 **Seq:** 1 **Gender:** M
Reported: 22/11/1996 **Weight:** 71.00
Hospitalisation: **Age:** 71Y
Onset Date: 19/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ASPIRIN (Other drug)			**Reason:**	
	150.0 Milligram	Daily		
Batch:	**Started:**	L TERM	**Stopped:**	
VOLTAREN (Other drug)			**Reason:**	
	20.0 Milligram	Daily		
Batch:	**Started:**	L TERM	**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression
	2.0 Dose Unspecified	Daily		
Batch:	**Started:** 19/09/1996		**Stopped:**	21/09/1996

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113298 **Seq:** 1 **Gender:** F
Reported: 21/11/1996 **Weight:** 66.00
Hospitalisation: **Age:** 36Y
Onset Date: 31/08/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Menstrual disorder		Periods are closer together	Oral contraceptive to control

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 14/08/1996 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113503 **Seq:** 1
Reported: 28/11/1996
Hospitalisation:
Onset Date: 22/11/1996
Outcome: Not yet recovered

Gender: F
Weight: 95.00
Age: 44Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Dizziness Flushing Nausea Vomiting			

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** 06/11/1996 **Stopped:**

PLENDIL ER (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 01/11/1994 **Stopped:**

INDERAL (Other drug) **Reason:** Essential benign hypertension
80.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:**

ENDONE (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 01/08/1996 **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. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 113503 **Seq:** 1
Reported: 28/11/1996
Hospitalisation:
Onset Date: 22/11/1996
Outcome: Not yet recovered

Gender: F
Weight: 95.00
Age: 44Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

INDOCID (Other drug)			**Reason:**	
	150.0 Milligram	Daily		
Batch:	**Started:** 21/11/1996		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression
	300.0 Milligram	Daily		
Batch:	**Started:** 22/11/1996		**Stopped:**	23/11/1996

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113537 **Seq:** 1 **Gender:** F
Reported: 29/11/1996 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: 11/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 13/12/1994 **Stopped:** 11/09/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was rechallenged on aurorix with same reaction occurring

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113689 **Seq:** 1 **Gender:** M
Reported: 05/12/1996 **Weight:**
Hospitalisation: **Age:** 41Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Epistaxis			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
900.0 Milligram Daily
Batch: **Started:** 01/10/1993 **Stopped:** 28/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient stopped medication two or three times over a three year period and epistaxis would commence again when resumed. 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: 113815 **Seq:** 1 **Gender:** M
Reported: 11/12/1996 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 12/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Insomnia			

Medicine Details:

ASPIRIN (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

CAPOTEN (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

LANOXIN (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

POTASSIUM CHLORIDE (Other drug) **Reason:**
0.0
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 113815 **Seq:** 1 **Gender:** M
Reported: 11/12/1996 **Weight:**
Hospitalisation: **Age:** 71Y
Onset Date: 12/09/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

UREX (Other drug)			**Reason:**		
	0.0				
Batch:	**Started:**	L TERM		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**	10/09/1996		**Stopped:**	20/09/1996

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114081 **Seq:** 1
Reported: 20/12/1996
Hospitalisation:
Onset Date: 01/11/1996
Outcome: Not yet recovered

Gender: F
Weight:
Age: 36Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dermographism Pruritus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

300.0 Milligram Daily

Batch: **Started:** 17/10/1996 **Stopped:** 13/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 114335 **Seq:** 1
Reported: 06/01/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: M
Weight:
Age: 74
DOB: 09/04/1922
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea Haemoptysis Lung infiltration Respiratory failure			Prednisone, transferred to icu, intubated and ventilated.

Medicine Details:

BECLOMETHASONE DIPROPIONATE (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AMIODARONE HYDROCHLORIDE (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** 17/09/1996 **Stopped:**

FRUSEMIDE (Suspected) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

SLOW-K (Suspected) **Reason:**
1.2 Gram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Histology				Autopsy 15/10/96: the interstitial pneumonitis is associated with a mostly sparse lumphocytic inflammatory cell infiltrate and focal alveolar haemorrhage

Additional Information:

This patient had a complicated past medical history which included longstand- ing cal with bullous emphysema, pneumonia, angina with a quadruple coronary artery bypass graft in 1981, carcinoma of the prostate treated with radiotherapy and a right total hip replacement. patient died 13/10/96. committee comment: include in annual fatality analysis

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114335 **Seq:** 1
Reported: 06/01/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death, maybe drug

Gender: M
Weight:
Age: 74
DOB: 09/04/1922
Causality: Causality possible

Reaction Detail

Medicine Details:

LANOXIN PG (Suspected)			**Reason:**	
	1.2	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
ISOSORBIDE MONONITRATE (Suspected)			**Reason:**	
	120.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
MOCLOBEMIDE (Suspected)			**Reason:**	
	300.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

This patient had a complicated past medical history which included longstand- ing cal with bullous emphysema, pneumonia, angina with a quadruple coronary artery bypass graft in 1981, carcinoma of the prostate treated with radiotherapy and a right total hip replacement. patient died 13/10/96. committee comment: include in annual fatality analysis

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114400 **Seq:** 1
Reported: 08/01/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chills Confusional state Flushing Serotonin syndrome Tremor Urinary incontinence			

Medicine Details:

ZOLOFT (Interaction) **Reason:**
Tablet 50.0 Milligram 1 time Oral
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:**
300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114533 **Seq:** 1 **Gender:** U
Reported: 14/01/1997 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain		Supra pubic pain	
Agitation			
Chills			
Dyspnoea			
Dysuria			
Hyperhidrosis			
Hypertension			Cessation of sertraline.
Nausea			
Pyrexia			

Medicine Details:

ZOLOFT (Suspected) **Reason:** Depression
100.0 Milligram Daily
Batch: **Started:** 14/12/1996 **Stopped:** 14/12/1996

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 12/12/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114584 **Seq:** 1 **Gender:** U
Reported: 16/01/1997 **Weight:** 95.00
Hospitalisation: **Age:** 88
Onset Date: **DOB:** 26/12/1908
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 01/09/1996 **Stopped:** 28/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	(1-40)	28/11/1996	849	
	ALT = SGPT	(1-40)	02/12/1996	295	
	AST = SGOT	(1-45)	28/11/1996	211	
	AST = SGOT	(1-45)	02/12/1996	57	
	GGT = SGGT =	(1-40)	28/11/1996	86	
	GGT = SGGT =	(1-40)	02/12/1996	121	

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114649 **Seq:** 1 **Gender:** M
Reported: 20/01/1997 **Weight:** 73.00
Hospitalisation: **Age:** 21Y
Onset Date: 09/01/2003 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Accommodation disorder Dyskinesia Dyskinesia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily Oral
Batch: **Started:** 14/11/1996 **Stopped:** 09/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Partical facial paresis and temporarylost of eye movement and focusint ability several times involuntary tongue swallowing.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114670 **Seq:** 1
Reported: 22/01/1997
Hospitalisation:
Onset Date: 28/12/1996
Outcome: Not yet recovered

Gender: F
Weight: 90.00
Age: 26
DOB: 24/07/1970
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Menstrual disorder			

Medicine Details:

AURORIX (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114803 **Seq:** 1 **Gender:** F
Reported: 28/01/1997 **Weight:** 58.00
Hospitalisation: **Age:** 77
Onset Date: **DOB:** 30/03/1919
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Diarrhoea Nocturia Pollakiuria		Falls, secondary to dizziness and pour	

Medicine Details:

PANADEINE (Other drug) **Reason:** Other rheumatoid arthritis
Tablet 0.0 As necessary Oral
Batch: **Started:** 01/01/1989 **Stopped:**

CARDIZEM CD (Other drug) **Reason:** Essential benign hypertension
Capsule 240.0 Milligram Daily Oral
Batch: **Started:** 01/03/1990 **Stopped:**

LIPEX (Other drug) **Reason:** Othr&unspec metabolic diseases
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 01/03/1995 **Stopped:**

PETHIDINE HYDROCHLORIDE (Other drug) **Reason:** Pain
40.0 Milligram Daily
Batch: **Started:** 01/11/1991 **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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114803 **Seq:** 1
Reported: 28/01/1997
Hospitalisation:
Onset Date:
Outcome: Unknown

Gender: F
Weight: 58.00
Age: 77
DOB: 30/03/1919
Causality: Causality probable

Reaction Detail

Medicine Details:

SEREPAX (Other drug)		**Reason:**	Anxiety neurosis	
	30.0 Milligram	Daily		
Batch:	**Started:** 01/01/1989		**Stopped:**	
AURORIX (Suspected)		**Reason:**	Depression	
Tablet	450.0 Milligram	Daily	Oral	
Batch:	**Started:** 12/10/1996		**Stopped:**	24/12/1996

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 114864 **Seq:** 1 **Gender:** F
Reported: 30/01/1997 **Weight:**
Hospitalisation: **Age:** 30Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Dizziness Fatigue			

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 114906 **Seq:** 1 **Gender:** F
Reported: 03/02/1997 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 29/01/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyskinesia		Numb tongue, difficulty speaking.	Cogentin

Medicine Details:

NORVASC (Other drug) **Reason:** Essential benign hypertension
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

SERENACE (Suspected) **Reason:**
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** 28/01/1997 **Stopped:** 29/01/1997

TEMAZE (Suspected) **Reason:**
Capsule 10.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115092 **Seq:** 1
Reported: 10/02/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight:
Age: 48Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Salivary hypersecretion Abdominal pain Nervousness Tremor		Patient was suffering serotonergic	

Medicine Details:

ZOLOFT (Interaction) **Reason:**

Tablet 100.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 3D

AURORIX (Interaction) **Reason:**

Tablet 900.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 1Y

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 115095 **Seq:** 1 **Gender:** F
Reported: 10/02/1997 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: 24/12/1996 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthritis		Costochondritis	
Antinuclear antibody positive		Postive ana antibodies	

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis
300.0 Milligram Daily
Batch: **Started:** 07/08/1996 **Stopped:** 30/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Antinuclear factor		06/01/1997	160	
	Antinuclear factor		24/01/1997	160	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115097 **Seq:** 1 **Gender:** F
Reported: 10/02/1997 **Weight:**
Hospitalisation: **Age:** 40Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Hyperkinesia Tremor			

Medicine Details:

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

AURORIX (Interaction) **Reason:**
900.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 115112 **Seq:** 1 **Gender:** M
Reported: 10/02/1997 **Weight:**
Hospitalisation: **Age:** 74
Onset Date: 31/12/1996 **DOB:** 24/10/1922
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dementia Hypertonia Hypoglycaemia Hypothermia			

Medicine Details:

NEOTIGASON (Suspected) **Reason:** Other psoriasis
Capsule 25.0 Milligram Daily Oral
Batch: **Started:** 01/04/1996 **Stopped:**

THIAMINE (Suspected) **Reason:** Unspecified psychosis
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

ASPIRIN (Suspected) **Reason:**
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

MOGADON (Suspected) **Reason:** Nausea and vomiting
Tablet 10.0 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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115112 **Seq:** 1
Reported: 10/02/1997
Hospitalisation:
Onset Date: 31/12/1996
Outcome: Not yet recovered

Gender: M
Weight:
Age: 74
DOB: 24/10/1922
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** **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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115190 **Seq:** 1
Reported: 13/02/1997
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: M
Weight: 50.00
Age: 74Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Osteoarthritis Arthralgia		Effusions in knees.	

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 115196 **Seq:** 1
Reported: 13/02/1997
Hospitalisation:
Onset Date: 31/12/1996
Outcome: Not yet recovered

Gender: M
Weight: 67.00
Age: 38Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression Drug ineffective			

Medicine Details:

ZANTAC (Suspected) **Reason:**

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 28/03/1994 **Stopped:** 28/12/1996

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115387 **Seq:** 1
Reported: 18/02/1997
Hospitalisation: Admitted to hospital
Onset Date: 14/01/1997
Outcome: Recovered

Gender: M
Weight: 65.00
Age: 42Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia		39.7 degrees c.	
Hypertonia		Extreme leadpipe rigidity.	
Tachycardia		Pulse rate ot 120 bpm	
Agitation			
Blood creatine phosphokinase in			
Hyperhidrosis			
Neuroleptic malignant syndrome			Dantrolene 2mg/kg x 4 doses, intubation for airway protection, diazepam
Serotonin syndrome			
Tremor			

Medicine Details:

LORAZEPAM (Other drug) **Reason:** Anxiety neurosis
4.0 Millilitre Daily
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **Stopped:** 14/01/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine phosphokinase				Initial creatinine kinase level was normal (317 u/l) but on the next day rose to 30139 u/l and peaked 46594 u/l.

Additional Information:

"altered mental state, rigidity, increased cpk to about 30,000 units/l, fever, and tachycardia. dantrolene terminated the syndrome - tachycardia, fever, rigidity resolved in 4 hours; mental state returned to normal in 1 day." reaction occurred after 4 days of extreme heat (max temp in excess of 40 degrees c). 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115450 **Seq:** 1
Reported: 21/02/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight: 45.00
Age: 25Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Crying Serotonin syndrome			

Medicine Details:

PROZAC (Interaction) **Reason:** Depression
40.0 Milligram Daily
Batch: **Started:** 13/03/1996 **Stopped:** 15/04/1996

AURORIX (Interaction) **Reason:** Depression
900.0 Milligram Daily
Batch: **Started:** 15/04/1996 **Stopped:**

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

AMOXIL (Other drug) **Reason:** Othr diseases of urinary tract
1.5 Gram Daily
Batch: **Started:** 12/04/1996 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 115450 **Seq:** 1
Reported: 21/02/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight: 45.00
Age: 25Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ZANTAC (Other drug) **Reason:** Gastritis&duodenitis

300.0 Milligram Daily

Batch: **Started:** 02/04/1996 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115462 **Seq:** 1 **Gender:** M
Reported: 19/02/1997 **Weight:** 72.00
Hospitalisation: **Age:** 34Y
Onset Date: 14/02/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delusion Paranoia	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 12/02/1997 **Stopped:** 14/02/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115469 **Seq:** 1 **Gender:** F
Reported: 24/02/1997 **Weight:** 72.00
Hospitalisation: **Age:** 42Y
Onset Date: 31/01/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension			

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** 01/02/1996 **Stopped:**

AURORIX (Suspected) **Reason:**
0.0
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. the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month.

Report Detail

Case Number: 115482 **Seq:** 1 **Gender:** M
Reported: 24/02/1997 **Weight:** 90.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erectile dysfunction			

Medicine Details:

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

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

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

ECOTRIN (Other drug) **Reason:**
0.0
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115482 **Seq:** 1
Reported: 24/02/1997
Hospitalisation:
Onset Date:
Outcome: Recovered

Gender: M
Weight: 90.00
Age: 49Y
DOB:
Causality: Causality probable

Reaction Detail

Medicine Details:

PANADEINE FORTE (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
PEPCIDINE (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
ZYLOPRIM (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**		**Stopped:**
AURORIX (Suspected)		**Reason:**	
	0.0		
Batch:	**Started:**	01/07/1996	**Stopped:**

Laboratory Investigations:

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115654 **Seq:** 1 **Gender:** F
Reported: 26/02/1997 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Agitation			

Medicine Details:

OESTROGENS NOS (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 115852 **Seq:** 1 **Gender:** M
Reported: 12/03/1997 **Weight:** 75.00
Hospitalisation: **Age:** 24Y
Onset Date: 17/02/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nausea Orthostatic hypotension Syncope			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 16/02/1997 **Stopped:** 06/03/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 115956 **Seq:** 1 **Gender:** M
Reported: 12/03/1997 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Neurosis Agitation Insomnia		Agoraphobia	

Medicine Details:

NORVASC (Suspected)		**Reason:**	
Tablet	0.0		Oral
Batch:	**Started:**	**Stopped:**	
AURORIX (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116051 **Seq:** 1 **Gender:** F
Reported: 18/03/1997 **Weight:** 48.00
Hospitalisation: **Age:** 74Y
Onset Date: 05/03/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Vertigo			Haloperidol, clonazepam.

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 116109 **Seq:** 1 **Gender:** M
Reported: 18/03/1997 **Weight:** 65.00
Hospitalisation: **Age:** 31Y
Onset Date: 08/03/1997 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Back pain Chest pain	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 08/03/1997 **Stopped:** 11/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Same reaction occurred 2 years ago on aurorix treatment.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116144 **Seq:** 1
Reported: 19/03/1997
Hospitalisation:
Onset Date: 09/03/1997
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 35Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash erythematous Skin exfoliation			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

300.0 Milligram Daily

Batch: **Started:** 07/03/1997 **Stopped:** 10/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 116194 **Seq:** 1 **Gender:** F
Reported: 20/03/1997 **Weight:**
Hospitalisation: **Age:** 50Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperkinesia			

Medicine Details:

THIORIDAZINE HYDROCHLORIDE (Other drug) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

DIAZEPAM (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 21/02/1997 **Stopped:** 03/03/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116262 **Seq:** 1 **Gender:** F
Reported: 21/03/1997 **Weight:**
Hospitalisation: **Age:** 36Y
Onset Date: 31/01/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Jaundice Pruritus			Cholestyramine, phenergan, oxazepam.

Medicine Details:

ORAL CONTRACEPTIVE NOS (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:** 28/10/1996

AURORIX (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:** 24/02/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	0 - 55 U/L	06/03/1997	150	
	ALT = SGPT	0 - 55 U/L	10/03/1997	170	
	ALT = SGPT	0 - 55 U/L	11/03/1997	194	
	ALT = SGPT	0 - 55 U/L	17/03/1997	122	
	AST = SGOT	0 - 45 U/L	06/03/1997	80	
	Bilirubin	6 - 24 UMOL/L	06/03/1997	57	
	Bilirubin	6 - 24 UMOL/L	10/03/1997	46	

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.

Report Detail

Case Number: 116262 **Seq:** 1 **Gender:** F
Reported: 21/03/1997 **Weight:**
Hospitalisation: **Age:** 36Y
Onset Date: 31/01/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ORAL CONTRACEPTIVE NOS (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**	**Stopped:**	28/10/1996
AURORIX (Suspected)		**Reason:**	
Tablet	0.0		Oral
Batch:	**Started:**	**Stopped:**	24/02/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Bilirubin	6 - 24 UMOL/L	11/03/1997	44	
	Bilirubin	6 - 24 UMOL/L	17/03/1997	29	
	GGT = SGGT =	0 - 60 U/L	06/03/1997	32	
	GGT = SGGT =	0 - 60 U/L	17/03/1997	27	
	GGT = SGGT =	0 - 60 U/L	10/03/1997	31	
	GGT = SGGT =	0 - 60 U/L	11/03/1997	32	
	Immunology				Anti-reticulin antibody: positive

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116262 **Seq:** 1
Reported: 21/03/1997
Hospitalisation:
Onset Date: 31/01/1997
Outcome: Not yet recovered

Gender: F
Weight:
Age: 36Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ORAL CONTRACEPTIVE NOS (Other drug)		**Reason:**	
	0.0		
Batch:	**Started:**	**Stopped:**	28/10/1996
AURORIX (Suspected)		**Reason:**	
Tablet	0.0		Oral
Batch:	**Started:**	**Stopped:**	24/02/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Liver biopsy				Shown on liver biopsy hepatitis cholestatic associated with lobular inflammation.
	SAP = ALP	30 - 110 U/L	06/03/1997	283	
	SAP = ALP	30 - 110 U/L	10/03/1997	253	
	SAP = ALP	30 - 110 U/L	11/03/1997	254	
	SAP = ALP	30 - 110 U/L	17/03/1997	213	

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116526 **Seq:** 1 **Gender:** M
Reported: 08/04/1997 **Weight:** 65.00
Hospitalisation: **Age:** 62Y
Onset Date: 02/04/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mania		Crawling around like a dog	
Speech disorder		Mumbling	
Apathy		Withdrawn	
Syncope			

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis
300.0 Milligram Daily
Batch: **Started:** 30/03/1997 **Stopped:** 02/04/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Started aurorix 5 days earlier for anxiety and sleeping. ok for 4 days than did not sleep over night, felt something 'snapin' his head. began crawling around lounge like a dog and than collapsed. mumbling and withdrawn.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116554 **Seq:** 1 **Gender:** M
Reported: 08/04/1997 **Weight:**
Hospitalisation: **Age:** AD
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Convulsion		Tonic-clonic seisure	

Medicine Details:

ZOLOFT (Interaction) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Past history of severe carotid vascular disease and recurrent strokes.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116574 **Seq:** 1 **Gender:** M
Reported: 09/04/1997 **Weight:** 98.00
Hospitalisation: **Age:** 56Y
Onset Date: 31/10/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia		Sentasation of skin crawling along back,	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 01/06/1996 **Stopped:** 28/02/1997

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116761 **Seq:** 1 **Gender:** F
Reported: 11/04/1997 **Weight:** 57.00
Hospitalisation: **Age:** 48Y
Onset Date: 18/04/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido increased			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 18/04/1996 **Stopped:** 18/07/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Aropax causes the opposite.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116816 **Seq:** 1 **Gender:** F
Reported: 16/04/1997 **Weight:** 65.00
Hospitalisation: **Age:** 31Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Talipes		Baby boy with bilateral club feet.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 900.0 Milligram Daily Oral

Batch: **Started:** 01/06/1995 **Stopped:** 28/01/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

"mother took aurorix 450 mg bd for the previous 7 months prior to delivery up to 2-3 weeks after conceiving her child." 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: 116929 **Seq:** 1 **Gender:** F
Reported: 21/04/1997 **Weight:** 58.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Urticaria			

Medicine Details:

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

VENTOLIN (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** L TERM **Stopped:**

FLIXOTIDE (Other drug) **Reason:**
6.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

RHINOCORT (Other drug) **Reason:**
1.0 Dose Unspecified 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 116929 **Seq:** 1 **Gender:** F
Reported: 21/04/1997 **Weight:** 58.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
1.0 Dose Unspecified Daily
Batch: **Started:** 01/04/1996 **Stopped:** 16/04/1996

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117221 **Seq:** 1 **Gender:** F
Reported: 05/05/1997 **Weight:** 43.00
Hospitalisation: **Age:** 78Y
Onset Date: 19/03/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vasculitis Erythema multiforme		Vasculitic rash of hands and feet.	

Medicine Details:

ESTRADERM (Other drug) **Reason:**
50.0 Milligram Weekly
Batch: **Started:** L TERM **Stopped:**

PARACETAMOL (Other drug) **Reason:** Pain
2.0 Dose Unspecified As necessary
Batch: **Started:** L TERM **Stopped:**

METOCLOPRAMIDE HYDROCHLORIDE (Other drug) **Reason:** Nausea and vomiting
30.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 21/03/1997

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 16/03/1997 **Stopped:** 20/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Skin biopsy showed spongiosis and vasculitis.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117528 **Seq:** 1 **Gender:** F
Reported: 13/05/1997 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Anorexia Gastritis Nausea Weight decreased			

Medicine Details:

LISINOPRIL (Other drug) **Reason:** Congestive heart failure
5.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

FRUSEMIDE (Other drug) **Reason:** Congestive heart failure
80.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

ASPIRIN (Other drug) **Reason:** Chron isch heart dis no hyper
100.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

NITRADISC (Other drug) **Reason:** Chron isch heart dis no hyper
Cream 25.0 Milligram Daily Transdermal
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Endoscopy: moderate chronic gastritis.

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: 117528 **Seq:** 1 **Gender:** F
Reported: 13/05/1997 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

FOSAMAX (Suspected)		**Reason:**	Osteoporosis
	10.0 Milligram	Daily	
Batch:	**Started:** 01/02/1997		**Stopped:** 27/03/1997
MOCLOBEMIDE (Suspected)		**Reason:**	Depression
	300.0 Milligram	Daily	
Batch:	**Started:** 01/01/1997		**Stopped:** 02/05/1997

Laboratory Investigations:

Additional Information:

the dosage start date is not necessarily accurate but indicates that start occurred sometime during the month. the dosage 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117581 **Seq:** 1
Reported: 15/05/1997
Hospitalisation:
Onset Date: 06/05/1997
Outcome: Not yet recovered

Gender: M
Weight: 80.00
Age: 68Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness			
Dyspnoea			
Headache			
Palpitations			
Tinnitus			
Visual disturbance			

Medicine Details:

ASPIRIN (Other drug) **Reason:**
0.5 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 06/05/1997 **Stopped:** 08/05/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117815 **Seq:** 1 **Gender:** F
Reported: 26/05/1997 **Weight:**
Hospitalisation: **Age:** 81Y
Onset Date: 09/05/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			

Medicine Details:

PAROXETINE HYDROCHLORIDE (Suspected) **Reason:**
0.0
Batch: **Started:** 18/04/1997 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/04/1997 **Stopped:**

LISINOPRIL (Suspected) **Reason:**
Oral application 5.0 Milligram Daily Oral
Batch: **Started:** 28/04/1997 **Stopped:**

AMLODIPINE (Suspected) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** 13/02/1997 **Stopped:** 22/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

On paroxetine previously and developed hyponatraemia 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117815 **Seq:** 1
Reported: 26/05/1997
Hospitalisation:
Onset Date: 09/05/1997
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

COLCHICINE (Suspected)			**Reason:**	
Tablet	1.0	Dose Unspecified	Daily	Oral
Batch:	**Started:**	13/02/1997	**Stopped:**	
FERRO-GRADUMET (Suspected)			**Reason:**	
Tablet	1.0	Dose Unspecified	Daily	Oral
Batch:	**Started:**	01/04/1997	**Stopped:**	
ASCORBIC ACID (Suspected)			**Reason:**	
Tablet	1.0	Dose Unspecified	Daily	Oral
Batch:	**Started:**	01/04/1997	**Stopped:**	

Laboratory Investigations:

Additional Information:

On paroxetine previously and developed hyponatraemia 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117880 **Seq:** 1 **Gender:** F
Reported: 27/05/1997 **Weight:** 65.00
Hospitalisation: **Age:** 70Y
Onset Date: 22/05/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Angina pectoris Anxiety Dizziness Headache Nausea			

Medicine Details:

NITROLINGUAL SPRAY (Other drug) **Reason:** Angina pectoris w/o hyperten
Inhalation 2.0 Dose Unspecified As necessary Sublingual
Batch: **Started:** **Stopped:**

TRANSIDERM-NITRO (Other drug) **Reason:** Angina pectoris w/o hyperten
50.0 Milligram Daily
Batch: **Started:** 16/05/1997 **Stopped:**

CORDARONE X (Other drug) **Reason:** Otr&nos disord of heart rhythm
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

NORVASC (Other drug) **Reason:** Essential benign hypertension
5.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117880 **Seq:** 1 **Gender:** F
Reported: 27/05/1997 **Weight:** 65.00
Hospitalisation: **Age:** 70Y
Onset Date: 22/05/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

BETOPTIC S (Other drug)			**Reason:**	Unspecified glaucoma	
Eye Drops	4.0	Dose Unspecified	Daily		Ophthalmic
Batch:	**Started:**			**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
	150.0	Milligram	1 time		
Batch:	**Started:**	22/05/1997		**Stopped:**	22/05/1997

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117910 **Seq:** 1 **Gender:** F
Reported: 28/05/1997 **Weight:**
Hospitalisation: **Age:** 30Y
Onset Date: 31/07/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Seborrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/07/1996 **Stopped:** 28/03/1997

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 117912 **Seq:** 1
Reported: 28/05/1997
Hospitalisation:
Onset Date: 12/05/1997
Outcome: Not yet recovered

Gender: M
Weight: 94.00
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Aggression Diarrhoea Disturbance in attention Nausea Pollakiuria Tinnitus Vertigo			

Medicine Details:

PRINIVIL (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 01/12/1996 **Stopped:**

LOSEC (Other drug) **Reason:** Other diseases of esophagus
10.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 10/05/1997 **Stopped:** 17/05/1997

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.

Report Detail

Case Number: 118001 **Seq:** 1 **Gender:** U
Reported: 02/06/1997 **Weight:**
Hospitalisation: **Age:** 68Y
Onset Date: 02/04/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 01/02/1997 **Stopped:** 02/04/1997

CAPOTEN (Suspected) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

ISOPTIN (Suspected) **Reason:** Essential benign hypertension
320.0 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118095 **Seq:** 1
Reported: 05/06/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight: 0.00
Age: 66Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			
Hyponatraemia			
Inappropriate antidiuretic hormon			
Syncope			
Confusional state			

Medicine Details:

LOMOTIL (Suspected) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:** 05/05/1997

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 05/05/1997

TRITACE (Suspected) **Reason:** Essential benign hypertension
2.5 Milligram Daily
Batch: **Started:** **Stopped:** 05/05/1997

DIAZEPAM (Suspected) **Reason:** Specific disorders of sleep
5.0 Milligram Daily
Batch: **Started:** **Stopped:** 05/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
16/12/2002	Sodium		10/09/2003	109	
16/12/2002	Sodium		10/09/2003	136	

Additional Information:

Report Detail

Case Number: 118123 **Seq:** 1
Reported: 06/06/1997
Hospitalisation: Admitted to hospital
Onset Date: 10/01/1997
Outcome: Recovered

Gender: M
Weight:
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Confusional state			

Medicine Details:

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

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

FERRO-GRADUMET (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient commenced on moclobemide 2 weeks prior to adverse event dose was increase from 150mg alternate days,to 150mg daily,3 days prior to adverse event. agitation rsolved when dose returned to 150mg alternate days.

Report Detail

Case Number: 118123 **Seq:** 1
Reported: 06/06/1997
Hospitalisation: Admitted to hospital
Onset Date: 10/01/1997
Outcome: Recovered

Gender: M
Weight:
Age: 75Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ASPIRIN (Other drug)		**Reason:**	
	1.0 Dose Unspecified	Daily	
Batch:	**Started:**		**Stopped:**
TIMOPTOL (Other drug)		**Reason:**	Unspecified glaucoma
	1.0 Dose Unspecified	Daily	
Batch:	**Started:**		**Stopped:**
MOCLOBEMIDE (Suspected)		**Reason:**	Depression
	150.0 Milligram	Daily	
Batch:	**Started:** 07/01/1997		**Stopped:**

Laboratory Investigations:

Additional Information:

Patient commenced on moclobemide 2 weeks prior to adverse event dose was increase from 150mg alternate days,to 150mg daily,3 days prior to adverse event. agitation rsolved when dose returned to 150mg alternate days.

Report Detail

Case Number: 118159 **Seq:** 1 **Gender:** F
Reported: 10/06/1997 **Weight:** 60.00
Hospitalisation: **Age:** 40Y
Onset Date: 28/05/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Insomnia Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:**
450.0 Milligram Daily
Batch: **Started:** 28/05/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient reported "feeling better" 6 days after commencement of drug.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118216 **Seq:** 1 **Gender:** F
Reported: 13/06/1997 **Weight:** 90.00
Hospitalisation: **Age:** 62Y
Onset Date: 22/04/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor Flushing Hypertension		Patient felt a bit "shaky"	8 days after increasing dose to 300mg bd bp:200/130 next day bp:230/120 aurorix reduced to 150mg bd & tenormin 50mg daily started. 2 days later bp:170/95, 7days later bp:160/95 aurorix ceased parnate restarted, 2 days later tenormin ceased.

Medicine Details:

SEREPAX (Other drug) **Reason:** Disturbance of sleep
60.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

ORUVAIL SR (Other drug) **Reason:** Unspecified arthritis
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** 08/03/1993 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 07/04/1997 **Stopped:**

AURORIX (Suspected) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 16/04/1997 **Stopped:** 05/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has family history of hypertension but has never had bp problems herself.

Report Detail

Case Number: 118227 **Seq:** 1 **Gender:** F
Reported: 13/06/1997 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise		Patient exhibited toxic effects	

Medicine Details:

MACRO E (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ATROVENT (Interaction) **Reason:**
Spray 2.0 Dose Unspecified As necessary Nasal
Batch: **Started:** 14/09/1996 **Stopped:**

AURORIX (Interaction) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** 27/03/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms ceased on cessation of macro e

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118291 **Seq:** 1 **Gender:** F
Reported: 16/06/1997 **Weight:** 63.00
Hospitalisation: **Age:** 28Y
Onset Date: 07/06/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Distortion of visual perception eg. walls	
Agitation			
Diarrhoea			
Paranoia			
Tachycardia			
Tremor			
Vomiting			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 07/06/1997 **Stopped:** 08/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 118353 **Seq:** 1 **Gender:** M
Reported: 18/06/1997 **Weight:** 57.00
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased Sexual dysfunction		Libido reduced on 300mg, absent on [illegible] Nausea with erection	Ceased aurorix

Medicine Details:

ADALAT (Other drug) **Reason:** Progressive systemic sclerosis
30.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:**

PENICILLAMINE (Other drug) **Reason:** Progressive systemic sclerosis
250.0 Milligram Daily
Batch: **Started:** 01/04/1997 **Stopped:**

OMEPRAZOLE (Other drug) **Reason:** Progressive systemic sclerosis
1.0 Dose Unspecified Daily
Batch: **Started:** 01/04/1997 **Stopped:**

AURORIX (Suspected) **Reason:**
0.0
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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118595 **Seq:** 1 **Gender:** F
Reported: 26/06/1997 **Weight:** 55.00
Hospitalisation: **Age:** 31Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Intra-uterine death			

Medicine Details:

AURORIX (Suspected) **Reason:**

0.0

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

2 x miscarriage when on aurorix in early pregnancy. both times aurorix was ceased 3 and 4 days after missed period. dose 1st time:150mg x 5/d 2nd time: 150 mg x 3/d.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118744 **Seq:** 1 **Gender:** M
Reported: 02/07/1997 **Weight:** 75.00
Hospitalisation: **Age:** 48Y
Onset Date: 15/12/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Disturbance in attention Dizziness Mydriasis Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 15/11/1996

EFEXOR (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 11/12/1996 **Stopped:** 15/12/1996

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118874 **Seq:** 1 **Gender:** F
Reported: 07/07/1997 **Weight:** 63.00
Hospitalisation: **Age:** 49Y
Onset Date: 30/06/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dry mouth		Dry mouth & throat	
Hypoaesthesia		Lip went slightly numb	
Dizziness			
Somnolence			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 150.0 Milligram 1 time Oral

Batch: **Started:** 29/06/1997 **Stopped:** 30/06/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 118878 **Seq:** 1 **Gender:** F
Reported: 07/07/1997 **Weight:** 60.00
Hospitalisation: **Age:** 68Y
Onset Date: 21/12/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia Alcohol intolerance Anorexia		Altered taste	

Medicine Details:

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

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 17/12/1996 **Stopped:** 16/04/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Dose reduced to 150mg daily in feb '97. taste recovered 30/5/1997

Report Detail

Case Number: 118972 **Seq:** 1 **Gender:** F
Reported: 10/07/1997 **Weight:** 70.00
Hospitalisation: **Age:** 39Y
Onset Date: 03/06/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting Diarrhoea Nausea		Vomiting every morning	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 27/05/1997 **Stopped:** 08/07/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 119128 **Seq:** 1 **Gender:** M
Reported: 16/07/1997 **Weight:** 90.00
Hospitalisation: **Age:** 14Y
Onset Date: 04/07/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Angioedema		Lump in throat	
Oedema peripheral		Swollen hands	
Dizziness			
Dyspnoea			
Fatigue			
Headache			Phenergan

Medicine Details:

PREDNISOLONE (Other drug) **Reason:** Asthma
0.0
Batch: **Started:** **Stopped:**

RITALIN (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** 04/07/1997 **Stopped:**

AURORIX (Suspected) **Reason:**
75.0 Milligram Daily
Batch: **Started:** 04/07/1997 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 119287 **Seq:** 1 **Gender:** F
Reported: 21/07/1997 **Weight:** 65.00
Hospitalisation: **Age:** 51Y
Onset Date: 01/07/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus		Itch on inner arms and thighs	
Dyspnoea		Slight shortness of breath on exertion	
Insomnia			
Palpitations			
Tremor			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 19/05/1997 **Stopped:** 10/07/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 119374 **Seq:** 1 **Gender:** F
Reported: 23/07/1997 **Weight:**
Hospitalisation: **Age:** 87Y
Onset Date: 22/05/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Fluid restriction, ceased

Medicine Details:

RANITIDINE (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

ASPIRIN (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

FELODIPINE (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

METOPROLOL TARTRATE (Other drug) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 119374 **Seq:** 1
Reported: 23/07/1997
Hospitalisation:
Onset Date: 22/05/1997
Outcome: Not yet recovered

Gender: F
Weight:
Age: 87Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

NITRAZEPAM (Other drug)			**Reason:**	
	10.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
MOCLOBEMIDE (Suspected)			**Reason:**	Depression
	2.0	Dose Unspecified	Daily	
Batch:	**Started:**			**Stopped:** 29/05/1997

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 119414 **Seq:** 1 **Gender:** F
Reported: 24/07/1997 **Weight:** 58.00
Hospitalisation: **Age:** 74Y
Onset Date: 31/12/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Somnolence		Excessive sleepiness	Aurorix ceased

Medicine Details:

LANOXIN (Other drug) **Reason:**
312.5 Microgram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 09/04/1996 **Stopped:** 06/07/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms have ceased since aurorix was ceased. - 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 119459 **Seq:** 1 **Gender:** M
Reported: 28/07/1997 **Weight:** 76.00
Hospitalisation: **Age:** 54Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Duodenal ulcer		Disturbs dudenal ulcer.	Zantac

Medicine Details:

DIAZEPAM (Other drug) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Regulary disturbs dudenal ulcer when taking moclobemide. zantac allows patient to continue taking moclobemide by controlling pain.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 119469 **Seq:** 1 **Gender:** M
Reported: 25/07/1997 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: 13/10/1996 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			Withdraw aurorix.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 29/09/1996 **Stopped:** 28/01/1997

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: 119475 **Seq:** 1 **Gender:** F
Reported: 28/07/1997 **Weight:** 85.00
Hospitalisation: **Age:** 41Y
Onset Date: 01/07/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness		Lightheadedness	
Paraesthesia		Perioral paraesthesia	

Medicine Details:

VOLTAREN (Other drug) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 25/06/1997 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 25/06/1997 **Stopped:** 14/07/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 119746 **Seq:** 1 **Gender:** F
Reported: 07/08/1997 **Weight:** 80.00
Hospitalisation: **Age:**
Onset Date: 23/07/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Pruritus			

Medicine Details:

CHLOTRIDE (Other drug) **Reason:** Essential benign hypertension
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 19/07/1997 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120005 **Seq:** 1 **Gender:** M
Reported: 18/08/1997 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 13/07/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Dizziness Dyspnoea Orthostatic hypotension		Chest discomfort.	

Medicine Details:

ANGININE (Other drug) **Reason:**
Tablet 0.0 As necessary Sublingual
Batch: **Started:** **Stopped:** CONTIN

BETALOC (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

ASPIRIN (Other drug) **Reason:**
150.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

LOSEC (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 120005 **Seq:** 1 **Gender:** M
Reported: 18/08/1997 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 13/07/1996 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

ZESTRIL (Suspected)			**Reason:**		
	20.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	
IMDUR (Suspected)			**Reason:**		
	30.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	
NORVASC (Suspected)			**Reason:**		
	5.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	CONTIN
AURORIX (Suspected)			**Reason:**	Depression	
	300.0	Milligram	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120158 **Seq:** 1
Reported: 22/08/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight: 70.00
Age: 71Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder			Admitted to hospital, nasogastric feeding,

Medicine Details:

ZANTAC (Other drug) **Reason:**
3.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

COLOXYL (Other drug) **Reason:**
3.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

TAZAC (Other drug) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 23/01/1997 **Stopped:**

LARGACTIL (Suspected) **Reason:** Otr spec symp psychopathol nec
Injection 50.0 Milligram 1 time Intramuscular
Batch: **Started:** 09/02/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient bedbound. prior to neuroleptic administration walking independently. patient discharged to nursing home 22/07/1997. developed pneumonia and died on 9/8/1997

Report Detail

Case Number: 120158 **Seq:** 1
Reported: 22/08/1997
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight: 70.00
Age: 71Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

FLUANXOL (Suspected)			**Reason:** Otr spec symp psychopathol nec	
Injection	20.0 Milligram		1 time	Intramuscular
Batch:	**Started:** 08/02/1997			**Stopped:**
HALDOL DECANOATE (Suspected)			**Reason:** Otr spec symp psychopathol nec	
Injection	150.0 Milligram		1 time	Intramuscular
Batch:	**Started:** 13/02/1997			**Stopped:**
AURORIX (Suspected)			**Reason:** Depression	
Tablet	600.0 Milligram		Daily	Oral
Batch:	**Started:** 13/02/1997			**Stopped:**

Laboratory Investigations:

Additional Information:

Patient bedbound. prior to neuroleptic administration walking independently. patient discharged to nursing home 22/07/1997. developed pneumonia and died on 9/8/1997

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120219 **Seq:** 1 **Gender:** F
Reported: 22/08/1997 **Weight:** 63.00
Hospitalisation: **Age:** 51Y
Onset Date: 19/08/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion			
Flushing			
Hypertension			
Malaise			
Serotonin syndrome			
Tachycardia			
Tremor			

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
0.0
Batch: **Started:** **Stopped:** 17/08/1997

ZOLOFT (Interaction) **Reason:** Depression
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 18/08/1997 **Stopped:** 19/08/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 120492 **Seq:** 1 **Gender:** F
Reported: 03/09/1997 **Weight:** 56.00
Hospitalisation: **Age:** 78Y
Onset Date: 21/08/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Vasculitis		Rash on trunk and back Vasculitis petechial on upper legs.	

Medicine Details:

MAREVAN (Other drug) **Reason:**
Tablet 3.0 Milligram Daily Oral
Batch: **Started:** 11/08/1997 **Stopped:** CONTIN

LANOXIN (Other drug) **Reason:**
Oral application 250.0 Milligram Daily Oral
Batch: **Started:** 26/07/1997 **Stopped:** CONTIN

PARACETAMOL (Other drug) **Reason:**
Oral application 4.0 Gram Daily Oral
Batch: **Started:** 25/07/1997 **Stopped:** CONTIN

PREMARIN (Other drug) **Reason:**
625.0 Microgram Daily
Batch: **Started:** 26/07/1997 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergy to cortimoxazole.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120492 **Seq:** 1 **Gender:** F
Reported: 03/09/1997 **Weight:** 56.00
Hospitalisation: **Age:** 78Y
Onset Date: 21/08/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

TEMAZEPAM (Other drug)		**Reason:**		
	10.0 Milligram	Daily		
Batch:	**Started:** 04/08/1997		**Stopped:**	CONTIN
PERIACTIN (Other drug)		**Reason:**		
	4.0 Milligram	Daily		
Batch:	**Started:** 06/08/1997		**Stopped:**	CONTIN
AURORIX (Suspected)		**Reason:**	Depression	
Tablet	900.0 Milligram	Daily	Oral	
Batch:	**Started:** 30/07/1997		**Stopped:**	22/08/1997

Laboratory Investigations:

Additional Information:

Allergy to cortimoxazole.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120579 **Seq:** 1
Reported: 08/09/1997
Hospitalisation:
Onset Date: 30/07/1997
Outcome: Not yet recovered

Gender: F
Weight: 96.00
Age: 45Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dry mouth			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 08/05/1996 **Stopped:** 30/07/1997

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 30/07/1997 **Stopped:** 03/08/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Mild moderate xerostomia on 150mg bd, severe on 300mg bd.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120701 **Seq:** 1 **Gender:** F
Reported: 11/09/1997 **Weight:** 80.00
Hospitalisation: **Age:** 71Y
Onset Date: 06/09/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Abdominal pain Hyperhidrosis Nausea Tachycardia		Bowel evacuation	

Medicine Details:

THIAMINE HYDROCHLORIDE (Other drug) **Reason:**
0.0
Batch: **Started:** L TERM **Stopped:**

DICHLOTRIDE (Other drug) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

MONOPRIL (Other drug) **Reason:**
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

PARACETAMOL (Other drug) **Reason:**
Oral application 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 120701 **Seq:** 1 **Gender:** F
Reported: 11/09/1997 **Weight:** 80.00
Hospitalisation: **Age:** 71Y
Onset Date: 06/09/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

BECLOFORTE (Other drug)			**Reason:**		
Inhalation	4.0	Dose Unspecified	Daily		Inhalation
Batch:	**Started:**	L TERM		**Stopped:**	
VENTOLIN (Other drug)			**Reason:**		
Inhalation	12.0	Dose Unspecified	Daily		Inhalation
Batch:	**Started:**	L TERM		**Stopped:**	
VOLTAREN (Other drug)			**Reason:**		
	100.0	Milligram	Daily		
Batch:	**Started:**	L TERM		**Stopped:**	
EFEXOR (Suspected)			**Reason:**	Depression	
Tablet	37.5	Milligram	1 time		Oral
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 120701 **Seq:** 1 **Gender:** F
Reported: 11/09/1997 **Weight:** 80.00
Hospitalisation: **Age:** 71Y
Onset Date: 06/09/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120726 **Seq:** 1 **Gender:** F
Reported: 12/09/1997 **Weight:**
Hospitalisation: **Age:** 37
Onset Date: **DOB:** 01/11/1959
Outcome: Not yet recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Micturition disorder Constipation Dizziness Headache Urinary retention		Micturation delyed.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 21/05/1997 **Stopped:** 15/08/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient retried aurorix and again got urinary frequency and burning but delayed micturition.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 120801 **Seq:** 1 **Gender:** M
Reported: 15/09/1997 **Weight:** 73.00
Hospitalisation: **Age:** 52Y
Onset Date: 26/08/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pollakiuria			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 120911 **Seq:** 1 **Gender:** M
Reported: 17/09/1997 **Weight:**
Hospitalisation: **Age:** 72
Onset Date: **DOB:** 21/04/1925
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular			Ceased aurorix

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 11/03/1997

TEGRETOL (Suspected) **Reason:** Other&unspecified epilepsy
400.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 20/03/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121121 **Seq:** 1 **Gender:** F
Reported: 26/09/1997 **Weight:** 72.00
Hospitalisation: **Age:** 57
Onset Date: **DOB:** 16/05/1940
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Pruritus		Grouped plaques on arms, elbows, knees.	

Medicine Details:

ESTRADERM (Other drug)		**Reason:** Menopausal symptoms		
Cream	100.0 Milligram	Weekly	Transdermal	
Batch:	**Started:** 01/01/1990		**Stopped:**	CONTIN
MERSYNDOL (Other drug)		**Reason:** Migraine		
Tablet	0.0	As necessary	Oral	
Batch:	**Started:** 01/01/1991		**Stopped:**	CONTIN
CALTRATE (Other drug)		**Reason:** Osteoporosis		
Tablet	600.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/04/1997		**Stopped:**	CONTIN
IMURAN (Other drug)		**Reason:**		
Tablet	50.0 Milligram	Daily	Oral	
Batch:	**Started:** 01/05/1997		**Stopped:**	28/08/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Supposed systemic lupus erythematous diagnosed elsewhere. serology not supportive of diagnosis.
	Skin biopsy				Biopsy - lymphomatoid dermatitis 20/08/97. immunoflourescence

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 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 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: 121121 **Seq:** 1
Reported: 26/09/1997
Hospitalisation:
Onset Date:
Outcome: Unknown

Gender: F
Weight: 72.00
Age: 57
DOB: 16/05/1940
Causality: Causality possible

Reaction Detail

Medicine Details:

PREDNISONE (Other drug)			**Reason:**	
Tablet	0.0		Daily	Oral
Batch:	**Started:**		**Stopped:**	
AURORIX (Suspected)			**Reason:**	
Tablet	600.0	Milligram	Daily	Oral
Batch:	**Started:**	01/03/1997	**Stopped:**	27/08/1997

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 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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121323 **Seq:** 1 **Gender:** F
Reported: 03/10/1997 **Weight:**
Hospitalisation: **Age:** 36Y
Onset Date: 08/08/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blepharospasm		Flickering eyelids	
Gait disturbance		Unable to walk properly	
Diplopia			
Dizziness			Cogentin, ciproheptadine
Euphoric mood			
Malaise			
Nausea			
Serotonin syndrome			

Medicine Details:

ZOLOFT (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:** 06/08/1997

AURORIX (Interaction) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 07/08/1997 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121332 **Seq:** 1 **Gender:** F
Reported: 06/10/1997 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/12/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination			

Medicine Details:

ZOLOFT (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:** 04D

AURORIX (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient on zoloft for about 4 days after coming off aurorix gradually after 3 weeks and experienced hallucinations. - the date of onset is not accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121457 **Seq:** 1 **Gender:** F
Reported: 09/10/1997 **Weight:** 55.00
Hospitalisation: **Age:** 75Y
Onset Date: 24/09/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral		Bilateral oedema to mid-calf.	

Medicine Details:

CLOZAPINE (Other drug) **Reason:** Other&unspecified epilepsy
10.0 Milligram Daily
Batch: **Started:** 18/09/1997 **Stopped:** CONTIN

ASPIRIN (Other drug) **Reason:** Angina pectoris w/o hyperten
150.0 Milligram Daily
Batch: **Started:** 11/08/1997 **Stopped:** CONTIN

DILTIAZEM HYDROCHLORIDE (Other drug) **Reason:** Angina pectoris w/o hyperten
180.0 Milligram Daily
Batch: **Started:** 11/08/1997 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 10/09/1997 **Stopped:** 29/09/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 121586 **Seq:** 1 **Gender:** M
Reported: 14/10/1997 **Weight:** 80.00
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Increased appetite Nausea Dizziness		Increased appetite for lollies - occasionally Queasy occasionally.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 01/09/1997 **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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121621 **Seq:** 1 **Gender:** F
Reported: 16/10/1997 **Weight:** 54.00
Hospitalisation: **Age:** 24Y
Onset Date: 21/09/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Hypertension			

Medicine Details:

SUDAFED (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:** Depression
450.0 Milligram Daily Oral
Batch: **Started:** 18/09/1997 **Stopped:**

RHINOCORT (Other drug) **Reason:** Chronic nasopharyngitis
0.0
Batch: **Started:** 18/02/1997 **Stopped:**

ZOLADEX IMPLANT (Other drug) **Reason:** Uterine endometriosis
0.0 Monthly
Batch: **Started:** 05/09/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/12/2002	Other data				Bp : 188/110, 118/115, 182/108, 148/92, 132/86 time :__1335 1340 1345 1350

Additional Information:

History of allergy to ees - rash, pethidine, omnopon - respiratory problems, metoclopramide - occulogyric crisis. bp time 188/110 - 13.35 188/115 - . 182/108 - 13.40 148/92 - 13.45 132/86 - 13.50

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121674 **Seq:** 1 **Gender:** F
Reported: 20/10/1997 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 05/08/1997 **Stopped:** 12/08/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121759 **Seq:** 1 **Gender:** F
Reported: 22/10/1997 **Weight:** 80.00
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Gravitational oedema			

Medicine Details:

OROXINE (Suspected) **Reason:**
100.0 Microgram Daily
Batch: **Started:** 01/01/1984 **Stopped:** 31/01/1997

AURORIX (Suspected) **Reason:**
250.0 Milligram Daily
Batch: **Started:** 01/01/1994 **Stopped:** 28/10/1997

AMOXIL (Suspected) **Reason:**
250.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:** 28/10/1997

MODURETIC (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/02/1997 **Stopped:** 28/10/1997

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. 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 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 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

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 121759 **Seq:** 1 **Gender:** F
Reported: 22/10/1997 **Weight:** 80.00
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detai

Medicine Details:

TAZAC (Suspected)		**Reason:**	
	2.0 Dose Unspecified Daily		
Batch:	**Started:** 01/02/1997	**Stopped:**	28/10/1997
TAMOXIFEN CITRATE (Suspected)		**Reason:**	
	20.0 Milligram Daily		
Batch:	**Started:** 01/06/1997	**Stopped:**	28/10/1997

Laboratory Investigations:

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 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 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

Report Detail

Case Number: 121885 **Seq:** 1 **Gender:** M
Reported: 24/10/1997 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: 15/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Vomiting			Stemetil

Medicine Details:

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

ASCORBIC ACID (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

OXYBUTYNIN HYDROCHLORIDE (Other drug) **Reason:**
Tablet 7.5 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
300.0 Milligram 1 time
Batch: **Started:** 15/10/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergy to penicillin, sulphur. reaction to tryptanol - rash

Report Detail

Case Number: 121966 **Seq:** 1 **Gender:** F
Reported: 28/10/1997 **Weight:**
Hospitalisation: **Age:** 32Y
Onset Date: 14/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pain Headache Nausea	Severe	Neck stiffness, spasm.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 09/10/1997 **Stopped:** 20/10/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 122107 **Seq:** 1
Reported: 03/11/1997
Hospitalisation:
Onset Date: 24/10/1997
Outcome: Not yet recovered

Gender: M
Weight: 90.00
Age: 58Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flatulence		Feeling of fullness/discomfort	

Medicine Details:

ZANTAC (Other drug) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/10/1997 **Stopped:** 25/10/1997

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 122114 **Seq:** 1 **Gender:** F
Reported: 04/11/1997 **Weight:** 60.00
Hospitalisation: **Age:** 53Y
Onset Date: 24/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			

Medicine Details:

OGEN (Other drug) **Reason:** Menopausal symptoms
1.2 Milligram Daily
Batch: **Started:** 24/07/1996 **Stopped:** CONTIN

PROVERA (Other drug) **Reason:** Menopausal symptoms
5.0 Milligram Daily
Batch: **Started:** 24/07/1997 **Stopped:** CONTIN

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

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 27/09/1996 **Stopped:** 23/10/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 122232 **Seq:** 1 **Gender:** M
Reported: 10/11/1997 **Weight:**
Hospitalisation: **Age:** 54Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hallucination		Visual and auditory hallucinations	

Medicine Details:

SEREPAX (Other drug) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 21/08/1997 **Stopped:** 15/09/1997

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 122238 **Seq:** 1 **Gender:** F
Reported: 10/11/1997 **Weight:** 50.00
Hospitalisation: **Age:** 49Y
Onset Date: 01/11/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Anorexia			
Anxiety			
Insomnia			
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/11/1997 **Stopped:** 06/11/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 122661 **Seq:** 1 **Gender:** F
Reported: 21/11/1997 **Weight:** 65.00
Hospitalisation: **Age:** 80Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Focal convulsiva episodes leading to	Sodium valproate

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 122753 **Seq:** 1 **Gender:** U
Reported: 26/11/1997 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle twitching		Muscle twitching	Aurorix ceased and zoloft commenced

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 122777 **Seq:** 1 **Gender:** F
Reported: 25/11/1997 **Weight:** 47.00
Hospitalisation: **Age:** 32Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Nausea			Ceased aurorix

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 09/10/1997 **Stopped:** 20/10/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123167 **Seq:** 1 **Gender:** M
Reported: 09/12/1997 **Weight:**
Hospitalisation: **Age:** 25
Onset Date: 31/10/1997 **DOB:** 05/08/1972
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Hives/allergic rash	Aurorix discontinued

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 0.0 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123265 **Seq:** 1 **Gender:** M
Reported: 11/12/1997 **Weight:** 79.00
Hospitalisation: **Age:** 21Y
Onset Date: 02/12/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Severe		

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/12/1997 **Stopped:** 05/12/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123474 **Seq:** 1 **Gender:** M
Reported: 17/12/1997 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 31/12/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**

0.0

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123783 **Seq:** 1 **Gender:** M
Reported: 07/01/1998 **Weight:** 80.00
Hospitalisation: **Age:** 35Y
Onset Date: 20/12/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 16/12/1997 **Stopped:** 24/12/1997

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123792 **Seq:** 1 **Gender:** F
Reported: 07/01/1997 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/12/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Libido decreased			

Medicine Details:

AURORIX (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: 123803 **Seq:** 1 **Gender:** F
Reported: 06/01/1998 **Weight:** 65.00
Hospitalisation: **Age:** 72Y
Onset Date: 01/12/1997 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Speech disorder Dysarthria		Slurred speech	Aurorix discontinued

Medicine Details:

PREPULSID (Other drug) **Reason:** Other diseases of esophagus
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

ROHYPNOL (Other drug) **Reason:** Specific disorders of sleep
1.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:**
450.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Slurred speech within 4 hrs of commencing aurorix. disappears by 24hrs after dose, returns after re-dosing. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 123834 **Seq:** 1 **Gender:** M
Reported: 09/01/1998 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: 31/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dyspnoea			Aurorix discontinued

Medicine Details:

DIAMICRON (Other drug) **Reason:**
Tablet 4.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

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

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

DIAFORMIN (Other drug) **Reason:**
500.0 Milligram Daily
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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 123834 **Seq:** 1 **Gender:** M
Reported: 09/01/1998 **Weight:**
Hospitalisation: **Age:** 65Y
Onset Date: 31/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis
300.0 Milligram Daily
Batch: **Started:** 13/10/1997 **Stopped:** 28/12/1997

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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124090 **Seq:** 1 **Gender:** U
Reported: 16/01/1998 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lipodystrophy acquired			

Medicine Details:

CRIXIVAN (Suspected) **Reason:** Hiv infection
Capsule 2.4 Gram Daily Oral
Batch: **Started:** 14/11/1996 **Stopped:** CONTIN

ZIDOVUDINE (Suspected) **Reason:** Hiv infection
500.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

3TC (Suspected) **Reason:** Hiv infection
300.0 Milligram Daily
Batch: **Started:** 01/01/1995 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/01/1996 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124090 **Seq:** 1
Reported: 16/01/1998
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: U
Weight:
Age:
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ACICLOVIR (Suspected) **Reason:** Hiv infection

800.0 Milligram Daily

Batch: **Started:** 01/01/1995 **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124247 **Seq:** 1
Reported: 19/01/1998
Hospitalisation: Admitted to hospital
Onset Date: 29/01/1997
Outcome: Unknown

Gender: F
Weight:
Age: 79Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression			

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
0.0
Batch: **Started:** **Stopped:**

PREDSOL (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

IOPIDINE (Interaction) **Reason:** Unspecified glaucoma
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124306 **Seq:** 1 **Gender:** F
Reported: 21/01/1998 **Weight:**
Hospitalisation: **Age:** 67
Onset Date: 21/10/1997 **DOB:** 16/05/1930
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Eosinophilia		Widespread rash erythrodermic	Betnovate ointment bd, sorbolene cream, sm-33 gel, prednisolone 25mg for 6 day then 20mg, hydrocortisone oin bd, liquid paraffin, wet dressing

Medicine Details:

ATENOLOL (Suspected) **Reason:**
25.0 Milligram Alternating-other
Batch: **Started:** L TERM **Stopped:**

RANITIDINE (Suspected) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

QUININE (Suspected) **Reason:**
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

SIMVASTATIN (Suspected) **Reason:**
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:** 19/11/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Skin biopsy c/w drug hypersensitivity.

Additional Information:

History of allergy to penicillin and perfumes.

Report Detail

Case Number: 124306 **Seq:** 1
Reported: 21/01/1998
Hospitalisation:
Onset Date: 21/10/1997
Outcome: Not yet recovered

Gender: F
Weight:
Age: 67
DOB: 16/05/1930
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**		
Tablet	600.0	Milligram	Daily		Oral
Batch:	**Started:**		L TERM	**Stopped:**	
F.G.F. (Suspected)			**Reason:**		
Oral application	1.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**		L TERM	**Stopped:**	
CALTRATE (Suspected)			**Reason:**		
Tablet	3.0	Dose Unspecified	Daily		Oral
Batch:	**Started:**		L TERM	**Stopped:**	
CALCITRIOL (Suspected)			**Reason:**		
	250.0	Microgram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	

Laboratory Investigations:

Additional Information:

History of allergy to penicillin and perfumes.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124306 **Seq:** 1
Reported: 21/01/1998
Hospitalisation:
Onset Date: 21/10/1997
Outcome: Not yet recovered

Gender: F
Weight:
Age: 67
DOB: 16/05/1930
Causality: Causality possible

Reaction Detail

Medicine Details:

AMLODIPINE (Suspected)			**Reason:**		
	10.0	Milligram	Daily		
Batch:	**Started:**		L TERM	**Stopped:**	19/11/1997
ACICLOVIR (Suspected)			**Reason:**		
Tablet	400.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
THYROXINE SODIUM (Suspected)			**Reason:**		
	25.0	Microgram	Alternate days		
Batch:	**Started:**			**Stopped:**	
EPOETIN ALFA (Suspected)			**Reason:**		
	4.0	Thousand Internat	Alternate days		
Batch:	**Started:**		L TERM	**Stopped:**	05/11/1997

Laboratory Investigations:

Additional Information:

History of allergy to penicillin and perfumes.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124410 **Seq:** 1 **Gender:** M
Reported: 28/01/1998 **Weight:** 90.00
Hospitalisation: **Age:** 47Y
Onset Date: 08/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 08/01/1998 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 124411 **Seq:** 1 **Gender:** F
Reported: 28/01/1998 **Weight:** 52.00
Hospitalisation: **Age:** 22Y
Onset Date: 09/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Galactorrhoea		Bilateral watery discharge from both	Aurorix ceased
Breast pain		Breast soreness	
Hyperprolactinaemia		Prolactin level increased.	

Medicine Details:

MAXOLON (Other drug) **Reason:** Nausea and vomiting
10.0 Milligram Daily
Batch: **Started:** 01/11/1997 **Stopped:**

PROZAC (Other drug) **Reason:** Depression
40.0 Milligram Daily
Batch: **Started:** 06/10/1997 **Stopped:** 01/01/1998

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 02/01/1998 **Stopped:** 16/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prolactin	(50-500)	12/01/1998	680	
	Prolactin	(50-500)	20/01/1998	434	

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124579 **Seq:** 1 **Gender:** F
Reported: 02/02/1998 **Weight:**
Hospitalisation: **Age:** 36
Onset Date: 15/07/1997 **DOB:** 23/11/1960
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast pain Hyperprolactinaemia			

Medicine Details:

CYCLOPHOSPHAMIDE (Suspected) **Reason:** Systemic lupus erythematosus
0.0
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 01/10/1996 **Stopped:** CONTIN

RISPERDAL (Suspected) **Reason:** Unspecified psychosis
Tablet 500.0 Microgram Daily Oral
Batch: **Started:** 01/10/1996 **Stopped:** CONTIN

PREDNISONE (Suspected) **Reason:** Systemic lupus erythematosus
50.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prolactin	(46-622)	15/07/1997	838	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124608 **Seq:** 1 **Gender:** F
Reported: 03/02/1998 **Weight:** 64.00
Hospitalisation: **Age:** 64Y
Onset Date: 07/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Aphasia		Confusion/expressive dyphasia	

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occured each day approx. 30 min after dose. see original report for other drugs - dilantin, tegretol, renitec, natrilix, pepcidine

Report Detail

Case Number: 124630 **Seq:** 1 **Gender:** F
Reported: 03/02/1998 **Weight:** 71.00
Hospitalisation: **Age:** 63Y
Onset Date: 07/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sensory disturbance Hypoaesthesia Neuropathy peripheral Pain		Decreased sensation in both legs.	 Ceased cozaar

Medicine Details:

PREPULSID FORTE (Other drug) **Reason:** Other diseases of esophagus
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

AMFAMOX (Other drug) **Reason:** Other diseases of esophagus
80.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ESTRADERM (Other drug) **Reason:** Menopausal symptoms
Cream 50.0 Microgram Daily Transdermal
Batch: **Started:** L TERM **Stopped:**

ASPIRIN (Other drug) **Reason:** Angina pectoris w/o hyperten
0.5 Dose Unspecified Daily
Batch: **Started:** 20/06/1997 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Decreased pinprick in stocking distribution to proximal lower leg.

Additional Information:

Rapid progression of peripheral neuropathy since on cozaar diabetes well controlled.

Report Detail

Case Number: 124630 **Seq:** 1 **Gender:** F
Reported: 03/02/1998 **Weight:** 71.00
Hospitalisation: **Age:** 63Y
Onset Date: 07/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

MINIDIAB (Other drug)		**Reason:**		
	2.5 Milligram	Daily		
Batch:	**Started:** 18/06/1997		**Stopped:**	
COZAAR (Suspected)		**Reason:**	Cardiomyopathy	
Tablet	50.0 Milligram	Daily		Oral
Batch:	**Started:** 15/12/1997		**Stopped:**	28/01/1998
AURORIX (Suspected)		**Reason:**		
	450.0 Milligram	Daily		
Batch:	**Started:** 17/11/1997		**Stopped:**	28/01/1998

Laboratory Investigations:

Additional Information:

Rapid progression of peripheral neuropathy since on cozaar diabetes well controlled.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 124941 **Seq:** 1 **Gender:** F
Reported: 13/02/1998 **Weight:** 57.00
Hospitalisation: **Age:** 36Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bradycardia Orthostatic hypotension			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 19/06/1997 **Stopped:** 11/01/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125095 **Seq:** 1 **Gender:** M
Reported: 18/02/1998 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular	Required Specialist Consultation		Diprosone topically; aurovix

Medicine Details:

THIORIDAZINE HYDROCHLORIDE (Other drug) **Reason:**
50.0 Milligram As necessary
Batch: **Started:** **Stopped:** CONTIN

VERAPAMIL HYDROCHLORIDE (Other drug) **Reason:**
Oral application 160.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

OXAZEPAM (Other drug) **Reason:**
30.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
900.0 Milligram Daily
Batch: **Started:** 13/01/1998 **Stopped:** 29/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

P/h allergy to penicillin.

Report Detail

Case Number: 125146 **Seq:** 1
Reported: 13/02/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight:
Age: 24Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor Convulsion Agitation Coordination abnormal Flushing Insomnia Serotonin syndrome Urinary retention Vomiting	Severe	Legs shaking violently and chattering of [illegible] Myoclonic jerking	

Medicine Details:

CLOMIPRAMINE HYDROCHLORIDE (Interaction) **Reason:**
50.0 Milligram 1 time
Batch: **Started:** **Stopped:**

FLUNITRAZEPAM (Interaction) **Reason:**
2.0 Milligram 1 time
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:**
750.0 Milligram Daily
Batch: **Started:** **Stopped:** 17D

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

International clinical psychopharmacology 1997, 12, 339-342, v12.

Report Detail

Case Number: 125259 **Seq:** 1 **Gender:** F
Reported: 24/02/1998 **Weight:**
Hospitalisation: **Age:** 92Y
Onset Date: 26/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SLOW-K (Other drug) **Reason:**
Tablet 1.2 Gram Daily Oral
Batch: **Started:** **Stopped:**

SOLPRIN (Other drug) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

PANAMAX (Other drug) **Reason:** Pain
Tablet 4.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium		04/02/1998	4.4	
	Potassium		10/02/1998	4.0	
	Sodium		26/01/1998	120	
	Sodium		01/02/1998	138	
	Sodium		04/02/1998	135	
	Sodium		10/02/1998	130	

Additional Information:

Patient noted to have siadh in past medical history.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125259 **Seq:** 1 **Gender:** F
Reported: 24/02/1998 **Weight:**
Hospitalisation: **Age:** 92Y
Onset Date: 26/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

RIVOTRIL (Suspected)			**Reason:**		
Tablet	500.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	28/01/1998
LASIX (Suspected)			**Reason:**	Essential benign hypertension	
Tablet	40.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	30/01/1998
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	900.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	28/01/1998

Laboratory Investigations:

Additional Information:

Patient noted to have siadh in past medical history.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125288 **Seq:** 1 **Gender:** F
Reported: 25/02/1998 **Weight:**
Hospitalisation: **Age:** 84
Onset Date: 26/01/1998 **DOB:** 21/02/1913
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Aurorix ceased

Medicine Details:

COZAAR (Other drug) **Reason:** Essential benign hypertension
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 15/01/1998 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 22/01/1998 **Stopped:** 03/02/1998

CARDOL (Suspected) **Reason:** Otr&nos disord of heart rhythm
Tablet 160.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 04/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium		26/01/1998	-	
	Potassium		05/02/1998	-	
	Potassium		09/02/1998	4.1	
	Potassium		06/02/1998	3.8	
	Sodium	133-145_MMOL/	26/01/1998	123	
	Sodium	133-145_MMOL/	05/02/1998	128	
	Sodium	133-145_MMOL/	06/02/1998	130	

Additional Information:

Patient noted to have siadh in post medical history.

Report Detail

Case Number: 125288 **Seq:** 1 **Gender:** F
Reported: 25/02/1998 **Weight:**
Hospitalisation: **Age:** 84
Onset Date: 26/01/1998 **DOB:** 21/02/1913
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

COZAAR (Other drug)			**Reason:**	Essential benign hypertension
Tablet	50.0	Milligram	Daily	Oral
Batch:	**Started:**	15/01/1998	**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression
Tablet	300.0	Milligram	Daily	Oral
Batch:	**Started:**	22/01/1998	**Stopped:**	03/02/1998
CARDOL (Suspected)			**Reason:**	Otr&nos disord of heart rhythm
Tablet	160.0	Milligram	Daily	Oral
Batch:	**Started:**		**Stopped:**	04/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium	133-145_MMOL/	09/02/1998	138	

Additional Information:

Patient noted to have siadh in post medical history.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125289 **Seq:** 1 **Gender:** F
Reported: 25/02/1998 **Weight:**
Hospitalisation: **Age:** 91
Onset Date: 26/01/1997 **DOB:** 18/06/1905
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			

Medicine Details:

RENITEC (Other drug) **Reason:** Essential benign hypertension
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SLOW-K (Other drug) **Reason:**
Tablet 1.2 Gram Daily Oral
Batch: **Started:** **Stopped:**

SOLPRIN (Other drug) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 28/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium	3.5-5.5 MMOL/L	04/02/1998	4.4	
	Potassium	3.5-5.5 MMOL/L	05/02/1998	4.4	
	Potassium	3.5-5.5 MMOL/L	07/02/1998	4.9	
	Potassium	3.5-5.5 MMOL/L	11/02/1998	4.0	
	Potassium	3.5-5.5 MMOL/L	10/02/1998	4.0	
	Sodium	135-145 MMOL/L	26/01/1998	120	
	Sodium	135-145 MMOL/L	01/02/1998	138	

Additional Information:

Patient noted to have siadh in past medical history

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125289 **Seq:** 1 **Gender:** F
Reported: 25/02/1998 **Weight:**
Hospitalisation: **Age:** 91
Onset Date: 26/01/1997 **DOB:** 18/06/1905
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

RIVOTRIL (Suspected)			**Reason:**		
Tablet	500.0	Microgram	Daily		Oral
Batch:	**Started:**			**Stopped:**	28/01/1998
LASIX (Suspected)			**Reason:**	Essential benign hypertension	
Tablet	40.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	30/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium	135-145 MMOL/L	04/02/1998	135	
	Sodium	135-145 MMOL/L	05/02/1998	135	
	Sodium	135-145 MMOL/L	07/02/1998	129	
	Sodium	135-145 MMOL/L	10/02/1998	130	
	Sodium	135-145 MMOL/L	11/02/1998	129	

Additional Information:

Patient noted to have siadh in past medical history

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125466 **Seq:** 1
Reported: 03/03/1998
Hospitalisation:
Onset Date: 30/01/1998
Outcome: Not yet recovered

Gender: F
Weight: 99.00
Age: 47Y
DOB:
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia		Thinning of hair of whole head. loss 1/3 of	Ceased aurorix.

Medicine Details:

CAPADEX (Other drug) **Reason:** Pain
8.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1991 **Stopped:** 07/02/1998

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 09/01/1998 **Stopped:** 01/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous exposure to aurorix in september 1996 to may 1997, symptoms started to occurred in this period. generalised thinning of hair of whole head. loss of 1/3 of previous hair mass. the dosage start date is not accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 125491 **Seq:** 1 **Gender:** F
Reported: 03/03/1998 **Weight:**
Hospitalisation: **Age:** 66
Onset Date: 09/02/1998 **DOB:** 17/10/1931
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare	Severe		Ceased moclobemide

Medicine Details:

IMDUR (Other drug) **Reason:**
60.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

MIXTARD 30/70 (Other drug) **Reason:** Diabetes mellitus
28.0 Unit Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

OMEPRAZOLE (Other drug) **Reason:** Ulcer of duodenum w/o ment perf
Capsule 40.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

MAGMIN (Other drug) **Reason:**
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 125491 **Seq:** 1 **Gender:** F
Reported: 03/03/1998 **Weight:**
Hospitalisation: **Age:** 66
Onset Date: 09/02/1998 **DOB:** 17/10/1931
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

PREDNISOLONE (Other drug)		**Reason:**	Otr mus rheum,fibrosit&myalgia
5.0 Milligram	Daily		
Batch:	**Started:** L TERM	**Stopped:**	
MOCLOBEMIDE (Suspected)		**Reason:**	Depression
150.0 Milligram	Daily		
Batch:	**Started:** 09/02/1998	**Stopped:**	16/02/1998

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125593 **Seq:** 1 **Gender:** M
Reported: 06/03/1998 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Agitation		Irritability, short tempered,easily	Gradually taken off aurorix over 3 weeks
Aggression		Short tempered, not coping well with stress	
Insomnia			

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SANDOMIGRAN (Interaction) **Reason:** Other prophylactic procedures
Tablet 3.0 Dose Unspecified Daily Oral
Batch: **Started:** **Stopped:**

KALMA (Other drug) **Reason:** Anxiety neurosis
Tablet 1.0 Milligram As necessary Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 125773 **Seq:** 1 **Gender:** F
Reported: 13/02/1998 **Weight:** 90.00
Hospitalisation: **Age:** 53Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bronchospasm		Wheezy	

Medicine Details:

STEMETIL (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

ESTRADERM (Other drug) **Reason:** Menopausal symptoms
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

FRUSEMIDE (Other drug) **Reason:**
50.0 Milligram Daily
Batch: **Started:** **Stopped:**

OROXINE (Other drug) **Reason:** Myxedema
0.0 Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Certain - rechallenge. increased aurorix from 300mg to 600 mg and became wheezy. delayed dose and settled. took 600mg and became wheezy again. same next day. today no aurorix and only slightly short of breath. audible wheeze over phone.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125773 **Seq:** 1 **Gender:** F
Reported: 13/02/1998 **Weight:** 90.00
Hospitalisation: **Age:** 53Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Medicine Details:

PROVERA (Other drug)			**Reason:**	
	0.0		Daily	
Batch:	**Started:**			**Stopped:**
SLOW-K (Other drug)			**Reason:**	
	0.0		Daily	
Batch:	**Started:**			**Stopped:**
SULINDAC (Other drug)			**Reason:**	
	200.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
AURORIX (Suspected)			**Reason:**	
Tablet	600.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Certain - rechallenge. increased aurorix from 300mg to 600 mg and became wheezy. delayed dose and settled. took 600mg and became wheezy again. same next day. today no aurorix and only slightly short of breath. audible wheeze over phone.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125823 **Seq:** 1 **Gender:** F
Reported: 17/03/1998 **Weight:** 60.00
Hospitalisation: **Age:** 68Y
Onset Date: 12/03/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache		Head felt heavy and swollen	
Agitation		Panic attack	
Paraesthesia		Skin felt crawly.	
Agitation			
Confusional state			
Insomnia			
Palpitations			

Medicine Details:

PANAMAX (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

BECONASE (Other drug) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:**

PRAVACHOL (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

OROXINE (Other drug) **Reason:**
100.0 Microgram 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125823 **Seq:** 1 **Gender:** F
Reported: 17/03/1998 **Weight:** 60.00
Hospitalisation: **Age:** 68Y
Onset Date: 12/03/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ALODORM (Other drug)			**Reason:**	
	5.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
AURORIX (Suspected)			**Reason:**	
Tablet	300.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125898 **Seq:** 1 **Gender:** M
Reported: 18/03/1998 **Weight:** 100.00
Hospitalisation: **Age:** 46Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sexual dysfunction	Severe	Severe sexual dysfunction	

Medicine Details:

VENTOLIN (Other drug) **Reason:** Asthma
Inhalation 0.0 Daily Inhalation
Batch: **Started:** **Stopped:**

BECOTIDE (Other drug) **Reason:** Asthma
Inhalation 0.0 Daily Inhalation
Batch: **Started:** **Stopped:**

SERZONE (Other drug) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 13/08/1997 **Stopped:** 28/09/1997

AURORIX (Suspected) **Reason:**
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** 17/11/1997 **Stopped:** CONTIN

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: 125919 **Seq:** 1 **Gender:** F
Reported: 19/03/1998 **Weight:** 79.00
Hospitalisation: **Age:** 49Y
Onset Date: 05/03/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Galactorrhoea			

Medicine Details:

CALCIUM CARBONATE (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** 01/01/1998 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/01/1994 **Stopped:**

COVERSYL (Suspected) **Reason:**
2.0 Milligram Daily
Batch: **Started:** 01/01/1997 **Stopped:**

THYROXINE SODIUM (Suspected) **Reason:**
0.2 Milligram Daily
Batch: **Started:** 01/01/1998 **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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125919 **Seq:** 1
Reported: 19/03/1998
Hospitalisation:
Onset Date: 05/03/1998
Outcome: Not yet recovered

Gender: F
Weight: 79.00
Age: 49Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

MAGNESIUM SULPHATE (Suspected)		**Reason:**	
Tablet	0.0	Daily	Oral
Batch:	**Started:** 01/01/1998		**Stopped:**

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 125933 **Seq:** 1 **Gender:** F
Reported: 19/03/1998 **Weight:** 70.00
Hospitalisation: **Age:** 61Y
Onset Date: 07/01/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dehydration Diarrhoea	Severe		Ceased aurorix

Medicine Details:

ROHYPNOL (Other drug) **Reason:** Specific disorders of sleep
0.0
Batch: **Started:** 01/02/1996 **Stopped:**

CARDIZEM CD (Other drug) **Reason:** Essential benign hypertension
Capsule 180.0 Milligram Daily Oral
Batch: **Started:** 01/03/1995 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 05/12/1997 **Stopped:** 09/01/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pt has allergies to lithium, keflex, zoloft and efexor 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: 126022 **Seq:** 1 **Gender:** F
Reported: 23/03/1998 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 27/02/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haematuria Prothrombin level decreased		Spontaneous bleeds	 Ceased warfarin

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 24/02/1998 **Stopped:**

WARFARIN SODIUM (Interaction) **Reason:**
1.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 27/02/1998

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

CARDIZEM (Other drug) **Reason:** Essential benign hypertension
180.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	International				Inr dramatically increased to 140.

Additional Information:

Report Detail

Case Number: 126022 **Seq:** 1 **Gender:** F
Reported: 23/03/1998 **Weight:**
Hospitalisation: **Age:** 85Y
Onset Date: 27/02/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

GLYCERYL TRINITRATE (Other drug)				**Reason:**		
Cream	1.0	Dose Unspecified	Daily		Transdermal	
Batch:	**Started:**		L TERM	**Stopped:**		
NEULACTIL (Other drug)				**Reason:**	Otr spec symp psychopathol nec	
	5.0	Milligram	Daily			
Batch:	**Started:**	03/02/1998		**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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126088 **Seq:** 1
Reported: 23/03/1998
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nausea Pyrexia Serotonin syndrome Tremor			Patient is stabilised on zoloft

Medicine Details:

ZOLOFT (Interaction)			**Reason:**		
Tablet	50.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	
AURORIX (Interaction)			**Reason:**		
Tablet	600.0	Milligram	Daily	Oral	
Batch:	**Started:**			**Stopped:**	3M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Zoloft commenced after 4 day washout.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126101 **Seq:** 1 **Gender:** F
Reported: 23/03/1998 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 11/12/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia		Sharp shooting pains from back of tongue	

Medicine Details:

PROVERA (Other drug) **Reason:** Menopausal symptoms
0.0
Batch: **Started:** L TERM **Stopped:** CONTIN

ENALAPRIL MALEATE (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

LASIX (Other drug) **Reason:** Essential benign hypertension
20.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

FEMTRAN (Other drug) **Reason:** Menopausal symptoms
Cream 0.0 Daily Topical
Batch: **Started:** L TERM **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126101 **Seq:** 1 **Gender:** F
Reported: 23/03/1998 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 11/12/1997 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** **Stopped:** 23/10/1997

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 126253 **Seq:** 1 **Gender:** F
Reported: 30/03/1998 **Weight:** 57.00
Hospitalisation: **Age:** 82Y
Onset Date: 14/03/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Severe		Ceased aurorix

Medicine Details:

AURORIX (Suspected) **Reason:**

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 13/03/1998 **Stopped:** 20/03/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126308 **Seq:** 1 **Gender:** F
Reported: 31/03/1998 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: 15/02/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria	Severe	Rash on ankles, legs & progressed to trunk	Aurorix ceased

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 14/02/1998 **Stopped:** 15/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Histology				Biopsy showed perivascular infiltration of lymphocytes and moderate eosinophils and occassional neutrophils.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126764 **Seq:** 1 **Gender:** M
Reported: 17/04/1998 **Weight:**
Hospitalisation: **Age:** 38
Onset Date: **DOB:** 12/02/1960
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thinking abnormal Disturbance in attention Hallucination Tremor		Low mental state. Unable to concentrate	 Stopped all medication. diazepam 5-10mg prn oral given

Medicine Details:

AROPAX (Suspected)			**Reason:**	
	20.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	1W
AURORIX (Suspected)			**Reason:**	
	300.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	1W
MELLERIL (Suspected)			**Reason:**	
	100.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	1W
NEULACTIL (Suspected)			**Reason:**	
	30.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	1W

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126785 **Seq:** 1 **Gender:** F
Reported: 20/04/1998 **Weight:** 50.00
Hospitalisation: **Age:** 79Y
Onset Date: 15/12/1997 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			

Medicine Details:

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

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

AMFAMOX (Other drug) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

SEREPAX (Other drug) **Reason:**
30.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126785 **Seq:** 1
Reported: 20/04/1998
Hospitalisation:
Onset Date: 15/12/1997
Outcome: Not yet recovered

Gender: F
Weight: 50.00
Age: 79Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 15/12/1997 **Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126844 **Seq:** 1 **Gender:** F
Reported: 22/04/1998 **Weight:** 60.00
Hospitalisation: **Age:** 71Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysarthria		Slurred speech	Ceased aurorix

Medicine Details:

GAVISCON (Other drug) **Reason:**
10.0 Millilitre As necessary
Batch: **Started:** **Stopped:**

PREPULSID (Other drug) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

LIPEX (Other drug) **Reason:**
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

HISMANAL (Other drug) **Reason:**
1.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Second attempt with this drug produced same result.

Report Detail

Case Number: 126844 **Seq:** 1 **Gender:** F
Reported: 22/04/1998 **Weight:** 60.00
Hospitalisation: **Age:** 71Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Medicine Details:

FLIXOTIDE (Other drug)			**Reason:**	
	1.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
AURORIX (Suspected)			**Reason:**	
	450.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Second attempt with this drug produced same result.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 126985 **Seq:** 1 **Gender:** F
Reported: 29/04/1998 **Weight:** 50.00
Hospitalisation: **Age:** 32Y
Onset Date: 05/04/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flatulence Rectal tenesmus		Feeling of fullness after 8 hrs after eating. Sensation of wanting to go to toilet all the	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 26/03/1998 **Stopped:** 10/04/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 127189 **Seq:** 1 **Gender:** F
Reported: 06/05/1998 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Death **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Agitation Sudden death		Very agitated	Zoloft ceased, aurorix commenced. unknown whether zoloft was gradually reduced or whether aurorix was commenced.

Medicine Details:

ZOLOFT (Interaction) **Reason:** Depression
200.0 Milligram Daily
Batch: **Started:** **Stopped:** 4W

AURORIX (Interaction) **Reason:** Depression
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Dose was titrated very quickly from 50mg to 200 mg. reaction occurred at this dosage.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 127487 **Seq:** 1 **Gender:** M
Reported: 15/05/1998 **Weight:** 70.00
Hospitalisation: **Age:** 30Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tachycardia		Heart racing	
Thinking abnormal		Racing thoughts	
Dysarthria		Slurring speach	
Abdominal pain			
Confusional state			
Diarrhoea			
Hyperhidrosis			
Myalgia			
Nervousness			
Serotonin syndrome			
Tremor			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 01/04/1998 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

13th reaction - emotional lability. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128018 **Seq:** 1
Reported: 01/06/1998
Hospitalisation:
Onset Date: 01/05/1998
Outcome: Not yet recovered

Gender: M
Weight: 80.00
Age: 51Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 12/09/1997 **Stopped:** CONTIN

MOGADON (Suspected) **Reason:**
5.0 Milligram Daily
Batch: **Started:** 01/01/1995 **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: 128026 **Seq:** 1
Reported: 01/06/1998
Hospitalisation: Admitted to hospital
Onset Date: 23/05/1998
Outcome: Recovered

Gender: F
Weight: 60.00
Age: 50Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing Orthostatic hypotension Somnolence Visual disturbance			

Medicine Details:

EFEXOR (Interaction) **Reason:**

Tablet 2.0 Dose Unspecified 1 time Oral

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

AURORIX (Interaction) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 23/05/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128170 **Seq:** 1
Reported: 09/06/1998
Hospitalisation:
Onset Date: 18/05/1998
Outcome: Not yet recovered

Gender: F
Weight: 102.00
Age: 36Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Osteoarthritis		Ache, heaviness in knees, elbows wrists & Stiffness in knees, elbows, wrists & ankles	Aurorix dose decreased to

Medicine Details:

XANAX (Other drug) **Reason:** Anxiety neurosis
1.5 Milligram Daily
Batch: **Started:** 23/01/1998 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Anxiety neurosis
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** 30/04/1998 **Stopped:** 20/05/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Multiple adverse reactions to most available antidepressants.

Report Detail

Case Number: 128229 **Seq:** 1 **Gender:** F
Reported: 09/06/1998 **Weight:** 47.00
Hospitalisation: **Age:** 29Y
Onset Date: 23/05/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Agitation Diarrhoea Nausea		Inability to sleep	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 16/05/1998 **Stopped:** 31/05/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 128253 **Seq:** 1 **Gender:** F
Reported: 10/06/1998 **Weight:** 56.00
Hospitalisation: **Age:** 34Y
Onset Date: 07/05/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia		Sleep disturbance	Aurorix reduced
Hyperhidrosis		Sweats	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 06/05/1998 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128320 **Seq:** 1 **Gender:** M
Reported: 12/06/1998 **Weight:** 65.00
Hospitalisation: **Age:** 30Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bruxism		Clenching jaws.	
Hyperhidrosis		Drenching sweats.	
Anxiety			
Insomnia			
Palpitations			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Total
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128358 **Seq:** 1 **Gender:** F
Reported: 15/06/1998 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: 03/06/1998 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Pruritus		Rash mostly on arms and legs, mild on	Periactin 4mg tds, aurorix and zyprexa ceased.

Medicine Details:

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

AURORIX (Suspected) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128388 **Seq:** 1 **Gender:** F
Reported: 17/06/1998 **Weight:** 80.00
Hospitalisation: **Age:** 52Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neutropenia	Required Visit to Doctor		Aurorix and panadol ceased

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 08/04/1998 **Stopped:** 03/05/1998

PANADOL (Suspected) **Reason:** Headache
Oral application 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 09/04/1998 **Stopped:** 27/04/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pt has had previous episode of neutropenia (only on panadol at the time).

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 128746 **Seq:** 1 **Gender:** F
Reported: 29/06/1998 **Weight:** 61.00
Hospitalisation: **Age:** 32Y
Onset Date: 19/06/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suicide attempt		Contemplated suicide	
Disturbance in attention		Felt vague	Ceased aurorix
Asthenia		Had to lie down	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 09/06/1998 **Stopped:** 21/06/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129251 **Seq:** 1 **Gender:** F
Reported: 17/07/1998 **Weight:** 60.00
Hospitalisation: **Age:** 53Y
Onset Date: 25/02/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flatulence			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/10/1997 **Stopped:** 24/02/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No previous drug allergies 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129344 **Seq:** 1 **Gender:** F
Reported: 21/07/1998 **Weight:** 87.00
Hospitalisation: **Age:** 48Y
Onset Date: 09/07/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Confusional state Hallucination Myalgia Vertigo		Throbbing headache	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 07/07/1998 **Stopped:** 14/07/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129533 **Seq:** 1 **Gender:** F
Reported: 27/07/1998 **Weight:** 68.00
Hospitalisation: **Age:** 76Y
Onset Date: 08/07/1998 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Diarrhoea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 25/06/1998 **Stopped:** 13/07/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient's symptoms recurred when rechallenged on aurorix. see original report for other drugs.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129564 **Seq:** 1 **Gender:** M
Reported: 27/07/1998 **Weight:** 90.00
Hospitalisation: **Age:** 36Y
Onset Date: 31/12/1997 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Agitation Nausea	Severe	Could not sleep.	Ceased aurorix.

Medicine Details:

RIVOTRIL (Other drug) **Reason:** Other disturbance of sensation
1.0 Dose Unspecified Daily
Batch: **Started:** 01/10/1997 **Stopped:** 28/02/1998

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/12/1997 **Stopped:** 28/07/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient suffered worse symptoms when taking aurorix 600mg. - 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129817 **Seq:** 1 **Gender:** M
Reported: 31/07/1998 **Weight:**
Hospitalisation: **Age:** 79Y
Onset Date: 19/06/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Aurorix ceased

Medicine Details:

ZANTAC (Other drug) **Reason:** Other diseases of esophagus
Tablet 300.0 Milligram As necessary Oral
Batch: **Started:** 01/01/1982 **Stopped:**

CISAPRIDE (Other drug) **Reason:** Other diseases of esophagus
0.0 As necessary
Batch: **Started:** 01/01/1996 **Stopped:**

DILANTIN (Other drug) **Reason:** Other&unspecified epilepsy
Capsule 400.0 Milligram Daily Oral
Batch: **Started:** 01/01/1982 **Stopped:**

ENALAPRIL MALEATE (Other drug) **Reason:** Essential benign hypertension
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 01/01/1996 **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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129817 **Seq:** 1 **Gender:** M
Reported: 31/07/1998 **Weight:**
Hospitalisation: **Age:** 79Y
Onset Date: 19/06/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 01/01/1995 **Stopped:** 19/06/1998

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129992 **Seq:** 1 **Gender:** M
Reported: 10/08/1998 **Weight:** 60.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gout		Pseudo gout - pain in right knee & big toe	Moclobemide ceased

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 28/07/1998 **Stopped:** 02/08/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous exposure in 1996 with similar effect.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 129994 **Seq:** 1
Reported: 10/08/1998
Hospitalisation:
Onset Date: 31/05/1998
Outcome: Not yet recovered

Gender: M
Weight: 80.00
Age: 69Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		Left macular vision blurring (intermittent)	Aurorix ceased

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 825.0 Milligram Daily Oral

Batch: **Started:** 01/01/1997 **Stopped:** 28/06/1998

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130118 **Seq:** 1 **Gender:** M
Reported: 11/08/1998 **Weight:** 75.00
Hospitalisation: **Age:** 62Y
Onset Date: 24/07/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Fatigue			
Flushing			
Hyperhidrosis			
Hypertension			
Insomnia			
Serotonin syndrome			Ceased both pseudoephedrine and moclobemide

Medicine Details:

NORVASC (Other drug) **Reason:** Essential benign hypertension
0.0
Batch: **Started:** L TERM **Stopped:**

PSEUDOEPHEDRINE HYDROCHLORIDE (Suspected) **Reason:** Acute nasopharyngitis,com cold
0.0
Batch: **Started:** 24/07/1998 **Stopped:** 25/07/1998

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/04/1996 **Stopped:** 26/07/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Prior tolerance to moclobemide good. symptoms subsided on ceasing both pseudoephedrine and moclobemide.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130209 **Seq:** 1 **Gender:** M
Reported: 17/08/1998 **Weight:**
Hospitalisation: **Age:** 27Y
Onset Date: 11/06/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus		Electric noises in the head.	
Anxiety		Head rush to the point eyes couldn't see.	
Coordination abnormal		Loss of balance	
Chest pain		Rapid heartbeat sometimes, pain	
Tongue oedema		Swollen tongue and mouth	
Fatigue			
Visual disturbance			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 05/05/1998 **Stopped:** 11/06/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130305 **Seq:** 1 **Gender:** F
Reported: 19/08/1998 **Weight:** 100.00
Hospitalisation: **Age:** 55Y
Onset Date: 11/08/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension		Bp 170/90	
Headache		Dull headache	
Confusional state			
Dizziness			
Malaise			
Nausea			
Palpitations			

Medicine Details:

CATAPRES (Other drug) **Reason:**
200.0 Microgram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 11/08/1998 **Stopped:** 14/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130514 **Seq:** 1
Reported: 24/08/1998
Hospitalisation: Admitted to hospital
Onset Date: 17/08/1998
Outcome: Recovered

Gender: F
Weight:
Age: 35Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pyrexia		Elevated body temperature	
Confusional state		Mental confusion	
Chills			
Diarrhoea			
Hyperhidrosis			
Mydriasis			
Serotonin syndrome			Observed overnight in hospital, given rehydration, 1/2 hour observations and anti-nausea drugs. discharged home the following morning.
Syncope			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/02/1998 **Stopped:** 17/08/1998

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 17/08/1998 **Stopped:** 17/08/1998

EFEXOR (Suspected) **Reason:** Depression
Tablet 75.0 Milligram Daily Oral
Batch: **Started:** 17/08/1998 **Stopped:** 17/08/1998

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.

Report Detail

Case Number: 130514 **Seq:** 1
Reported: 24/08/1998
Hospitalisation: Admitted to hospital
Onset Date: 17/08/1998
Outcome: Recovered

Gender: F
Weight:
Age: 35Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting			

Medicine Details:

AURORIX (Suspected)			**Reason:** Depression	
Tablet	600.0 Milligram		Daily	Oral
Batch:	**Started:** 01/02/1998		**Stopped:**	17/08/1998
AURORIX (Suspected)			**Reason:** Depression	
Tablet	300.0 Milligram		Daily	Oral
Batch:	**Started:** 17/08/1998		**Stopped:**	17/08/1998
EFEXOR (Suspected)			**Reason:** Depression	
Tablet	75.0 Milligram		Daily	Oral
Batch:	**Started:** 17/08/1998		**Stopped:**	17/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130590 **Seq:** 1 **Gender:** M
Reported: 27/08/1998 **Weight:** 81.00
Hospitalisation: **Age:** 68Y
Onset Date: 18/08/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Choreoathetosis		Involuntary movements of limbs of	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 14/08/1998 **Stopped:** 21/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 130762 **Seq:** 1 **Gender:** F
Reported: 03/09/1998 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: 13/08/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysarthria Diarrhoea Headache Nausea	Severe	Slurred speech	

Medicine Details:

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

PREMARIN (Other drug) **Reason:** Menopausal symptoms
Tablet 925.0 Microgram Daily Oral
Batch: **Started:** L TERM **Stopped:**

MODURETIC (Other drug) **Reason:** Essential benign hypertension
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** L TERM **Stopped:**

PROVERA (Other drug) **Reason:** Menopausal symptoms
Tablet 5.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms started within 30 minutes of taking aurorix.

Report Detail

Case Number: 130762 **Seq:** 1 **Gender:** F
Reported: 03/09/1998 **Weight:**
Hospitalisation: **Age:** 61Y
Onset Date: 13/08/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

SLOW-K (Other drug)			**Reason:**	Essential benign hypertension	
Tablet	600.0	Milligram	Daily	Oral	
Batch:	**Started:**	L TERM	**Stopped:**		
NORVASC (Other drug)			**Reason:**	Essential benign hypertension	
Tablet	5.0	Milligram	Daily	Oral	
Batch:	**Started:**	31/07/1998	**Stopped:**		
VALIUM (Other drug)			**Reason:**	Anxiety neurosis	
Tablet	5.0	Milligram	Daily	Oral	
Batch:	**Started:**	07/04/1998	**Stopped:**		
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	300.0	Milligram	Daily	Oral	
Batch:	**Started:**	13/08/1998	**Stopped:**	13/08/1998	

Laboratory Investigations:

Additional Information:

Symptoms started within 30 minutes of taking aurorix.

Report Detail

Case Number: 131030 **Seq:** 1 **Gender:** M
Reported: 11/09/1998 **Weight:**
Hospitalisation: **Age:** AD
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor		Symptoms similar to electric shock	

Medicine Details:

CODRAL DAYTIME/NIGHTIME TABLETS (Interaction) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 131516 **Seq:** 1 **Gender:** F
Reported: 25/09/1998 **Weight:**
Hospitalisation: **Age:** 34Y
Onset Date: 28/08/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dystonia Hypertension Malaise Serotonin syndrome Tachycardia			

Medicine Details:

ARIMA (Interaction) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/02/1998 **Stopped:** 26/08/1998

PAROXETINE HYDROCHLORIDE (Interaction) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** 28/08/1998 **Stopped:** 28/08/1998

TEMAZEPAM (Interaction) **Reason:** Specific disorders of sleep
20.0 Milligram Daily
Batch: **Started:** 28/08/1998 **Stopped:** 28/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 131524 **Seq:** 1
Reported: 25/09/1998
Hospitalisation:
Onset Date:
Outcome: Not yet recovered

Gender: F
Weight: 56.00
Age: 25Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		Blurred vision cant focus on distance vision	
Rash maculo-papular		Fine rash papular on face	
Dizziness			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/08/1998 **Stopped:** 28/09/1998

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. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 131749 **Seq:** 1 **Gender:** F
Reported: 01/10/1998 **Weight:**
Hospitalisation: **Age:** 52Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia			Aurorix ceased and changed to

Medicine Details:

AURORIX (Suspected) **Reason:**
450.0 Milligram Daily
Batch: **Started:** 21/08/1998 **Stopped:** 24/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 131827 **Seq:** 1 **Gender:** F
Reported: 07/10/1998 **Weight:** 75.00
Hospitalisation: **Age:** 70Y
Onset Date: 22/09/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Agitation Confusional state Diarrhoea		Unsteadiness	

Medicine Details:

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

MOGADON (Other drug) **Reason:** Specific disorders of sleep
2.5 Milligram Daily
Batch: **Started:** 01/01/1980 **Stopped:**

ALCOHOL (Other drug) **Reason:**
100.0 Millilitre Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 21/09/1998 **Stopped:** 26/09/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 132210 **Seq:** 1
Reported: 21/10/1998
Hospitalisation:
Onset Date: 14/10/1998
Outcome: Not yet recovered

Gender: F
Weight: 64.00
Age: 43Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Palpitations			Aurorix decreased to 150mg from 300mg daily.

Medicine Details:

MESALAZINE (Other drug) **Reason:** Ulcerative colitis
4.0 Dose Unspecified Daily
Batch: **Started:** 07/10/1998 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 21/10/1992 **Stopped:** 14/10/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 132243 **Seq:** 1 **Gender:** F
Reported: 22/10/1998 **Weight:** 43.00
Hospitalisation: **Age:** 38Y
Onset Date: 14/10/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypoaesthesia Anxiety Dizziness Malaise	Severe	Numbness of fingers and toes.	

Medicine Details:

SERZONE (Suspected) **Reason:**
100.0 Milligram 1 time
Batch: **Started:** 14/10/1998 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/08/1998 **Stopped:** 12/10/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 132788 **Seq:** 1 **Gender:** M
Reported: 04/11/1998 **Weight:**
Hospitalisation: **Age:** 44Y
Onset Date: 31/05/1998 **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia Hypoaesthesia		Parasthesia of fingertips.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Ceased aurorix and symptoms resolved, re-started aurorix and recurrence of symptoms. - 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 132847 **Seq:** 1
Reported: 04/11/1998
Hospitalisation: Admitted to hospital
Onset Date: 07/08/1998
Outcome: Recovered

Gender: F
Weight: 50.00
Age: 70Y
DOB:
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety Dry mouth Palpitations			Ceased moclobemide

Medicine Details:

SINEMET CR (Other drug) **Reason:** Paralysis agitans
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 01/01/1997 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 1.2 Gram Daily Oral
Batch: **Started:** 14/07/1998 **Stopped:** 07/08/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 132909 **Seq:** 1 **Gender:** M
Reported: 06/11/1998 **Weight:**
Hospitalisation: **Age:** 53Y
Onset Date: 29/10/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Disturbance in attention		Poor concentration	
Somnolence		Sleepy	

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 23/10/1998 **Stopped:** 29/10/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 133206 **Seq:** 1 **Gender:** M
Reported: 16/11/1998 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 06/11/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agranulocytosis			Timentin, gentamicin, g-csf 263 mcg (2 doses).

Medicine Details:

TICLID (Suspected) **Reason:** Transient cereb isch no hypert
500.0 Milligram Daily
Batch: **Started:** 06/10/1998 **Stopped:** 06/11/1998

ACCUPRIL (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:** 06/11/1998

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 13/10/1998 **Stopped:** 06/11/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Haemoglobin		06/11/1998	117	
	Lymphocytes		06/11/1998	1.2	
	Monocytes		06/11/1998	1.0	
	Neutrophils		06/11/1998	0.0	
	Neutrophils		08/11/1998	8.9	
	Neutrophils		07/11/1998	0.6	
	Red blood cells		06/11/1998	3.98	

Additional Information:

Report Detail

Case Number: 133206 **Seq:** 1 **Gender:** M
Reported: 16/11/1998 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 06/11/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

TICLID (Suspected) **Reason:** Transient cereb isch no hypert
500.0 Milligram Daily
Batch: **Started:** 06/10/1998 **Stopped:** 06/11/1998

ACCUPRIL (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:** 06/11/1998

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 13/10/1998 **Stopped:** 06/11/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	White blood cells		06/11/1998	2.3	
	White blood cells		07/11/1998	4.0	
	White blood cells		08/11/1998	13	

Additional Information:

Report Detail

Case Number: 133448 **Seq:** 1 **Gender:** F
Reported: 20/11/1998 **Weight:** 134.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective			Transferred to cipramil.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 133651 **Seq:** 1
Reported: 23/11/1998
Hospitalisation:
Onset Date: 15/09/1998
Outcome: Not yet recovered

Gender: F
Weight: 57.00
Age: 39Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation		Spacey feeling	
Fatigue		Very tired and drained.	
Confusional state			
Depression			
Dizziness			
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:**

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 08/09/1998 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 133734 **Seq:** 1 **Gender:** F
Reported: 26/11/1998 **Weight:** 62.00
Hospitalisation: **Age:** 56Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Glossitis			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms to moclobemide cleared within 10 days of withdrawal and reappeared after commencing drug 6months later.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 133750 **Seq:** 1 **Gender:** F
Reported: 27/11/1998 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: **DOB:**
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis Jaundice	Severe		

Medicine Details:

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

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

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Antinuclear factor Bilirubin				Positive ana Bilirubin up to 400

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 134146 **Seq:** 1 **Gender:** M
Reported: 10/12/1998 **Weight:** 95.00
Hospitalisation: **Age:** 48Y
Onset Date: 31/10/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Back pain			

Medicine Details:

ENDEP (Other drug) **Reason:** Anxiety neurosis
50.0 Milligram Daily
Batch: **Started:** **Stopped:** CONTIN

ZESTRIL (Other drug) **Reason:** Essential benign hypertension
10.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:**

PLENDIL (Other drug) **Reason:** Essential benign hypertension
5.0 Milligram Daily
Batch: **Started:** 01/01/1996 **Stopped:**

AURORIX (Suspected) **Reason:** Anxiety neurosis
1.2 Gram Daily
Batch: **Started:** 01/10/1998 **Stopped:** 28/11/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Dose related back pain - ceased 1 week after reducing dose. - 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.

Report Detail

Case Number: 134392 **Seq:** 1
Reported: 18/12/1998
Hospitalisation: Admitted to hospital
Onset Date: 31/12/1998
Outcome: Recovered

Gender: F
Weight:
Age: 45Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Confusional state Depersonalisation Tachycardia			

Medicine Details:

ZOLOFT (Suspected) **Reason:** Depression

Tablet 50.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 18M

AURORIX (Suspected) **Reason:** Depression

Tablet 300.0 Milligram Daily Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Moclobemide was started after 1 day of ceasing zoloft. - the date of onset is not accurate but indicates that onset occurred sometime during the year.

Report Detail

Case Number: 134527 **Seq:** 1 **Gender:** F
Reported: 21/12/1998 **Weight:** 52.00
Hospitalisation: **Age:** 38Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia		Unable to sit down for any period of time.	 Aurorix ceased.

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 1M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 134716 **Seq:** 1 **Gender:** M
Reported: 04/01/1999 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypersensitivity			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram 1 time Oral
Batch: **Started:** 27/12/1998 **Stopped:** 27/12/1998

AROPAX (Suspected) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** 18/11/1998 **Stopped:** 26/12/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pt suffered symptoms half hour after taking first tab of aurorix. known allergy to coconut, peanut and penicillin.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135058 **Seq:** 1 **Gender:** F
Reported: 11/01/1999 **Weight:**
Hospitalisation: **Age:** 18Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Convulsion			
Hypertonia			
Hypotension			
Muscle twitching			
Serotonin syndrome			12mg cyproheptadine.
Speech disorder			

Medicine Details:

CLOMIPRAMINE HYDROCHLORIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:** 3M

MOCLOBEMIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Mja vol 169 16/11/98 p. 523 changed to moclobemide with no washout period.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135062 **Seq:** 1 **Gender:** M
Reported: 11/01/1999 **Weight:**
Hospitalisation: **Age:** 32Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension		Bp 162/107	
Hypertonia		Muscle rigidity	
Agitation			
Dizziness			
Dystonia			
Hallucination			
Hyperhidrosis			
Mydriasis			
Nystagmus			
Serotonin syndrome			Diazepam 10mg and chlorpromazine 12.5mg.

Medicine Details:

VENLAFAXINE HYDROCHLORIDE (Interaction) **Reason:**
150.0 Milligram 1 time
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

DIAZEPAM (Other drug) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

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Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135062 **Seq:** 1 **Gender:** M
Reported: 11/01/1999 **Weight:**
Hospitalisation: **Age:** 32Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting			

Medicine Details:

VENLAFAXINE HYDROCHLORIDE (Interaction) **Reason:**
150.0 Milligram 1 time
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:**
40.0 Milligram Daily
Batch: **Started:** **Stopped:**

DIAZEPAM (Other drug) **Reason:**
15.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Additional Information:

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Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135063 **Seq:** 1
Reported: 11/01/1999
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 82Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Speech disorder Agitation Confusional state Serotonin syndrome Tremor		Could only answer yes or no to questions	4mg cyproheptadine.

Medicine Details:

FLUOXETINE HYDROCHLORIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

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THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135106 **Seq:** 1 **Gender:** F
Reported: 14/01/1999 **Weight:** 65.00
Hospitalisation: **Age:** 38Y
Onset Date: 05/01/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus Rash			Phenergan 20mg and stopped aurorix.

Medicine Details:

INTERFERON ALFA-2A (Other drug) **Reason:** Hiv infection
3.0 Dose Unspecified Weekly
Batch: **Started:** 01/09/1998 **Stopped:**

QUININE (Other drug) **Reason:** Pain in limb
300.0 Milligram Daily
Batch: **Started:** 01/12/1998 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 03/01/1999 **Stopped:** 06/01/1999

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 135758 **Seq:** 1 **Gender:** F
Reported: 02/02/1999 **Weight:**
Hospitalisation: **Age:** 77
Onset Date: **DOB:** 02/03/1921
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia Polydipsia			Mild fluid restriction.

Medicine Details:

MOCLOBEMIDE (Suspected) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 136417 **Seq:** 1 **Gender:** F
Reported: 19/02/1999 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/01/1999 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation		Felt very excited and uncomfortable.	Was suggested reducing the dose of zolft from 50mg to

Medicine Details:

ZOLOFT (Suspected)			**Reason:**	Depression	
Tablet	50.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression	
Tablet	300.0	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 136532 **Seq:** 1 **Gender:** F
Reported: 23/02/1999 **Weight:** 55.00
Hospitalisation: **Age:** 35Y
Onset Date: 19/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Nausea Nystagmus Palpitations Tinnitus Vertigo		Unbalance	

Medicine Details:

DEMAZIN TABLET (Interaction) **Reason:** Chronic nasopharyngitis
Tablet 1.0 Dose Unspecified 1 time Oral
Batch: **Started:** 18/02/1999 **Stopped:**

AURORIX (Interaction) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/12/1998 **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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 136633 **Seq:** 1 **Gender:** F
Reported: 25/02/1999 **Weight:** 50.00
Hospitalisation: **Age:** 86Y
Onset Date: 02/11/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Moclobemide ceased.

Medicine Details:

COLOXYL WITH SENNA (Other drug) **Reason:**
0.0
Batch: **Started:** 25/10/1998 **Stopped:**

TEGRETOL (Other drug) **Reason:** Other&unspecified epilepsy
Tablet 400.0 Milligram Daily Oral
Batch: **Started:** 30/10/1998 **Stopped:** 02/11/1998

ASPIRIN (Other drug) **Reason:**
Oral application 150.0 Milligram Daily Oral
Batch: **Started:** 25/10/1998 **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium		26/10/1998	135	
	Sodium		02/11/1998	122	
	Sodium		05/11/1998	130	
	Sodium		06/11/1998	135	

Additional Information:

Report Detail

Case Number: 136633 **Seq:** 1 **Gender:** F
Reported: 25/02/1999 **Weight:** 50.00
Hospitalisation: **Age:** 86Y
Onset Date: 02/11/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

ATORVASTATIN (Other drug)			**Reason:**		
Tablet	10.0 Milligram		Daily	Oral	
Batch:	**Started:** 25/10/1998			**Stopped:**	CONTIN
MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
	600.0 Milligram		Daily		
Batch:	**Started:** 29/10/1998			**Stopped:**	04/11/1998

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 137069 **Seq:** 1 **Gender:** M
Reported: 08/03/1999 **Weight:**
Hospitalisation: **Age:** 74Y
Onset Date: 31/12/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Flushing			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

0.0

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient was started on aurorix which controls his symptoms. patient has developed flushing, which he finds quite irritating. - the date of onset is not accurate but indicates that onset occurred sometime during the year.

Report Detail

Case Number: 137268 **Seq:** 1 **Gender:** F
Reported: 12/03/1999 **Weight:**
Hospitalisation: **Age:** 57Y
Onset Date: 18/02/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysgeusia Nausea Tinnitus			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 11/02/1999 **Stopped:** 08/03/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergic to penicillian

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 137581 **Seq:** 1 **Gender:** F
Reported: 19/03/1999 **Weight:** 0.00
Hospitalisation: **Age:** 28
Onset Date: **DOB:** 06/11/1970
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia			

Medicine Details:

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

TEGRETOL CR (Suspected) **Reason:** Unspecifid affective psychosis
600.0 Milligram Daily Oral
Batch: **Started:** 01/10/1998 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/12/2002	GGT = SGGT =				Elevation of ggtp to 69u/l.

Additional Information:

See original report for other drug details - moclabemide. patient complained of marked short term memory problems, eg forgetting the content of recent conversations which occur only minutes beforehand. tegretol cr was lowered to 500mg/day, the to 400mg/day in march 99. memory problems resolved but relapse of mood symptoms. on 24 march 99 mood symptoms settled but memory problem reutrned. by mistake patient had increased the dose of tegretol cr to 600mg/day instead of 500mg/day. when decreased to 500mg/day memory problem was a lot better. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 137762 **Seq:** 1 **Gender:** F
Reported: 23/03/1999 **Weight:** 58.00
Hospitalisation: **Age:** 75Y
Onset Date: 26/11/1998 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Trunk and upper arm	Ceased moclobemide

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 25/11/1998 **Stopped:** 26/11/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Please see original report for details of other drugs - uremide, arthrotec, digoxin (lanoxin), irbesartan, famotidine (pepcidine), caltrate, gtn patches, pravastatin (pravachol), pulmicort.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 138013 **Seq:** 1 **Gender:** F
Reported: 31/03/1999 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myalgia Paraesthesia			

Medicine Details:

MACRODANTIN (Suspected) **Reason:** Pyuria and bacteriuria
Capsule 100.0 Milligram Daily Oral
Batch: **Started:** 01/12/1998 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 0.0 Daily Oral
Batch: **Started:** **Stopped:** CONTIN

COVERSYL (Suspected) **Reason:** Essential benign hypertension
Tablet 0.0 Daily Oral
Batch: **Started:** **Stopped:** CONTIN

LASIX (Suspected) **Reason:**
Oral application 0.0 Daily Oral
Batch: **Started:** **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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 138013 **Seq:** 1 **Gender:** F
Reported: 31/03/1999 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Medicine Details:

TEMAZE (Suspected)		**Reason:**		
Oral application	0.0	Daily	Oral	
Batch:	**Started:**		**Stopped:**	CONTIN

Laboratory Investigations:

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 138411 **Seq:** 1 **Gender:** F
Reported: 13/04/1999 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: 09/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tongue disorder Dysphagia Tongue oedema		Dysaesthesia of tongue, extrapyramidal	Moclobemide ceased. cogentin 1mg given.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 05/02/1999 **Stopped:** 10/02/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Prothiaden - extrapyramidal side effect. previous allergy reaction. see original report for details of other drugs. methadone, ranitidine, lorazepam, digesic, proladone supp and amphotericin.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 138787 **Seq:** 1 **Gender:** M
Reported: 27/04/1999 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 14/04/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatitis Renal failure acute			

Medicine Details:

AMOXYCILLIN SODIUM (Suspected) **Reason:** Pneumonia,unspecified
0.0
Batch: **Started:** 06/04/1999 **Stopped:** 14/04/1999

AURORIX (Suspected) **Reason:** Depression
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 20/04/1999

WARFARIN SODIUM (Suspected) **Reason:**
Tablet 2.0 Milligram Daily Oral
Batch: **Started:** 01/05/1998 **Stopped:** 14/04/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		14/04/1999	2540	
	AST = SGOT		14/04/1999	1930	
	Albumin		14/04/1999	31	
	Bilirubin		14/04/1999	27	
	Bilirubin		20/04/1999	67	
	Biochemistry				All hepatic serology negative.
	GGT = SGGT =		14/04/1999	183	

Additional Information:

See original report for other drugs: prednisolone, allopurinol (progout), famotidine. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 138787 **Seq:** 1 **Gender:** M
Reported: 27/04/1999 **Weight:**
Hospitalisation: **Age:** 76Y
Onset Date: 14/04/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

AMOXYCILLIN SODIUM (Suspected)			**Reason:**	Pneumonia,unspecified
	0.0			
Batch:	**Started:** 06/04/1999		**Stopped:**	14/04/1999
AURORIX (Suspected)			**Reason:**	Depression
Tablet	60.0 Milligram	Daily		Oral
Batch:	**Started:**		**Stopped:**	20/04/1999
WARFARIN SODIUM (Suspected)			**Reason:**	
Tablet	2.0 Milligram	Daily		Oral
Batch:	**Started:** 01/05/1998		**Stopped:**	14/04/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	LDH		14/04/1999	1131	
	Protein		14/04/1999	49	
	SAP = ALP		14/04/1999	183	

Additional Information:

See original report for other drugs: prednisolone, allopurinol (progout), famotidine. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139167 **Seq:** 1 **Gender:** F
Reported: 03/05/1999 **Weight:**
Hospitalisation: **Age:** 32Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Insomnia Migraine			

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139252 **Seq:** 1 **Gender:** M
Reported: 06/05/1999 **Weight:**
Hospitalisation: **Age:** 23Y
Onset Date: 03/05/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Agitation Chills Confusional state Dizziness Migraine Nausea Vomiting			Ceased zoloft

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

ZOLOFT (Interaction) **Reason:** Depression
5.0 Milligram Daily
Batch: **Started:** 03/05/1999 **Stopped:** 04/05/1999

STELAZINE (Interaction) **Reason:**
5.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for details of other drug therapy: serepax.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139467 **Seq:** 1 **Gender:** M
Reported: 13/05/1999 **Weight:** 72.00
Hospitalisation: **Age:** 83Y
Onset Date: 26/03/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia		Hair loss	
Vomiting		Occational vomiting	
Rash		Rash on hairline and temple.	
Abdominal pain			
Diarrhoea			
Nausea			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 22/03/1999 **Stopped:** 13/04/1999

TICLID (Suspected) **Reason:** Prophylaxis
Tablet 500.0 Milligram Daily Oral
Batch: **Started:** 12/02/1999 **Stopped:** 13/04/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drug details: zantac & claratyne.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139516 **Seq:** 1 **Gender:** F
Reported: 14/05/1999 **Weight:**
Hospitalisation: **Age:**
Onset Date: 09/05/1999 **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia		Developed aching joints	

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Commenced on aurorix 150mg morning. by 3pm patient developed reaction. this lasted until about 12 midnight to 3am. this occurred for the 4 consecutive days the patient took the aurorix.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139735 **Seq:** 1 **Gender:** F
Reported: 21/05/1999 **Weight:**
Hospitalisation: **Age:** 22Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast pain		Patient experienced breast pain.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139748 **Seq:** 1 **Gender:** F
Reported: 21/05/1999 **Weight:**
Hospitalisation: **Age:** 32
Onset Date: 01/02/1999 **DOB:** 01/10/1966
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Patient developed a skin rash feb 99.	Has not improved on

Medicine Details:

AURORIX (Suspected) **Reason:** Phobic neurosis
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/12/1998 **Stopped:** 30/03/1999

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139784 **Seq:** 1 **Gender:** F
Reported: 24/05/1999 **Weight:**
Hospitalisation: **Age:** 70Y
Onset Date: 31/01/1999 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anorexia Ageusia		Loss of appetite	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 10/12/1998 **Stopped:** 17/05/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drug: aspirin - the date of onset is not necessarily accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 139817 **Seq:** 1 **Gender:** F
Reported: 24/05/1999 **Weight:** 43.00
Hospitalisation: **Age:** 15Y
Onset Date: 01/04/1999 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Insomnia		Patient complained of abdo discomfort. Worsening sleep disturbance.	Ceased aurorix - also because been on it for 10/52 oral with no real improvement in mood or sleep, temazepam 10mg given with effect.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 12/04/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient history: 24.2.99 not menstrated fr last 2 months, anorexia for approx 1 year folloing advice to lose wt for knee injury (rapid wt loss of 5kg in 1 month) & now symptomatic, nka, no other medications, vit other than aurorix.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 140251 **Seq:** 1 **Gender:** M
Reported: 04/06/1999 **Weight:**
Hospitalisation: **Age:** 56Y
Onset Date: 05/05/1999 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/03/1999 **Stopped:** 28/05/1999

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 140353 **Seq:** 1 **Gender:** F
Reported: 08/06/1999 **Weight:**
Hospitalisation: **Age:** 42
Onset Date: **DOB:** 22/04/1957
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amenorrhoea	Required Specialist Consultation		
Galactorrhoea	Required Specialist Consultation		
Menstrual disorder	Required Specialist Consultation		

Medicine Details:

AROPAX (Suspected) **Reason:** Depression
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 18/02/1999 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/01/1999 **Stopped:** 18/02/1999

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.

Report Detail

Case Number: 142002 **Seq:** 1 **Gender:** M
Reported: 28/07/1999 **Weight:** 90.00
Hospitalisation: **Age:** 69Y
Onset Date: 31/07/1998 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypoaesthesia Neuropathy peripheral Paraesthesia			Ceased aurorix

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 30/09/1997 **Stopped:** 15/06/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrographics				Nerve conduction studies confirm neuropathy peripheral

Additional Information:

Please see original report for details of other drugs - naprosyn, progout, timoptol. - the date of onset is not necessarily accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 142820 **Seq:** 1 **Gender:** M
Reported: 20/08/1999 **Weight:** 69.00
Hospitalisation: **Age:** 62Y
Onset Date: 12/08/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			
Dizziness			
Hyperhidrosis			
Hyperreflexia			
Hypertonia			
Nausea			
Serotonin syndrome			Treated with iv hydration. ceased moclobemide. started on bromocriptine.
Thinking abnormal			
Tremor			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 29/07/1999 **Stopped:** 12/08/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Please see original report for details of other drugs - sinemet cr, domperidone,sinemet 100/25, zantac.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 142918 **Seq:** 1 **Gender:** F
Reported: 23/08/1999 **Weight:**
Hospitalisation: **Age:** 42Y
Onset Date: 31/12/1999 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Paraesthesia			Zoloft dose was reduced from 100mg to 50mg/d. patient treated skin reaction with cold water and sorbelene cream.

Medicine Details:

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

AURORIX (Suspected) **Reason:**
Tablet 1.2 Gram Daily Oral
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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 143076 **Seq:** 1 **Gender:** M
Reported: 27/08/1999 **Weight:**
Hospitalisation: **Age:** 37Y
Onset Date: **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder		Eps due to the use of neuroleptics.	

Medicine Details:

FLUPENTHIXOL DECANOATE (Suspected) **Reason:**
0.0
Batch: **Started:** 01/05/1999 **Stopped:** CONTIN

MOCLOBEMIDE (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 01/01/1998 **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 accurate but indicates that start occurred sometime during the year.

Report Detail

Case Number: 143492 **Seq:** 1 **Gender:** M
Reported: 06/09/1999 **Weight:**
Hospitalisation: **Age:**
Onset Date: 31/01/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erythema multiforme		Lesions on entire body	Emolients, sorbolene, glycerin.

Medicine Details:

COLOXYL WITH SENNA (Suspected) **Reason:**
0.0 As necessary
Batch: **Started:** **Stopped:** CONTIN

ASPIRIN (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 11/12/1998 **Stopped:** 12/07/1999

TEMAZEPAM (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:** 12/07/1999

RANITIDINE (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 14/07/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Skin biopsy				Erythema multiforme proven on biopsy

Additional Information:

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

Report Detail

Case Number: 143492 **Seq:** 1
Reported: 06/09/1999
Hospitalisation:
Onset Date: 31/01/1999
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

AMIODARONE HYDROCHLORIDE (Suspected)			**Reason:**	Otr&nos disord of heart rhythm		
	200.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**		CONTIN
FLUDROCORTISONE (Suspected)			**Reason:**			
	100.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**		CONTIN
MOCLOBEMIDE (Suspected)			**Reason:**			
	300.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**	12/07/1999	
CELLUFRESH (Suspected)			**Reason:**			
Eye Drops	8.0	Dose Unspecified	Daily		Ophthalmic	
Batch:	**Started:**			**Stopped:**		CONTIN

Laboratory Investigations:

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 143492 **Seq:** 1
Reported: 06/09/1999
Hospitalisation:
Onset Date: 31/01/1999
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

SLOW-K (Suspected) **Reason:**

4.0 Dose Unspecified Daily

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

Laboratory Investigations:

Additional Information:

- the date of onset is not necessarily accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 143508 **Seq:** 1 **Gender:** M
Reported: 08/09/1999 **Weight:** 80.00
Hospitalisation: **Age:** 44Y
Onset Date: 01/06/1999 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia		Gynaecomastia after 3-4 months of use the	

Medicine Details:

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** 01/02/1999 **Stopped:** 01/07/1999

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 143519 **Seq:** 1 **Gender:** F
Reported: 09/09/1999 **Weight:** 80.00
Hospitalisation: **Age:** 49Y
Onset Date: 27/08/1999 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anxiety Tremor		Felt anxious and over stimulated.	

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 23/08/1999 **Stopped:** 28/08/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Rechallenge:- stopped for four days- felt better, then restarted it & the same symptoms returned.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 143627 **Seq:** 1 **Gender:** F
Reported: 13/09/1999 **Weight:** 70.00
Hospitalisation: **Age:** 43Y
Onset Date: 14/08/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation		Panic and feelings of fear.	
Anxiety		Severe exacerbation of anxiety symptoms.	Augmentin stopped
Agitation			
Insomnia			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 17/09/1998 **Stopped:**

AUGMENTIN DUO FORTE (Suspected) **Reason:** Chronic sinusitis
1.0 Dose Unspecified Daily
Batch: **Started:** 12/08/1999 **Stopped:** 16/08/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms resolved within 1 week of stopping augmentin.

Report Detail

Case Number: 143801 **Seq:** 1 **Gender:** M
Reported: 17/09/1999 **Weight:** 115.00
Hospitalisation: **Age:** 33Y
Onset Date: 31/07/1999 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain Myalgia		Chest wall ache on rising lasting 1-2hours.	Aurorix therapy ceased.

Medicine Details:

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

DIFFLAM GEL (Other drug) **Reason:**
0.0
Batch: **Started:** **Stopped:**

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

AURORIX (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 30/06/1999 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms occurred seven to ten days after therapy with aurorix commenced. - 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 145078 **Seq:** 1 **Gender:** M
Reported: 27/10/1999 **Weight:** 84.00
Hospitalisation: **Age:** 60Y
Onset Date: 31/05/1999 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Erectile dysfunction Libido decreased			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 900.0 Milligram Daily Oral

Batch: **Started:** 23/03/1999 **Stopped:** 21/06/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Aurorix increased from 300mg bd around 7/6/99. noted some change on 300mg bd but rapidly worsened when dose increased to 450mg bd. - the date of onset is not necessarily accurate but indicates that onset occurred sometime during the month.

Report Detail

Case Number: 145771 **Seq:** 1
Reported: 09/11/1999
Hospitalisation: Admitted to hospital
Onset Date: 20/07/1999
Outcome: Recovered

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Syncope Confusional state Hyponatraemia Inappropriate antidiuretic hormon		Falls	Ceased dilantin, solprin, aurorix & lipex. aspirin restarted.

Medicine Details:

DILANTIN (Suspected) **Reason:** Other&unspecified epilepsy
400.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** 20/07/1999 25Y

SOLPRIN (Suspected) **Reason:** Acute,ill-def cerebrovascular disease w/o hyperte
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 10/06/1999 **Stopped:** 20/07/1999

LIPEX (Suspected) **Reason:** Othr&unspec metabolic diseases
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level		20/07/1999	147	Phenytoin level
	Sodium		11/02/1999	136	
	Sodium		10/07/1999	125	
	Sodium		20/07/1999	121	
	Sodium		22/07/1999	124	
	Sodium		23/07/1999	124	
	Sodium		26/07/1999	133	

Additional Information:

Report Detail

Case Number: 145771 **Seq:** 1
Reported: 09/11/1999
Hospitalisation: Admitted to hospital
Onset Date: 20/07/1999
Outcome: Recovered

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

Reaction Detail

Medicine Details:

DILANTIN (Suspected)			**Reason:**	Other&unspecified epilepsy
	400.0 Milligram	Daily		
Batch:	**Started:**	L TERM	**Stopped:**	20/07/1999 25Y
SOLPRIN (Suspected)			**Reason:**	Acute,ill-def cerebrovascular disease w/o hyperte
	150.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	
AURORIX (Suspected)			**Reason:**	Depression
	300.0 Milligram	Daily		
Batch:	**Started:** 10/06/1999		**Stopped:**	20/07/1999
LIPEX (Suspected)			**Reason:**	Othr&unspec metabolic diseases
	20.0 Milligram	Daily		
Batch:	**Started:**		**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium		28/07/1999	134	
	Sodium		30/07/1999	136	
	Sodium		17/08/1999	138	
	Urine Sodium		22/07/1999	< 10	
	Urine osmolality		22/07/1999	494	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 145936 **Seq:** 1 **Gender:** M
Reported: 17/11/1999 **Weight:**
Hospitalisation: **Age:** 38Y
Onset Date: 18/10/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Confusional state Dystonia Pyrexia		Severe muscle dystona	

Medicine Details:

PAROXETINE HYDROCHLORIDE (Interaction) **Reason:**
0.0
Batch: **Started:** 10/10/1999 **Stopped:** 18/10/1999

MOCLOBEMIDE (Interaction) **Reason:**
300.0 Milligram Daily
Batch: **Started:** 18/10/1999 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for details on other drugs: tegretol, epilim, murelax. had been told to stop paroxetine for 2 days prior to commencing moclobemide.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 146140 **Seq:** 1 **Gender:** F
Reported: 23/11/1999 **Weight:** 70.00
Hospitalisation: **Age:** 42Y
Onset Date: 18/10/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nightmare			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** 11/10/1999 **Stopped:** 23/10/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Higher dose of 900mg aurorix caused symptoms, previously ok at 600mg.

Report Detail

Case Number: 146559 **Seq:** 1 **Gender:** F
Reported: 06/12/1999 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypertension Migraine		Bp 190/120	

Medicine Details:

CODRAL FLU AND COLD TABLETS (Interaction) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 15/10/1999 **Stopped:** 10/11/1999

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 146671 **Seq:** 1 **Gender:** F
Reported: 08/12/1999 **Weight:**
Hospitalisation: **Age:** 40Y
Onset Date: 31/12/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective		Depression not effectively treated.	Aurorix discontinued

Medicine Details:

AURORIX (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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 146751 **Seq:** 1 **Gender:** F
Reported: 10/12/1999 **Weight:** 64.00
Hospitalisation: **Age:** 75Y
Onset Date: 29/11/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Abdominal pain Dry mouth Tinnitus		Aggravation of reflux	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
75.0 Milligram Daily
Batch: **Started:** 29/11/1999 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pt also taking oroxine, methotrexate, valium, zantac.

Report Detail

Case Number: 146817 **Seq:** 1 **Gender:** M
Reported: 13/12/1999 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 30/11/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal			

Medicine Details:

CALTRATE (Other drug) **Reason:** Other diseases of bone
1.8 Gram Daily
Batch: **Started:** 01/01/1998 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:**
0.0
Batch: **Started:** **Stopped:** 28/10/1999

OLANZAPINE (Suspected) **Reason:** Depression
1.0 Dose Unspecified 1 time
Batch: **Started:** 01/10/1999 **Stopped:** 29/11/1999

VALIUM (Suspected) **Reason:** Depression
1.0 Dose Unspecified 1 time
Batch: **Started:** 01/10/1999 **Stopped:** 29/11/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT	(0-55)	30/11/1999	114	
	ALT = SGPT	(0-55)	07/12/1999	70	
	GGT = SGGT =	(0-60)	30/11/1999	113	
	GGT = SGGT =	(0-60)	07/12/1999	86	
	SAP = ALP	(30-110)	30/11/1999	109	
	SAP = ALP	(30-110)	07/12/1999	117	

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 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 146817 **Seq:** 1
Reported: 13/12/1999
Hospitalisation:
Onset Date: 30/11/1999
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

AROPAX (Suspected)		**Reason:**	Depression
	40.0 Milligram	Daily	
Batch:	**Started:** 01/10/1999	**Stopped:**	CONTIN
RANITIDINE (Suspected)		**Reason:**	Otr disorders stomach function
	300.0 Milligram	Daily	
Batch:	**Started:** 01/10/1999	**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 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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 146929 **Seq:** 1 **Gender:** F
Reported: 14/12/1999 **Weight:** 57.00
Hospitalisation: **Age:** 25Y
Onset Date: 01/12/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photosensitivity reaction Herpes simplex Skin exfoliation		Severe photosensitivity reaction	

Medicine Details:

NORIMIN (Other drug) **Reason:** Contraception
0.0
Batch: **Started:** L TERM **Stopped:** CONTIN

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 01/11/1999 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 147907 **Seq:** 1 **Gender:** M
Reported: 12/01/2000 **Weight:**
Hospitalisation: **Age:** 36
Onset Date: 08/12/1999 **DOB:** 08/12/1963
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thinking abnormal		Mind racing	
Anxiety			
Convulsion			
Dizziness			
Flushing			
Headache			
Insomnia			
Muscle spasms			
Serotonin syndrome			

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 20/09/1999 **Stopped:** 27/11/1999

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 147954 **Seq:** 1 **Gender:** F
Reported: 24/12/1999 **Weight:**
Hospitalisation: **Age:** 87Y
Onset Date: 11/10/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Iv frusemide, n/saline, iv hydrocortisone.

Medicine Details:

ARIMA (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 18/09/1999 **Stopped:** 15/10/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium	MMOL/L	17/09/1999	134	
	Sodium	MMOL/L	22/09/1999	130	
	Sodium	MMOL/L	02/10/1999	136	
	Sodium	MMOL/L	04/10/1999	136	
	Sodium	MMOL/L	11/10/1999	124	
	Sodium	MMOL/L	14/10/1999	119	
	Sodium	MMOL/L	18/10/1999	133	

Additional Information:

Pt also taking aspirin, coversyl, uremide.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 148256 **Seq:** 1 **Gender:** F
Reported: 21/01/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Coordination abnormal Dizziness		Lack of balance	

Medicine Details:

ALCOHOL (Suspected) **Reason:**
1.0 Dose Unspecified 1 time
Batch: **Started:** **Stopped:**

DOSTINEX (Suspected) **Reason:**
Tablet 1.0 Milligram Weekly Oral
Batch: **Started:** 01/12/1999 **Stopped:**

AURORIX (Suspected) **Reason:**
Tablet 0.0 Daily Oral
Batch: **Started:** **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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 148420 **Seq:** 1 **Gender:** F
Reported: 31/01/2000 **Weight:** 65.00
Hospitalisation: **Age:** 40Y
Onset Date: 05/10/1999 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperprolactinaemia			Change of medication

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Anxiety neurosis
900.0 Milligram Daily
Batch: **Started:** 05/10/1999 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: wt gain and hirsuitism.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 148846 **Seq:** 1 **Gender:** F
Reported: 11/02/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle contractions involuntary		Pt experienced singular muscle twitches.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient is also taking hormone replacement therapy.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 149254 **Seq:** 1 **Gender:** F
Reported: 17/02/2000 **Weight:**
Hospitalisation: **Age:** 59Y
Onset Date: 10/02/2000 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium Hyponatraemia	Severe	Hyponatraemia was associated with a Onset over 48h of acute severe	Fluid restriction, iv normal

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 07/02/2000 **Stopped:** 10/02/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium				Sodium down to 118.

Additional Information:

Patient also taking caltrate with vitamin d, atrovent, pulmicort, panadol, provera, progynova, sinemet sr, sinemet, artane and captopril.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 149612 **Seq:** 1 **Gender:** M
Reported: 25/02/2000 **Weight:** 63.00
Hospitalisation: **Age:** 80Y
Onset Date: 31/01/2000 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pruritus		Itching of skin esp legs, scratching till	Ceased aurorix.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 18/01/2000 **Stopped:** 05/02/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking vastin, lanoxin pg, lasix. - the date of onset is not necessarily accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 149627 **Seq:** 1 **Gender:** M
Reported: 25/02/2000 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gynaecomastia			

Medicine Details:

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

AURORIX (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 149634 **Seq:** 1 **Gender:** U
Reported: 25/02/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Moderately elevated lfts.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Capsule 600.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				Moderately elevated lfts.

Additional Information:

Report Detail

Case Number: 149667 **Seq:** 1 **Gender:** F
Reported: 25/02/2000 **Weight:**
Hospitalisation: Admitted to hospital **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Psychotic disorder		Patieht has a psychotic episode.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150027 **Seq:** 1 **Gender:** M
Reported: 06/03/2000 **Weight:** 79.00
Hospitalisation: **Age:** 51
Onset Date: **DOB:** 08/11/1948
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tongue disorder		Coated tongue	
Paraesthesia		Raw burning feeling of tongue	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 27/04/1999 **Stopped:** 14/01/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms returned when aurorix was recommenced.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150429 **Seq:** 1 **Gender:** M
Reported: 15/03/2000 **Weight:** 70.00
Hospitalisation: **Age:** 42
Onset Date: 10/03/2000 **DOB:** 09/02/1958
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Selltled with iv cogenytin - 1-2 minutes (1mg).

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 09/03/2000 **Stopped:** 11/03/2000

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150540 **Seq:** 1
Reported: 21/03/2000
Hospitalisation: Admitted to hospital
Onset Date: 31/12/2000
Outcome: Recovered

Gender: F
Weight:
Age: 48Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea	Severe	Patient felt ill immediately, with severe	Patient hospitalised, zoloft discontinued.
Malaise		Patient was acutely unwell.	
Diarrhoea	Severe	Severe diarrhoea.	
Serotonin syndrome			

Medicine Details:

ZOLOFT (Interaction) **Reason:** Depression
25.0 Milligram Daily
Batch: **Started:** **Stopped:** 3D

AURORIX (Interaction) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 2W

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Moclobemide was discontinued, and after a 1 week washout period, commenced zoloft 25mg/day. reactions then occurred. - the date of onset is not accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150605 **Seq:** 1 **Gender:** F
Reported: 21/03/2000 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Menstrual disorder Abdominal pain		Menstrual blood flow is lighter now	

Medicine Details:

XANAX (Suspected)		**Reason:**	
Tablet	0.0		Oral
Batch:	**Started:** 01/08/1999	**Stopped:**	CONTIN
AURORIX (Suspected)		**Reason:**	Depression
Tablet	0.0		Oral
Batch:	**Started:** 01/12/1999	**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.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150667 **Seq:** 1 **Gender:** M
Reported: 23/03/2000 **Weight:** 75.00
Hospitalisation: **Age:** 24Y
Onset Date: 15/03/2000 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Skin odour abnormal		Patient noticed a bad smell (wheat like)	Aurorix ceased.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 14/03/2000 **Stopped:** 18/03/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred within a couple of hours of taking aurorix. patient also taking rifampicin, amoxil and fusidic acid.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150727 **Seq:** 1
Reported: 23/03/2000
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: M
Weight:
Age: 60Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysphagia Extrapyramidal disorder Abdominal pain		Developed swallowing symptoms. Developed worsening movement	

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 150804 **Seq:** 1
Reported: 27/03/2000
Hospitalisation: Admitted to hospital
Onset Date: 20/12/1999
Outcome: Recovered

Gender: F
Weight: 65.00
Age: 61
DOB: 05/06/1938
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Severe		Aurorix ceased

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/07/1998 **Stopped:** 01/03/2000

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: 151331 **Seq:** 1 **Gender:** F
Reported: 06/04/2000 **Weight:**
Hospitalisation: **Age:** AD
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Speech disorder		Could not speak.	
Paresis		Pt could not move limbs at all	
Agitation			
Dizziness			
Tremor			
Visual disturbance			

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 151802 **Seq:** 1 **Gender:** F
Reported: 17/04/2000 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 16/02/2000 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hepatic function abnormal		Elevated liver enzymes.	Zyprexa ceased.

Medicine Details:

ZYPREXA (Suspected) **Reason:** Otr spec symp psychopathol nec
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 28/02/2000 12M

AURORIX (Suspected) **Reason:**
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

CAPTOPRIL (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

LUVOX (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT		16/02/2000	79	
	ALT = SGPT		13/03/2000	61	
	AST = SGOT		16/02/2000	82	
	AST = SGOT		13/03/2000	58	
	GGT = SGGT =		16/02/2000	493	
	GGT = SGGT =		13/03/2000	409	
	SAP = ALP		16/02/2000	175	

Additional Information:

Patient has just been diagnosed with diabetes and the doctor now believes this is the cause of the elevated lft's.

Report Detail

Case Number: 151802 **Seq:** 1
Reported: 17/04/2000
Hospitalisation:
Onset Date: 16/02/2000
Outcome: Not yet recovered

Gender: F
Weight:
Age: 48Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

ZYPREXA (Suspected)			**Reason:**	Otr spec symp psychopathol nec		
Tablet	10.0	Milligram	Daily		Oral	
Batch:	**Started:**			**Stopped:**	28/02/2000	12M
AURORIX (Suspected)			**Reason:**			
	600.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**		
CAPTOPRIL (Suspected)			**Reason:**			
	100.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**		
LUVOX (Suspected)			**Reason:**			
	100.0	Milligram	Daily			
Batch:	**Started:**			**Stopped:**		

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	SAP = ALP		13/03/2000	155	

Additional Information:

Patient has just been diagnosed with diabetes and the doctor now believes this is the cause of the elevated lft's.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 152193 **Seq:** 1 **Gender:** F
Reported: 01/05/2000 **Weight:**
Hospitalisation: **Age:** 72
Onset Date: 26/04/2000 **DOB:** 14/08/1927
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Hepatic function abnormal		Low frequency - jaw, face, upper & lower limbs	

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected) **Reason:**
410.0 Milligram Total
Batch: **Started:** 17/04/2000 **Stopped:** 24/04/2000

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 25/04/2000 **Stopped:** 27/04/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	ALT = SGPT AST = SGOT GGT = SGGT = SAP = ALP				Alt 10 Ast 29. Ggt 374. Alp 282.

Additional Information:

Patient also taking ciproxen, diflucan, losec, oroxine, calciparine and lasix. allergic to morphine and codeine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 152740 **Seq:** 1 **Gender:** F
Reported: 18/05/2000 **Weight:** 75.00
Hospitalisation: **Age:** 31
Onset Date: 03/04/2000 **DOB:** 24/11/1968
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Agitation Hyperkinesia Insomnia		Severe headaches for5 2-3 weeks.	Panadeine forte given, itirate aurorix dose more slowly.

Medicine Details:

AURORIX (Suspected) **Reason:**
900.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was currently taking aurorix 300mg tds and experienced symptoms, has now commenced on lower dose 3/4/00.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 152741 **Seq:** 1 **Gender:** F
Reported: 18/05/2000 **Weight:** 72.00
Hospitalisation: **Age:** 16Y
Onset Date: 30/03/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache Insomnia		Headaches for one week Insomnia for two weeks.	Panadol.

Medicine Details:

AURORIX (Suspected) **Reason:**
900.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

After commencing aurorix 450mg bd patient experienced symptoms. commenced 150mg bd on 30/3/00.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 152800 **Seq:** 1 **Gender:** M
Reported: 22/05/2000 **Weight:** 70.00
Hospitalisation: **Age:** 29Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache			Moclobemide ceased. switched to venlafaxine sr.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 16/05/2000 3W

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 152801 **Seq:** 1
Reported: 22/05/2000
Hospitalisation:
Onset Date: 16/05/2000
Outcome: Not yet recovered

Gender: M
Weight: 70.00
Age: 29Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Malaise Tremor		3-4 beat clonus.	 Maxolon 10mg ivi, cyproheptadine 8mg. cease both drugs

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** **Stopped:** 16/05/2000 3W

VENLAFAXINE HYDROCHLORIDE (Interaction) **Reason:** Depression

75.0 Milligram Daily

Batch: **Started:** 16/05/2000 **Stopped:** 16/05/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 153469 **Seq:** 1
Reported: 01/06/2000
Hospitalisation: Hospitalisation prolonged
Onset Date: 13/11/1999
Outcome: Unknown

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion		Myoclonic jerks while prone.	
Chills		Shaking uncontrollably,unable to work.	Treatment with prescription drug. congentin iv then po.
Serotonin syndrome			

Medicine Details:

EFEXOR (Interaction) **Reason:**

Tablet 37.5 Milligram 1 time Oral

Batch: **Started:** 12/11/1999 **Stopped:**

AURORIX (Interaction) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 01/07/1999 **Stopped:** 11/12/1999

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symtoms occurred 12 hours after taking a single dose of venlafaxine therapy. 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 154123 **Seq:** 1 **Gender:** F
Reported: 21/06/2000 **Weight:**
Hospitalisation: **Age:** 34
Onset Date: 01/03/1998 **DOB:** 24/06/1963
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthritis		Pain in toes, right wrist and fingers/psoriatic	Aurorix ceased

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 31/12/1997 **Stopped:** 28/12/1998

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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 154664 **Seq:** 1 **Gender:** M
Reported: 12/07/2000 **Weight:**
Hospitalisation: **Age:** 30Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Photophobia Headache		" sensitivity to light."	 Aurorix discontinued.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Symptoms occurred a few days after aurorix therapy was administered.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 154667 **Seq:** 1 **Gender:** F
Reported: 12/07/2000 **Weight:**
Hospitalisation: **Age:** 58Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urine analysis abnormal			Aurorix therapy was discontinued and levels

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Microbiology				Elevated urinary normetanephrine.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155147 **Seq:** 1 **Gender:** F
Reported: 20/07/2000 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 09/02/2000 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Back pain Extrapyramidal disorder Hyponatraemia			Moclobemide ceased. fluid

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression

Oral application 300.0 Milligram Daily Oral

Batch: **Started:** 08/02/2000 **Stopped:** 10/02/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum osmolality	270 - 290	08/02/2000	265	
	Serum osmolality	270 - 290	09/02/2000	254	
	Serum osmolality	270 - 290	10/02/2000	239	
	Serum osmolality	270 - 290	12/06/2000	271	
	Serum osmolality	270 - 290	11/02/2000	250	
	Sodium	135 - 145	08/02/2000	134	
	Sodium	135 - 145	09/02/2000	127	

Additional Information:

Previous adr's codeine - nausea, vomiting, bromocryptine -severe vomiting, sertraline - worsening of parkinson's disease. patient also taking provera, progynova, captopril, sinemet, kinson, benzhexol, atrovent, pulmicort.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155147 **Seq:** 1 **Gender:** F
Reported: 20/07/2000 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 09/02/2000 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Oral application 300.0 Milligram Daily Oral
Batch: **Started:** 08/02/2000 **Stopped:** 10/02/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium	135 - 145	10/02/2000	119	
	Sodium	135 - 145	11/02/2000	126	
	Sodium	135 - 145	12/06/2000	136	

Additional Information:

Previous adr's codeine - nausea, vomiting, bromocryptine -severe vomiting, sertraline - worsening of parkinson's disease. patient also taking provera, progynova, captopril, sinemet, kinson, benzhexol, atrovent, pulmicort.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155224 **Seq:** 1 **Gender:** M
Reported: 21/07/2000 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Thrombocytopenia			Aurorix ceased.

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Platelets				Platelets 64

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155328 **Seq:** 1 **Gender:** F
Reported: 12/07/2000 **Weight:** 72.00
Hospitalisation: **Age:** 42Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance			

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 09/09/1999 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking sorbilax, pluravit, panamax, oroxine, indocid, famotidine, diazepam, coloxyl with senna, bricanyl, beconase.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155377 **Seq:** 1
Reported: 21/06/2000
Hospitalisation:
Onset Date: 31/12/1999
Outcome: Not yet recovered

Gender: F
Weight: 63.00
Age: 37
DOB: 16/10/1962
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Alopecia		Diffuse hair loss one week after starting	Aurorix ceased.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 15/12/1999 **Stopped:** 15/01/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient had previously experienced diffuse hair loss to ssri's. - the date of onset is not necessarily accurate but indicates that onset 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 155904 **Seq:** 1
Reported: 07/08/2000
Hospitalisation:
Onset Date: 24/07/2000
Outcome: Not yet recovered

Gender: F
Weight: 65.00
Age: 66
DOB: 05/04/1934
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Insomnia Nervousness Thinking abnormal			Zantac ceased.

Medicine Details:

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

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 25/08/1999 **Stopped:** 28/07/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking dyazide, murelax and rhinocort. cns stimulation, unable to sleep, dreaming all the time, brain on overdrive nerves jumping. history: feldene - dyspnoea, macrodantin - tingling and aurorix - cns stimulation.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 156530 **Seq:** 1
Reported: 21/08/2000
Hospitalisation:
Onset Date: 20/07/2000
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 72Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Syncope Flatulence		Bad diarrhoea Fainting/falls.	Required a visit to doctor.

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/07/2000 **Stopped:** 10/08/2000

GREENRIDGE ANTI-STRESS (Interaction) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:** 6M

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Ceased and started avrorix again but whenever dose > 300mg daily patient experiences symptoms. pt has long history of irritable bowel but "not this bad".

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 158410 **Seq:** 1
Reported: 27/09/2000
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: M
Weight: 85.00
Age: 63
DOB: 22/01/1937
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pancreatitis			Required a visit to doctor. patient was hospitalised. date

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 01/02/2000 **Stopped:** 28/04/2000

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 159108 **Seq:** 1 **Gender:** F
Reported: 06/11/2000 **Weight:**
Hospitalisation: **Age:** 32
Onset Date: 21/10/2000 **DOB:** 25/09/1968
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium			
Hallucination			
Headache			
Nausea			
Palpitations			
Vertigo			

Medicine Details:

TELFAST DECONGESTANT (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 01/01/1988 **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.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 159143 **Seq:** 1 **Gender:** M
Reported: 08/11/2000 **Weight:**
Hospitalisation: **Age:** 56
Onset Date: 27/10/2000 **DOB:** 13/02/1944
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Ventricular tachycardia			Had ceased moclobemide.

Medicine Details:

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

PRAVACHOL (Suspected) **Reason:** Othr&unspec metabolic diseases
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

CARTIA (Suspected) **Reason:** Chron isch heart dis no hyper
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

DILTIAZEM HYDROCHLORIDE (Suspected) **Reason:** Chron isch heart dis no hyper
Oral application 180.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 159143 **Seq:** 1 **Gender:** M
Reported: 08/11/2000 **Weight:**
Hospitalisation: **Age:** 56
Onset Date: 27/10/2000 **DOB:** 13/02/1944
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

IRBESARTAN (Suspected)			**Reason:**	Essential benign hypertension	
Tablet	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
EFEXOR (Suspected)			**Reason:**	Depression	
	112.0	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 159299 **Seq:** 1 **Gender:** F
Reported: 16/11/2000 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 23/10/2000 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Confusional state Headache Hyponatraemia Nausea Tremor			Venlafaxime and moclobemide ceased.

Medicine Details:

VENLAFAXINE HYDROCHLORIDE (Suspected) **Reason:**
75.0 Milligram 1 time
Batch: **Started:** 23/10/2000 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 25/10/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium		25/10/2000	110	

Additional Information:

Patient also taking stemetil, nifedipine, gliclazide, thyroxine, imdur, atenol ol, aspirin, clopidogrel, lisinopril, atorvastatin.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 159583 **Seq:** 1 **Gender:** F
Reported: 29/11/2000 **Weight:** 57.00
Hospitalisation: **Age:** 17Y
Onset Date: 24/09/2000 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Chills Convulsion Diarrhoea Hyperhidrosis Serotonin syndrome Tremor			Phenergan 25mg stat im. ceased tramadol.

Medicine Details:

TRAMADOL HYDROCHLORIDE (Interaction) **Reason:** Pain
Injection 200.0 Milligram Total Intramuscular
Batch: **Started:** 24/09/2000 **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also tkaing capadex, diazepam, panadeine forte, metoclopramide. history of depression and recurrent tract infections.

Report Detail

Case Number: 159678 **Seq:** 1 **Gender:** M
Reported: 01/12/2000 **Weight:**
Hospitalisation: **Age:** 64
Onset Date: 23/11/2000 **DOB:** 05/08/1936
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pancytopenia		Pancytopenia on bone marrow.	

Medicine Details:

PRAZOSIN HYDROCHLORIDE (Other drug) **Reason:** Hyperplasia of prostate
Tablet 1.0 Milligram Daily Oral
Batch: **Started:** L TERM **Stopped:**

SOLPRIN (Other drug) **Reason:**
0.5 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** **Stopped:** 23/11/2000 3W

CELEBREX (Suspected) **Reason:** Unspecified arthritis
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:** 23/11/2000 3W

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 159978 **Seq:** 1
Reported: 13/12/2000
Hospitalisation: Admitted to hospital
Onset Date: 26/10/2000
Outcome: Unknown

Gender: F
Weight:
Age: 79
DOB: 27/09/1921
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Hyperventilation			Aurorix ceased, admitted to hospital for observation.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

450.0 Milligram Daily

Batch: **Started:** 24/10/2000 **Stopped:** 26/10/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient started on aurorix 300mg daily for 9 days and then it was increased to 450mg for 2 days. patient was also taking ranitidine, irbesartan, imdur, simvastatin, atenolol, digoxin, sandomigran, oxazepam.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160111 **Seq:** 1
Reported: 20/12/2000
Hospitalisation: Admitted to hospital
Onset Date: 18/10/2000
Outcome: Not yet recovered

Gender: F
Weight: 98.00
Age: 78
DOB: 20/11/1921
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Cerebrovascular disorder			Patient was hospitalised.

Medicine Details:

AURORIX (Suspected) **Reason:** Acute,ill-def cerebrovascular disease w/o hyperte

600.0 Milligram Daily

Batch: **Started:** 10/10/2000 **Stopped:** 18/10/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking capoten, natrilix, indocid, zantac. patient experienced symptoms 8 days after commencing aurorix.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160359 **Seq:** 1 **Gender:** M
Reported: 03/01/2001 **Weight:**
Hospitalisation: **Age:** 71
Onset Date: 05/12/2000 **DOB:** 03/05/1929
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression Suicide attempt			Zyban sr ceased.

Medicine Details:

ZYBAN SR (Suspected) **Reason:** Smoker
1.0 Dose Unspecified Daily
Batch: **Started:** 26/11/2000 **Stopped:** 05/12/2000

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** **Stopped:** 23/11/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient history: aaa repair, emphysema, coad and depression.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160764 **Seq:** 1
Reported: 24/11/2000
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Death

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Sudden death		Sudden death.	Patient was hospitalised.

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 0.0 Oral
Batch: **Started:** **Stopped:**

PETHIDINE HYDROCHLORIDE (Suspected) **Reason:**
Injection 0.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160874 **Seq:** 1 **Gender:** F
Reported: 24/01/2001 **Weight:**
Hospitalisation: **Age:** 55Y
Onset Date: 14/01/2001 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Delirium Electrolyte imbalance Hallucination Hypokalaemia Nervousness Serotonin syndrome			Hospital admission. remove antideprassants.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
0.0
Batch: **Started:** **Stopped:**

CIPRAMIL (Suspected) **Reason:** Depression
0.0
Batch: **Started:** **Stopped:**

XANAX (Suspected) **Reason:** Anxiety neurosis
0.0
Batch: **Started:** **Stopped:**

SERZONE (Suspected) **Reason:** Depression
Tablet 200.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Potassium				K dropped to 2.8 mmol/l.

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160874 **Seq:** 1
Reported: 24/01/2001
Hospitalisation:
Onset Date: 14/01/2001
Outcome: Not yet recovered

Gender: F
Weight:
Age: 55Y
DOB:
Causality: Causality possible

Reaction Detail

Medicine Details:

SOTALOL HYDROCHLORIDE (Suspected)				**Reason:**	Essential benign hypertension
Oral application	320.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
MONOPLUS NOS (Suspected)				**Reason:**	Essential benign hypertension
Tablet	1.0	Dose Unspecified	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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 160998 **Seq:** 1 **Gender:** F
Reported: 25/01/2001 **Weight:** 87.00
Hospitalisation: **Age:** 30Y
Onset Date: 20/01/2001 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea Vomiting			

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 20/01/2001 **Stopped:** 21/01/2001

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 161128 **Seq:** 1 **Gender:** F
Reported: 02/02/2001 **Weight:**
Hospitalisation: **Age:**
Onset Date: 25/01/2001 **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Angioedema Dysphagia Headache Pharyngitis Tongue oedema		Swollen mouth	Aurorix stopped.

Medicine Details:

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had similar reaction 1 month ago to same medication

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 161153 **Seq:** 1
Reported: 02/02/2001
Hospitalisation: Admitted to hospital
Onset Date: 01/11/2000
Outcome: Recovered

Gender: F
Weight:
Age: 51
DOB: 19/06/1949
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Purpura			Patient hospitalised. aurorix dose reduced to 600mg then

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 900.0 Milligram Daily Oral

Batch: **Started:** 27/09/2000 **Stopped:** 30/11/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient started aurorix 300mg bd and experienced no problems.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 161388 **Seq:** 1 **Gender:** F
Reported: 13/02/2001 **Weight:**
Hospitalisation: **Age:** 86Y
Onset Date: 20/11/2000 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Consciousness fluctuating		Unable to respond to questions, feed,	

Medicine Details:

CIPRAMIL (Suspected) **Reason:** Depression
20.0 Milligram Daily
Batch: **Started:** **Stopped:** 20/11/2000

AURORIX (Suspected) **Reason:**
150.0 Milligram Daily
Batch: **Started:** 20/11/2000 **Stopped:** 21/11/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient unable to respond to questions, feed, dress herself. did not recognise people.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 161430 **Seq:** 1
Reported: 12/02/2001
Hospitalisation: Admitted to hospital
Onset Date: 24/01/2001
Outcome: Recovered

Gender: F
Weight: 68.00
Age: 43
DOB: 22/04/1957
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Intra-uterine death			Dilantin and curettage. hospitalised for 2 nights. date of recovery 05/02/01. arima stopped

Medicine Details:

ARIMA (Suspected) **Reason:**

0.0

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient started arima. on day 1 vaginal bleeding began, day 5 miscarried 8 week old foetus. patient didnt know she was pregnant (had sterilization operation in 1998. history of 2 normal pregnances. children 13 and 15. no known allergies. history of reflux oesophagitis. patient also taking ventolin, pulmicort and ranitidine.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 162246 **Seq:** 1 **Gender:** M
Reported: 05/03/2001 **Weight:** 83.00
Hospitalisation: **Age:** 17
Onset Date: 20/01/2001 **DOB:** 03/09/1983
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nightmare			

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 162963 **Seq:** 1
Reported: 21/03/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

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

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Hypotension Syncope			

Medicine Details:

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

AURORIX (Suspected) **Reason:**
1.5 Gram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Sequelae: patient fainted and fractured her ankle.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 163077 **Seq:** 1 **Gender:** M
Reported: 23/03/2001 **Weight:** 64.00
Hospitalisation: **Age:** 52Y
Onset Date: 24/02/2001 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Mania			Aurorix ceased. required a visit to doctor.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
0.0
Batch: **Started:** 15/08/1997 **Stopped:** 12/03/2001

INTERLEUKIN 2 (Suspected) **Reason:**
0.0
Batch: **Started:** 21/02/2001 **Stopped:** 24/02/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking stavudine, lamivudine, saquinavir, ritonavir.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 163878 **Seq:** 1 **Gender:** U
Reported: 20/04/2001 **Weight:** 58.00
Hospitalisation: **Age:** 31
Onset Date: 01/04/2001 **DOB:** 14/03/1970
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dysphonia Vertigo			Date of recovery 03/04/2001.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 20/03/2001 **Stopped:** 01/04/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 164205 **Seq:** 1 **Gender:** F
Reported: 03/05/2001 **Weight:**
Hospitalisation: **Age:** 84
Onset Date: 18/09/2000 **DOB:** 28/03/1916
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			Withhold moclobemide. rehydration with n. saline.

Medicine Details:

RANITIDINE (Other drug) **Reason:**
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

ALENDRONATE SODIUM (Other drug) **Reason:**
10.0 Milligram Daily
Batch: **Started:** **Stopped:**

CALTRATE (Other drug) **Reason:**
600.0 Milligram Daily
Batch: **Started:** **Stopped:**

TIMOPTOL (Other drug) **Reason:**
Eye Drops 4.0 Dose Unspecified Daily Ophthalmic
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Sodium		06/09/2000	133	
	Sodium		16/09/2000	113	
	Sodium		18/09/2000	112	
	Sodium		18/09/2000	108	
	Sodium		19/09/2000	116	
	Sodium		20/09/2000	125	
	Sodium		29/09/2000	134	

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 164205 **Seq:** 1
Reported: 03/05/2001
Hospitalisation:
Onset Date: 18/09/2000
Outcome: Not yet recovered

Gender: F
Weight:
Age: 84
DOB: 28/03/1916
Causality: Causality possible

Reaction Detail

Medicine Details:

ALPHAGAN EYE DROPS (Other drug)			**Reason:**		
Eye Drops	4.0	Dose Unspecified	Daily		Ophthalmic
Batch:	**Started:**			**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**		
Oral application	300.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	18/09/2000

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 164446 **Seq:** 1 **Gender:** F
Reported: 11/05/2001 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 14/03/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Red hot rash on inner tighs, arms, back,	Promethazine given. moclobemide stopped.

Medicine Details:

TAMOXIFEN CITRATE (Other drug) **Reason:** Malignant neoplasm of breast
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

METOPROLOL TARTRATE (Other drug) **Reason:** Essential benign hypertension
100.0 Milligram Daily
Batch: **Started:** **Stopped:**

OMEPRAZOLE (Other drug) **Reason:** Gastritis&duodenitis
Oral application 20.0 Milligram Daily Oral
Batch: **Started:** 05/02/2001 **Stopped:**

OLANZAPINE (Other drug) **Reason:** Unspecified psychosis
5.0 Milligram Daily
Batch: **Started:** 26/02/2001 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 164446 **Seq:** 1 **Gender:** F
Reported: 11/05/2001 **Weight:**
Hospitalisation: **Age:** 69Y
Onset Date: 14/03/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

OXAZEPAM (Other drug)			**Reason:**	Otr spec symp psychopathol nec	
	120.0	Milligram	Daily		
Batch:	**Started:**	L TERM		**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
Oral application	900.0	Milligram	Daily		Oral
Batch:	**Started:**	03/03/2001		**Stopped:**	16/03/2001

Laboratory Investigations:

Additional Information:

Report Detail

Case Number: 164722 **Seq:** 1 **Gender:** F
Reported: 22/05/2001 **Weight:** 50.00
Hospitalisation: **Age:** 87
Onset Date: 12/05/2001 **DOB:** 07/07/1913
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Confusional state Somnolence		Macular truncal and limb rash.	Vioxx and arima ceased.

Medicine Details:

PANAMAX (Other drug) **Reason:**
1.0 Gram As necessary
Batch: **Started:** L TERM **Stopped:**

KENACOMB (Other drug) **Reason:**
2.0 Dose Unspecified Daily
Batch: **Started:** L TERM **Stopped:**

OXAZEPAM (Other drug) **Reason:**
45.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

VIOXX (Suspected) **Reason:** Osteoarthritis
25.0 Milligram Daily
Batch: **Started:** 30/06/2000 **Stopped:** 15/05/2001

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 164722 **Seq:** 1 **Gender:** F
Reported: 22/05/2001 **Weight:** 50.00
Hospitalisation: **Age:** 87
Onset Date: 12/05/2001 **DOB:** 07/07/1913
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

ARIMA (Suspected) **Reason:**
450.0 Milligram Daily
Batch: **Started:** 30/06/2000 **Stopped:** 15/05/2001

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 165839 **Seq:** 1
Reported: 22/06/2001
Hospitalisation: Admitted to hospital
Onset Date: 14/05/2001
Outcome: Recovered

Gender: F
Weight:
Age: 77
DOB: 23/07/1923
Causality: Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Inappropriate antidiuretic hormon			

Medicine Details:

ASPIRIN (Other drug) **Reason:** Other coagulation defects
150.0 Milligram Daily
Batch: **Started:** **Stopped:**

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

ZOTON (Other drug) **Reason:**
30.0 Milligram Daily
Batch: **Started:** **Stopped:**

ROCALTROL (Other drug) **Reason:**
0.5 Microgram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 165839 **Seq:** 1
Reported: 22/06/2001
Hospitalisation: Admitted to hospital
Onset Date: 14/05/2001
Outcome: Recovered

Gender: F
Weight:
Age: 77
DOB: 23/07/1923
Causality: Causality probable

Reaction Detail

Medicine Details:

AURORIX (Other drug)			**Reason:**	Depression
	300.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
ZYPREXA (Other drug)			**Reason:**	
	7.5	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
MOCLOBEMIDE (Suspected)			**Reason:**	
	0.0			
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 166627 **Seq:** 1 **Gender:** M
Reported: 17/07/2001 **Weight:**
Hospitalisation: **Age:** 68
Onset Date: **DOB:** 18/02/1933
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash			Seen by dermatologist. given celestone and aqueous. required a visit to doctor frusemide ceased

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/03/2001 **Stopped:** CONTIN

LISINOPRIL (Suspected) **Reason:**
10.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:** CONTIN

FRUSEMIDE (Suspected) **Reason:**
240.0 Milligram Daily
Batch: **Started:** 01/03/2001 L TERM **Stopped:** CONTIN

FENTANYL TRANSDERMAL (Suspected) **Reason:** Pain
Cream 2.5 Milligram Daily Transdermal
Batch: **Started:** 01/03/2001 **Stopped:** CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking - gliprcide, clpoidogrel, digoxin, spronolacture, fesq and ranitidine. the dose start date for the drugs moclobemide, lisinopril frusemide and fentanyl are not necessarily accurate but indicate drugs started sometime in the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 167264 **Seq:** 1
Reported: 07/08/2001
Hospitalisation:
Onset Date: 28/03/2001
Outcome: Death, maybe drug

Gender: F
Weight: 0.00
Age: 45
DOB: 14/08/1955
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug level increased			

Medicine Details:

ALCOHOL (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:** Depression
300.0 Milligram Daily Oral
Batch: **Started:** 15/10/1999 **Stopped:** 15/03/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
17/12/2002	Serum drug level				Autopsy showed a blood level of aurorix 3.8mg/kg (expected range from dose 150mg tds was reported as 0.35-1.8mg/kg) and a blood ethanol 239 mg/kg

Additional Information:

Patient had a medical history of drinking alcohol. the date of onset is not necessarily accurate but indicates onset sometime in the month.

Report Detail

Case Number: 167779 **Seq:** 1 **Gender:** M
Reported: 22/08/2001 **Weight:**
Hospitalisation: **Age:** 89Y
Onset Date: 20/08/2001 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia			

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 13/08/2001 **Stopped:** 20/08/2001

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 167803 **Seq:** 1 **Gender:** U
Reported: 23/08/2001 **Weight:**
Hospitalisation: **Age:**
Onset Date: 15/03/2001 **DOB:**
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Dry mouth Orthostatic hypotension			Required a visit to doctor.

Medicine Details:

PREMARIN (Other drug) **Reason:** Menopausal symptoms
0.6 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

PROVERA (Other drug) **Reason:** Menopausal symptoms
0.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 01/11/2000 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dose start date for the drug aurorix is not necessarily accurate but indicates the drug was started sometime in the month. aurorix was rechallenged march 2000 at 300mg/daily causing the same reaction to occur.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 168258 **Seq:** 1
Reported: 06/09/2001
Hospitalisation:
Onset Date: 16/08/2001
Outcome: Not yet recovered

Gender: F
Weight: 70.00
Age: 50
DOB: 02/10/1950
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Arthralgia Urticaria		Affecting hand and elbow joints	Patient required to visit a doctor; aurorix and voltaren

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 13/08/2001 **Stopped:** 31/08/2001

VOLTAREN (Suspected) **Reason:**

100.0 Milligram Daily

Batch: **Started:** 13/08/2001 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 168759 **Seq:** 1
Reported: 27/09/2001
Hospitalisation:
Onset Date: 28/04/2001
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 56
DOB: 08/12/1944
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyperhidrosis		Severe swaeting in the middle of the night.	Required a visit to doctor.

Medicine Details:

INDERAL (Other drug) **Reason:**
60.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

QUININE BISULPHATE (Other drug) **Reason:** Pain in limb
300.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

CALTRATE (Other drug) **Reason:**
1.2 Gram Daily
Batch: **Started:** L TERM **Stopped:**

ROCALTROL (Other drug) **Reason:**
0.2 Microgram Daily
Batch: **Started:** L TERM **Stopped:**

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 is not necessarily accurate but indicates the drug was sta rted sometime in the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 168759 **Seq:** 1
Reported: 27/09/2001
Hospitalisation:
Onset Date: 28/04/2001
Outcome: Not yet recovered

Gender: F
Weight: 53.00
Age: 56
DOB: 08/12/1944
Causality: Causality possible

Reaction Detail

Medicine Details:

OROXINE (Other drug)			**Reason:**	
	100.0 Microgram		Daily	
Batch:	**Started:**	L TERM		**Stopped:**
ARIMA (Suspected)			**Reason:**	Depression
	750.0 Milligram		Daily	
Batch:	**Started:** 01/01/1999			**Stopped:**

Laboratory Investigations:

Additional Information:

The date of onset is not necessarily accurate but indicates onset sometime in the month. the dose start date is not necessarily accurate but indicates the drug was sta rted sometime in the month.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 169321 **Seq:** 1
Reported: 22/10/2001
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: U
Weight:
Age:
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coma Therapeutic response increased			Patient hospitalised

Medicine Details:

STOCRIN (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Serum drug level				Very high levels of a efavirenz.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 170028 **Seq:** 1 **Gender:** M
Reported: 19/11/2001 **Weight:** 64.00
Hospitalisation: **Age:** 51
Onset Date: 13/10/2001 **DOB:** 27/12/1949
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash Diarrhoea		Trunk, inner upper arms, groin, lower	Date of recovery 1.11.01. required a visit to doctor.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 12/10/2001 **Stopped:** 22/10/2001

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 170253 **Seq:** 1 **Gender:** F
Reported: 23/11/2001 **Weight:**
Hospitalisation: **Age:** 77Y
Onset Date: 31/05/2001 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		Developed red, urticarial rash.	Treated with hydrocortisone 1% cream. moclobemide ceased.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 29/05/2001 **Stopped:** 31/05/2001

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 170806 **Seq:** 1 **Gender:** F
Reported: 03/12/2001 **Weight:** 56.00
Hospitalisation: **Age:** 81
Onset Date: 15/10/2001 **DOB:** 22/06/1920
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria			Given promethazine. required a visit to doctor.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 16/07/2001 **Stopped:** 19/10/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 170821 **Seq:** 1 **Gender:** F
Reported: 03/12/2001 **Weight:** 53.00
Hospitalisation: **Age:** 14
Onset Date: 03/03/2001 **DOB:** 20/04/1986
Outcome: Unknown **Causality:** Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Malaise Nausea		General feeling of malaise.	Changed to moclobemide.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Unspecified leukemia

Oral application 450.0 Milligram Daily Oral

Batch: **Started:** 03/03/2001 **Stopped:** 23/04/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pateint experienced symptoms with each dose of moclobemide

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 172555 **Seq:** 1 **Gender:** M
Reported: 11/02/2002 **Weight:** 70.00
Hospitalisation: **Age:** 35Y
Onset Date: 28/01/2002 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Drug withdrawal syndrome Nausea			Required a visit to doctor. symptoms stopped approx. 1 hour after taking moclobemide again

Medicine Details:

BACTRIM (Other drug) **Reason:** Prophylaxis
0.0
Batch: **Started:** **Stopped:**

COMBIVIR (Other drug) **Reason:** Hiv infection
0.0
Batch: **Started:** **Stopped:**

NEVIRAPINE (Other drug) **Reason:** Hiv infection
0.0
Batch: **Started:** **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 01/05/2001 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Within 24 hours of ceasing moclobemide patient experienced symptoms the dose of onset is not necessarily accurate but indicates onset sometime in the month. the dose start date is not necessarily accurate but indicates moclobemide was stared sometime in the month.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 173257 **Seq:** 1 **Gender:** F
Reported: 18/02/2002 **Weight:** 64.00
Hospitalisation: **Age:** 47
Onset Date: 30/01/2002 **DOB:** 20/05/1954
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Neuralgia		Pain left temple	Date of recovery 12/02/02.

Medicine Details:

VIOXX (Other drug) **Reason:** Otr&unsp vertebrogen pain synd
25.0 Milligram Daily
Batch: **Started:** **Stopped:**

LOSEC (Other drug) **Reason:** Otr disorders stomach function
20.0 Milligram Daily
Batch: **Started:** **Stopped:**

TRAMAL SR (Other drug) **Reason:** Lumbalgia
200.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
150.0 Milligram Daily
Batch: **Started:** 29/01/2002 **Stopped:** 11/02/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 173403 **Seq:** 1 **Gender:** F
Reported: 06/03/2002 **Weight:**
Hospitalisation: **Age:** 63
Onset Date: 10/02/2002 **DOB:** 18/05/1938
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tetany		Jaw spasm	
Convulsion		Myoclonic jerks	Diazepam. ceased citalopram.
Hypertonia		Stiffness	
Serotonin syndrome			

Medicine Details:

CITALOPRAM HYDROBROMIDE (Suspected) **Reason:** Depression
Tablet 20.0 Milligram Daily Oral
Batch: **Started:** 10/02/2002 **Stopped:** 10/02/2002

GALANTAMINE HYDROBROMIDE (Suspected) **Reason:**
Tablet 8.0 Milligram Daily Oral
Batch: **Started:** 21/01/2002 **Stopped:** 11/02/2002

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 900.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has early alzheimer. the diagnostic criteria are not fulfilled.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 173618 **Seq:** 1 **Gender:** M
Reported: 14/03/2002 **Weight:**
Hospitalisation: **Age:** 70
Onset Date: 21/01/2002 **DOB:** 19/11/1931
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea			Reduced dose aurorix while on doxycycline, resumed previous dose day after ceasing doxycycline

Medicine Details:

INZA (Other drug) **Reason:** Osteoarthritis
750.0 Milligram Daily
Batch: **Started:** 01/01/2000 **Stopped:**

SOLPRIN (Other drug) **Reason:** Transient cereb isch no hypert
150.0 Milligram Daily
Batch: **Started:** 01/01/2000 **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 01/01/2001 **Stopped:**

DOXYCYCLINE HYDROCHLORIDE (Suspected) **Reason:** Prophylaxis
Oral application 100.0 Milligram Daily Oral
Batch: **Started:** 21/01/2000 **Stopped:** 07/03/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dose start date is not necessarily accurate but indicates aurorix was started sometime in the year. patient recovered 8/3/02.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 173879 **Seq:** 1
Reported: 25/03/2002
Hospitalisation:
Onset Date: 17/03/2002
Outcome: Not yet recovered

Gender: F
Weight: 80.00
Age: 28
DOB: 24/01/1974
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting		Vomiting ++.	Required a visit to doctor. cease aurorix.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
300.0 Milligram Daily
Batch: **Started:** 17/03/2002 **Stopped:** 21/03/2002

FEFOL (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/03/2002 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 174189 **Seq:** 1
Reported: 03/04/2002
Hospitalisation:
Onset Date: 31/12/2001
Outcome: Recovered

Gender: F
Weight: 60.00
Age: 34
DOB: 12/10/1967
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amenorrhoea			Aurorix ceased. date of recovery 27.3.02. required a

Medicine Details:

AURORIX (Suspected) **Reason:**
1.2 Gram Daily
Batch: **Started:** 01/01/2001 **Stopped:** 01/01/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Prolactin	<500.			Serum prolactin at 839.

Additional Information:

The dose start and stop dates are not accurate but indicates the drug was started and stopped sometime in the year.

Report Detail

Case Number: 174330 **Seq:** 1 **Gender:** F
Reported: 09/04/2002 **Weight:** 59.00
Hospitalisation: **Age:** 34
Onset Date: 03/04/2002 **DOB:** 09/11/1967
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depersonalisation Insomnia Nausea		Felt out of body.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** 03/04/2002 **Stopped:** 04/04/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 174554 **Seq:** 1
Reported: 12/04/2002
Hospitalisation:
Onset Date: 28/03/2002
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 34
DOB: 12/10/1967
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Galactorrhoea Hyperprolactinaemia			Aurorix ceased. required a visit to doctor.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

150.0 Milligram Daily

Batch: **Started:** 01/01/2000 **Stopped:** 01/01/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The dose start and stop dates are not accurate but indicates the drug was started and stopped sometime in the year. sequelae: delayed fertility.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 174763 **Seq:** 1
Reported: 22/04/2002
Hospitalisation:
Onset Date: 12/03/2002
Outcome: Not yet recovered

Gender: F
Weight:
Age: 22
DOB: 15/06/1979
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Asthenia Bronchospasm Dyspnoea Pruritus		Sluggishness.	Tapering off aurorix over 2 weeks. required a visit to doctor

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

2.0 Dose Unspecified Daily

Batch: **Started:** 05/03/2002 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Exercise - induced asthma occuring with much less exertion.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 175000 **Seq:** 1
Reported: 01/05/2002
Hospitalisation: Admitted to hospital
Onset Date: 16/04/2002
Outcome: Not yet recovered

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

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Pancytopenia		White cells, red cells & platelets all [illegible]	Patient was hospitalised. discontinued both moclobemide [illegible]

Medicine Details:

VITAMINORUM B.P.C. (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

ASCORBIC ACID (Other drug) **Reason:**
500.0 Milligram Daily
Batch: **Started:** **Stopped:**

PRAZOSIN HYDROCHLORIDE (Other drug) **Reason:**
2.0 Milligram Daily
Batch: **Started:** **Stopped:**

NILSTAT (Other drug) **Reason:**
4.0 Millilitre Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No previous history of allergy.

Report Detail

Case Number: 175000 **Seq:** 1
Reported: 01/05/2002
Hospitalisation: Admitted to hospital
Onset Date: 16/04/2002
Outcome: Not yet recovered

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

Reaction Detail

Medicine Details:

THIAMINE (Other drug)			**Reason:**	
	100.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
MOCLOBEMIDE (Suspected)			**Reason:**	
	600.0	Milligram	Daily	
Batch:	**Started:**			**Stopped:**
RISPERIDONE (Suspected)			**Reason:**	
	2.0	Milligram	Daily	
Batch:	**Started:**	16/04/2002		**Stopped:** 19/04/2002

Laboratory Investigations:

Additional Information:

No previous history of allergy.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 175001 **Seq:** 1 **Gender:** M
Reported: 01/05/2002 **Weight:**
Hospitalisation: **Age:** 51Y
Onset Date: 15/04/2002 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state			Ceased tramal sr and aurorix and started avanza instead.

Medicine Details:

AURORIX (Interaction) **Reason:** Depression
0.0
Batch: **Started:** L TERM **Stopped:** 17/04/2002

TRAMAL SR (Interaction) **Reason:** Pain
300.0 Milligram Daily
Batch: **Started:** 13/04/2002 **Stopped:** 17/04/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 175118 **Seq:** 1
Reported: 06/05/2002
Hospitalisation: Admitted to hospital
Onset Date: 07/05/2001
Outcome: Recovered

Gender: F
Weight:
Age: 78
DOB: 31/05/1922
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Coma Orthostatic hypotension Syncope			Stopped moclobamide and changed to venlafaxine sr. verapamil held temporarily. admitted to hospital

Medicine Details:

CYANOCOBALAMIN (Other drug) **Reason:**
1.0 Milligram Monthly
Batch: **Started:** L TERM **Stopped:**

VERAPAMIL HYDROCHLORIDE (Other drug) **Reason:**
160.0 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

FOLIC ACID (Other drug) **Reason:**
0.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

ZOPICLONE (Other drug) **Reason:**
7.5 Milligram Daily
Batch: **Started:** L TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient history - parkinson's diseased; ?alzheimers; depression/anxiety; hypertension; tachycardia and pernicious anaemia.

Report Detail

Case Number: 175118 **Seq:** 1
Reported: 06/05/2002
Hospitalisation: Admitted to hospital
Onset Date: 07/05/2001
Outcome: Recovered

Gender: F
Weight:
Age: 78
DOB: 31/05/1922
Causality: Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
	300.0	Milligram	Daily		
Batch:	**Started:**			**Stopped:**	3D
MADOPAR HBS (Suspected)			**Reason:**	Paralysis agitans	
	2.0	Dose Unspecified	Daily		
Batch:	**Started:**			**Stopped:**	

Laboratory Investigations:

Additional Information:

Patient history - parkinson's diseased; ?alzheimers; depression/anxiety; hypertension; tachycardia and pernicious anaemia.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 175962 **Seq:** 1
Reported: 06/06/2002
Hospitalisation:
Onset Date: 01/06/2002
Outcome: Not yet recovered

Gender: F
Weight: 60.00
Age: 39
DOB: 05/06/1962
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Insomnia Nausea			Required a visit to doctor.

Medicine Details:

MICROGYNON 20 ED (Other drug) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** 01/01/1999 **Stopped:** CONTIN

AURORIX (Suspected) **Reason:**
300.0 Milligram 1 time
Batch: **Started:** 31/05/2002 **Stopped:** 31/05/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Myoclonic jerks increased to several per minute day and night after the aurorix (one only) 300mg, with vocal grunting.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 176025 **Seq:** 1
Reported: 07/06/2002
Hospitalisation: Admitted to hospital
Onset Date: 16/05/2002
Outcome: Recovered

Gender: M
Weight:
Age: A5
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Muscle contractions involuntary Blood pressure fluctuation Abdominal pain Agitation Confusional state Dizziness Hyperhidrosis Serotonin syndrome Tremor		Clonus. Unstable blood pressure.	Aurorix and zoloft ceased. pateint was hospitalised. date of recovery 17/05/02.

Medicine Details:

AURORIX (Interaction) **Reason:**
0.0
Batch: **Started:** **Stopped:**

ZOLOFT (Interaction) **Reason:** Depression
Tablet 25.0 Milligram Daily Oral
Batch: **Started:** 16/05/2002 **Stopped:** 17/05/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Also taking long list of herbal preparations (not specified). auorix ceased and the next day the patient was commenced on zoloft.

Report Detail

Case Number: 176025 **Seq:** 1
Reported: 07/06/2002
Hospitalisation: Admitted to hospital
Onset Date: 16/05/2002
Outcome: Recovered

Gender: M
Weight:
Age: A5
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Vomiting			

Medicine Details:

AURORIX (Interaction) **Reason:**

0.0

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

ZOLOFT (Interaction) **Reason:** Depression

Tablet 25.0 Milligram Daily Oral

Batch: **Started:** 16/05/2002 **Stopped:** 17/05/2002

Laboratory Investigations:

Additional Information:

Also taking long list of herbal preparations (not specified). auorix ceased and the next day the patient was commenced on zoloft.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 176293 **Seq:** 1 **Gender:** F
Reported: 14/06/2002 **Weight:**
Hospitalisation: **Age:** 15Y
Onset Date: 15/10/2000 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Serotonin syndrome Suicide attempt			Patient treated with

Medicine Details:

EFEXOR (Suspected) **Reason:**
750.0 Milligram Total
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of depression and past overdoses. serotonin toxicity characterised by sustained ankle clonus, ocular clonus, tachycardia, low-grade fever, hyperreflexia, tremor, myoclonic spasms and agitation.

Report Detail

Case Number: 176949 **Seq:** 1
Reported: 26/06/2002
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Recovered

Gender: F
Weight:
Age: 87Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diplopia Hallucination			Patient was hospitalised.

Medicine Details:

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

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

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

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient recently had a cataract operation.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 178973 **Seq:** 1
Reported: 12/09/2002
Hospitalisation: Admitted to hospital
Onset Date: 31/07/2002
Outcome: Recovered

Gender: M
Weight:
Age: 41
DOB: 16/08/1960
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation			
Confusional state			
Diarrhoea			
Dyspnoea			
Hypertonia			
Myalgia			Admitted to hospital.
Tremor			

Medicine Details:

AVANZA (Interaction) **Reason:** Depression

Tablet 30.0 Milligram Daily Oral

Batch: **Started:** 26/05/2002 **Stopped:** 29/07/2002

AURORIX (Interaction) **Reason:** Depression

Tablet 450.0 Milligram Daily Oral

Batch: **Started:** 31/07/2002 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Serotonin syndrome.

Report Detail

Case Number: 180284 **Seq:** 1 **Gender:** M
Reported: 04/11/2002 **Weight:**
Hospitalisation: **Age:** 86
Onset Date: 29/10/2002 **DOB:** 11/11/1915
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Rash		Rash face and neck.	Required a visit to doctor. hydrocortisone 1% cream bd.

Medicine Details:

ATROVENT (Other drug) **Reason:**
Inhalation 500.0 Microgram As necessary Inhalation
Batch: **Started:** 10/10/2002 **Stopped:**

VENTOLIN (Other drug) **Reason:**
Inhalation 5.0 Milligram As necessary Inhalation
Batch: **Started:** 10/10/2002 **Stopped:**

COLOXYL WITH SENNA (Other drug) **Reason:** Constipation
Tablet 4.0 Dose Unspecified Daily Oral
Batch: **Started:** 10/10/2002 **Stopped:**

BETAMIN (Suspected) **Reason:**
100.0 Milligram Daily
Batch: **Started:** 13/10/2002 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 180284 **Seq:** 1
Reported: 04/11/2002
Hospitalisation:
Onset Date: 29/10/2002
Outcome: Not yet recovered

Gender: M
Weight:
Age: 86
DOB: 11/11/1915
Causality: Causality possible

Reaction Detail

Medicine Details:

VITAMINORUM B.P.C. (Suspected)				**Reason:**		
Tablet	2.0	Dose Unspecified	Daily		Oral	
Batch:	**Started:**	18/10/2002		**Stopped:**		
SERETIDE ACCUHALER NOS (Suspected)				**Reason:**		
Inhalation	2.0	Dose Unspecified	Daily		Inhalation	
Batch:	**Started:**	23/10/2002		**Stopped:**		
PANAFCORTELONE (Suspected)				**Reason:**	Pneumonia,unspecified	
Tablet	5.0	Milligram	Daily		Oral	
Batch:	**Started:**	12/10/2002		**Stopped:**	27/10/2002	
ARIMA (Suspected)				**Reason:**	Depression	
Tablet	300.0	Milligram	Daily		Oral	
Batch:	**Started:**	20/10/2002		**Stopped:**	30/10/2002	

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 180284 **Seq:** 1
Reported: 04/11/2002
Hospitalisation:
Onset Date: 29/10/2002
Outcome: Not yet recovered

Gender: M
Weight:
Age: 86
DOB: 11/11/1915
Causality: Causality possible

Reaction Detail

Medicine Details:

ACIMAX (Suspected)			**Reason:**		
Tablet	40.0	Milligram	Daily		Oral
Batch:	**Started:**	23/10/2002		**Stopped:**	
CALCIPARINE (Suspected)			**Reason:**		
Injection	10.0	Thousand Internat	Daily		Subcutaneous
Batch:	**Started:**	21/10/2002		**Stopped:**	29/10/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 180710 **Seq:** 1 **Gender:** F
Reported: 17/12/2002 **Weight:** 77.00
Hospitalisation: **Age:** 49
Onset Date: 09/11/2002 **DOB:** 10/11/1952
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea Lethargy		Nausea and extreme lethargy	

Medicine Details:

HYDRENE 25/50 (Other drug) **Reason:** Essential benign hypertension
Batch: **Started:** **Stopped:**

CATAPRES (Other drug) **Reason:** Migraine
300.0 Milligram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** 08/11/2002 **Stopped:** 09/11/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181213 **Seq:** 1 **Gender:** F
Reported: 08/01/2003 **Weight:** 0.00
Hospitalisation: **Age:** 34Y
Onset Date: 04/04/2002 **DOB:**
Outcome: Recovered 12/04/2002 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state Delusion		Confusion and Delusional state	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 04/04/2002 **Stopped:** 10/04/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181247 **Seq:** 1 **Gender:** U
Reported: 09/01/2003 **Weight:** 0.00
Hospitalisation: **Age:** 55
Onset Date: **DOB:** 14/12/1947
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Glossodynia	Required Visit to Doctor	With higher dose of Moclobemide from 300mg/bd to 375mg/bd gradual over weeks of burning sensation of tongue	

Medicine Details:

ATENOLOL (Other drug) **Reason:** Essential benign hypertension

Tablet	50.0	Milligram	Daily	Oral

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

MOCLOBEMIDE (Suspected) **Reason:** Depression

Tablet	750.0	Milligram	Daily	Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Dose start and stop dates for the drug Moclobemide - March 2002 to May 2002. Date of onset - 04/02.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181324 **Seq:** 1
Reported: 09/01/2003
Hospitalisation:
Onset Date: 01/11/2002
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 55
DOB: 21/10/1947
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Testicular pain		Testicular pain	Diclofenac 50mg TID. Aurorix and Hiforc for men were

Medicine Details:

HIFORCE FOR MEN (Suspected) **Reason:**

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

AURORIX (Suspected) **Reason:** Depression

Tablet 150.0 Milligram Daily Oral

Batch: **Started:** 25/10/2002 **Stopped:** 04/11/2002

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181350 **Seq:** 1 **Gender:** M
Reported: 10/01/2003 **Weight:** 0.00
Hospitalisation: **Age:** 44
Onset Date: 26/10/2002 **DOB:** 16/12/1957
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Nausea	Caused or prolonged inpatient hospitalisation	Developed nausea, diarrhoea, dizziness shortness of breath, thight chest.	oxygen, was put on drip, aspirin
Flushing	Caused or prolonged inpatient hospitalisation	Possible serotonin reaction, felt warm, faint	
Syncope	Caused or prolonged inpatient hospitalisation	Sweating bradycardia of 30 BPM, heart rate of 70 BPM .	

Medicine Details:

AURORIX (Interaction) **Reason:**
Tablet 300.0 Milligram Oral
Batch: **Started:** **Stopped:** 22/10/2002 0

ZOLOFT (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 26/10/2002 **Stopped:** 26/10/2002

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient cycled 45.5km

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181350 **Seq:** 1 **Gender:** M
Reported: 10/01/2003 **Weight:** 0.00
Hospitalisation: **Age:** 44
Onset Date: 26/10/2002 **DOB:** 16/12/1957
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bradycardia	Caused or prolonged inpatient hospitalisation		
Chest discomfort	Caused or prolonged inpatient hospitalisation		
Chest pain	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AURORIX (Interaction) **Reason:**
Tablet 300.0 Milligram Oral
Batch: **Started:** **Stopped:** 22/10/2002 0

ZOLOFT (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 26/10/2002 **Stopped:** 26/10/2002

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

Laboratory Investigations:

Additional Information:

Patient cycled 45.5km

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 181350 **Seq:** 1 **Gender:** M
Reported: 10/01/2003 **Weight:** 0.00
Hospitalisation: **Age:** 44
Onset Date: 26/10/2002 **DOB:** 16/12/1957
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Caused or prolonged inpatient hospitalisation		
Dizziness	Caused or prolonged inpatient hospitalisation		
Hyperhidrosis	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AURORIX (Interaction) **Reason:**
Tablet 300.0 Milligram Oral
Batch: **Started:** **Stopped:** 22/10/2002 0

ZOLOFT (Interaction) **Reason:**
Tablet 50.0 Milligram Daily Oral
Batch: **Started:** 26/10/2002 **Stopped:** 26/10/2002

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

Laboratory Investigations:

Additional Information:

Patient cycled 45.5km

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 182379 **Seq:** 1 **Gender:** U
Reported: 05/02/2003 **Weight:** 0.00
Hospitalisation: **Age:** 99U
Onset Date: 14/01/2003 **DOB:**
Outcome: Recovered 19/01/2003 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agranulocytosis Diarrhoea	Severe Severe	Agranulocytosis and mild diarrhoea.	Moclobemide ceased.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
05/02/2003	Neutrophils		14/01/2003	0.0	

Additional Information:

Patient also taking Digoxin, Seretide, Salbutamol and Aspirin.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 182423 **Seq:** 1 **Gender:** M
Reported: 06/02/2003 **Weight:** 0.00
Hospitalisation: **Age:** 45
Onset Date: 22/09/2002 **DOB:** 18/10/1956
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia		Compulsive cleaning lasted hours. Claimed amnesia after the event	
Obsessive-compulsive disorder		Compulsive cleaning lasted hours. Claimed amnesia after the event	

Medicine Details:

STILNOX (Interaction) **Reason:** Specific disorders of sleep
15.0 Milligram Daily
Batch: **Started:** 22/09/2002 **Stopped:**

ARIMA (Interaction) **Reason:** Depression
450.0 Milligram Daily
Batch: **Started:** 06/09/2002 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Stilnox without antidepressant had been OK. Patient also taking valpro, quinine, zantac, zocor,murelax

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 182521 **Seq:** 1 **Gender:** F
Reported: 07/02/2003 **Weight:** 60.00
Hospitalisation: **Age:** 48
Onset Date: 17/10/2002 **DOB:** 07/06/1954
Outcome: Recovered 18/10/2002 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Depression		Echinacea appeared to counteract the patient's moclobemide therapy, symptoms of depression returned when patient commenced herbal tratment for symptoms of	Echinacea ceased recovered within few hours.
Drug ineffective		Echinacea appeared to counteract the patient's moclobemide therapy, symptoms of depression returned when patient commenced herbal tratment for symptoms of	

Medicine Details:

CENOVIS ECHINACEA 4500 (Suspected) **Reason:** Acute nasopharyngitis,com cold
Capsule 200.0 Milligram Daily Oral
Batch: **Started:** 12/10/2002 **Stopped:** 17/10/2002

ARIMA (Suspected) **Reason:** Depression
600.0 Milligram Daily
Batch: **Started:** L TERM **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 183025 **Seq:** 1 **Gender:** M
Reported: 14/02/2003 **Weight:** 0.00
Hospitalisation: **Age:** 41
Onset Date: **DOB:** 21/07/1961
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug ineffective	Required Visit to Doctor	Recurrences of depression	Resume Aurorix.

Medicine Details:

ARIMA (Suspected) **Reason:** Convulsions
120.0 Milligram Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred sometime in Dec 2002.
Arima was started sometime in December 2002.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 183791 **Seq:** 1 **Gender:** M
Reported: 23/12/2002 **Weight:** 85.00
Hospitalisation: **Age:** 99U
Onset Date: 01/11/2002 **DOB:**
Outcome: Recovered 13/11/2002 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Amnesia Thinking abnormal		Sense of being "lost" - would forget ahat he was doing there, por prioritisation	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 03/10/2002 **Stopped:** 08/11/2002

MINOMYCIN (Suspected) **Reason:** Other acne of sebaceous glands
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** 03/10/2002 **Stopped:** 08/11/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous similar effect secondary to Zoloft. 181709 is a duplicate.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 184786 **Seq:** 1 **Gender:** M
Reported: 08/04/2003 **Weight:** 0.00
Hospitalisation: **Age:** 78Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia		Hyponatraemia (SIADH).	

Medicine Details:

MOCLOBEMIDE (Suspected) **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 185715 **Seq:** 1
Reported: 09/05/2003
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: M
Weight: 0.00
Age: 99U
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Blood pressure increased		Patient commenced Aropax the next day after discontinuing Moclobemide and experienced an elevation in blood pressure from 160/90 to 180/90.	

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

AROPAX (Interaction) **Reason:**
Daily Oral
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 186945 **Seq:** 1
Reported: 17/06/2003
Hospitalisation: Admitted to hospital
Onset Date: 24/04/2003
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 26
DOB: 02/07/1976
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urinary retention	Required Visit to Doctor	Urinary retention - unable to urinate.	Aurorix stopped

Medicine Details:

AMOXYCILLIN TRIHYDRATE (Other drug) **Reason:** Acute apical periodontitis
1.5 Gram Daily
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Batch: **Started:** 08/11/2002 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 189137 **Seq:** 1 **Gender:** M
Reported: 31/07/2003 **Weight:** 70.00
Hospitalisation: **Age:** 75
Onset Date: 14/07/2003 **DOB:** 13/11/1927
Outcome: Recovered 29/07/2003 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest pain	Caused or prolonged inpatient hospitalisation		
Dizziness	Caused or prolonged inpatient hospitalisation		
Glucose tolerance impaired	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis

Tablet 900.0 Milligram Daily Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Lopid, Avapro, Quinate, Temazepam.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 189137 **Seq:** 1 **Gender:** M
Reported: 31/07/2003 **Weight:** 70.00
Hospitalisation: **Age:** 75
Onset Date: 14/07/2003 **DOB:** 13/11/1927
Outcome: Recovered 29/07/2003 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Headache	Caused or prolonged inpatient hospitalisation		
Tremor	Caused or prolonged inpatient hospitalisation		
Serotonin syndrome	Caused or prolonged inpatient hospitalisation	Serotonin syndrome, headache, chest pain, dizzy, tremors, cough, impaired glucose tolerance.	

Medicine Details:

AURORIX (Suspected) **Reason:** Anxiety neurosis

Tablet 900.0 Milligram Daily Oral

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

Laboratory Investigations:

Additional Information:

Patient was also taking Lopid, Avapro, Quinate, Temazepam.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 190767 **Seq:** 1 **Gender:** F
Reported: 23/09/2003 **Weight:** 0.00
Hospitalisation: **Age:** 84
Onset Date: 08/06/2003 **DOB:** 05/12/1918
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Confusional state	Caused or prolonged inpatient hospitalisation	Confusion, agitation, serotonin syndrome, bilateral gross chorea	ICU admission. Implicated medications ceased.
Rhabdomyolysis	Caused or prolonged inpatient hospitalisation	Rhabdomyolysis, acute renal failure, after changing from moclobemide to citalopram without washout period or possibly both taken together	
Agitation	Caused or prolonged inpatient hospitalisation		

Medicine Details:

CAPTOPRIL (Interaction) **Reason:**
Tablet 50.0 Milligram 3 times Oral
Batch: **Started:** **Stopped:** 08/06/2003

CELEBREX (Interaction) **Reason:**
Capsule 200.0 Milligram 2 times Oral
Batch: **Started:** **Stopped:** 08/06/2003

AURORIX (Interaction) **Reason:**
300.0 Milligram 2 times Oral
Batch: **Started:** **Stopped:**

CITALOPRAM HYDROBROMIDE (Interaction) **Reason:**
Tablet 20.0 Milligram 1 time Oral
Batch: **Started:** 22/05/2003 **Stopped:** 08/06/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
23/09/2003	Albumin				Creatinine Kinase 950u/L, serum creatinine 318 micromol/L.

Additional Information:

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 190767 **Seq:** 1 **Gender:** F
Reported: 23/09/2003 **Weight:** 0.00
Hospitalisation: **Age:** 84
Onset Date: 08/06/2003 **DOB:** 05/12/1918
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chorea	Caused or prolonged inpatient hospitalisation		
Renal failure acute	Caused or prolonged inpatient hospitalisation		

Medicine Details:

CAPTOPRIL (Interaction) **Reason:**
Tablet 50.0 Milligram 3 times Oral
Batch: **Started:** **Stopped:** 08/06/2003

CELEBREX (Interaction) **Reason:**
Capsule 200.0 Milligram 2 times Oral
Batch: **Started:** **Stopped:** 08/06/2003

AURORIX (Interaction) **Reason:**
300.0 Milligram 2 times Oral
Batch: **Started:** **Stopped:**

CITALOPRAM HYDROBROMIDE (Interaction) **Reason:**
Tablet 20.0 Milligram 1 time Oral
Batch: **Started:** 22/05/2003 **Stopped:** 08/06/2003

Laboratory Investigations:

Additional Information:

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 191071 **Seq:** 1 **Gender:** F
Reported: 02/10/2003 **Weight:** 0.00
Hospitalisation: **Age:** 66
Onset Date: **DOB:** 06/12/1936
Outcome: Unknown **Causality:** Causality possible

Reaction Detai

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Therapeutic response unexpecte		Patient experienced unexpected orgasm.	Nil.

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 2.0 Dose Unspecified Daily Oral
Batch: **Started:** 15/06/2002 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Date of onset of reaction 7/2002.
Aurorix ceased 7/2002.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 192088 **Seq:** 1
Reported: 11/11/2003
Hospitalisation:
Onset Date: 30/09/2003
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 24
DOB: 11/01/1979
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tremor Headache Hyperreflexia Muscle spasms Nausea Trismus Vision blurred		Patient developed nausea, spasm in leg, headache, also tremor in right leg/ frontal headache/ right eye blur vision/ clenchy jaw	

Medicine Details:

MOCLOBEMIDE (Interaction) **Reason:** Depression
1.2 Gram Daily
Batch: **Started:** 30/05/2003 **Stopped:** 30/09/2003

VENLAFAXINE HYDROCHLORIDE (Interaction) **Reason:** Depression
37.5 Milligram Daily
Batch: **Started:** 30/09/2003 **Stopped:**

DIAZEPAM (Other drug) **Reason:**
1.0 Dose Unspecified As necessary
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 192793 **Seq:** 1 **Gender:** F
Reported: 03/12/2003 **Weight:** 0.00
Hospitalisation: **Age:** 99U
Onset Date: 27/10/2003 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug withdrawal syndrome neona Neonatal disorder		Drug withdrawal syndrome. Neonatal disorder NOS - abnormality.	

Medicine Details:

RISPERDAL (Suspected) **Reason:**
Tablet 4.0 Milligram Daily Intrauterine
Batch: **Started:** **Stopped:**

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

MOCLOBEMIDE (Suspected) **Reason:**
1.0 Dose Unspecified Daily
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also administered: Marcain and Fentanyl.
Baby was exposed from 20+ weeks. The baby experienced withdrawl symptoms (from risperidone) for 2 days, on days 3 and 4, however, did not require medication to control it.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 193459 **Seq:** 1 **Gender:** F
Reported: 22/12/2003 **Weight:** 0.00
Hospitalisation: **Age:** 39
Onset Date: **DOB:** 06/03/1964
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Chest discomfort Arteriospasm coronary Dyspnoea Hyperhidrosis Nausea Palpitations		Chest heaviness, shortness of breath, palpitations, nausea, sweatiness and coronary spasm.	Morphine, betablockers and a GTN infusion.

Medicine Details:

IMIGRAN (Suspected) **Reason:**
Dose Unspecified
Batch: **Started:** L TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 15/10/2003 **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occured twice - Oct 2003 and Dec 2003.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 193619 **Seq:** 1 **Gender:** F
Reported: 06/01/2004 **Weight:** 68.00
Hospitalisation: **Age:** 30
Onset Date: 19/12/2003 **DOB:** 12/05/1973
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
White blood cell count decreased	Required Visit to Doctor	WCC decreased.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 27/11/2003 **Stopped:** 19/12/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/01/2004	White blood cells		19/12/2003	2.0	
06/01/2004	White blood cells		22/12/2003	4.4	

Additional Information:

No previous exposure to this drug. No other explanation for decreased WCC.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 193832 **Seq:** 1
Reported: 12/01/2004
Hospitalisation: Admitted to hospital
Onset Date:
Outcome: Unknown

Gender: F
Weight: 0.00
Age: 76Y
DOB:
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia	Caused or prolonged inpatient hospitalisation	Hyponatraemia	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 600.0 Milligram Daily Oral

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other drugs: Verapamil, Atorvastatin, Tamoxifen and Prochlorperazine. There was insufficient information to determine the outcome of the hyponatraemia. History: Depression.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 196053 **Seq:** 1 **Gender:** M
Reported: 29/03/2004 **Weight:** 85.00
Hospitalisation: **Age:** 21
Onset Date: 28/02/2004 **DOB:** 22/11/1982
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus	Required Visit to Doctor	Patient developed ringing in ears, stomach pains, dizzy.	Moclobemide ceased.
Abdominal pain upper	Required Visit to Doctor		
Dizziness	Required Visit to Doctor		

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 25/02/2004 S TERM **Stopped:** 2 weeks

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 196427 **Seq:** 1 **Gender:** F
Reported: 13/04/2004 **Weight:** 0.00
Hospitalisation: **Age:** 48
Onset Date: **DOB:**
Outcome: Death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug toxicity Completed suicide Death		Overdose of moclobemide of acute	Flumazenil administered by the ambulance staff during attempted resuscitation.

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
13/04/2004	Serum drug level				

Additional Information:

A long term history of depression and had attempted suicide on nine previous occasions. An autopsy revealed a large volume of tablet residue in her stomach (236gm) pulmonary oedema and mild lymphocytic thyroiditis. Other drug therapy diazepam, trifluoroperazine. Death was considered to be due to acute poisoning by moclobemide alone. Camaris C and Little D. A fatality due to moclobemide. J Forensic SCI 1997, 242 (5): 854 - 955

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 196467 **Seq:** 1 **Gender:** M
Reported: 14/04/2004 **Weight:** 0.00
Hospitalisation: **Age:** 22
Onset Date: **DOB:**
Outcome: Death **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Drug interaction Death Drug toxicity Intentional overdose		Drug interaction, drug toxicity due to intentional overdose, death.	Given Charcoal.

Medicine Details:

MOCLOBEMIDE (Interaction)		**Reason:**	
Batch:	**Started:**		**Stopped:**
SERTRALINE HYDROCHLORIDE (Interaction)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PIMOZIDE (Interaction)		**Reason:**	
Batch:	**Started:**		**Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
14/04/2004	Biochemistry				
14/04/2004	Haematology				
14/04/2004	Other data				

Additional Information:

J.Forensic Science 1997, 42(5): 951-953.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 197522 **Seq:** 1 **Gender:** F
Reported: 21/05/2004 **Weight:** 0.00
Hospitalisation: **Age:** 49Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Fear Insomnia Restlessness		Patient became over-stimulated, restless, frightened and could not sleep.	Aurorix ceased.

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 750.0 Milligram Daily Oral

Batch: **Started:** S TERM **Stopped:** 03/05/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was started on 300mg twice daily and did not experience any problems. Involved significant disability or incapacity.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 199111 **Seq:** 1
Reported: 19/07/2004
Hospitalisation: Admitted to hospital
Onset Date: 13/07/2004
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 57
DOB: 26/09/1946
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Medication error	Caused or prolonged inpatient hospitalisation	Inadvertant concomitant use of Venlafaxine SR and Moclobemide, low blood pressure 90/60, hypotension, twitching of limbs, lethargy and loss of interest.	Venlafaxine and Moclobemide ceased by patient.
Decreased interest	Caused or prolonged inpatient hospitalisation		
Hypotension	Caused or prolonged inpatient hospitalisation		

Medicine Details:

VENLAFAXINE HYDROCHLORIDE (Suspected) **Reason:**
Dose Unspecified
Batch: **Started:** **Stopped:** 11 days

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 02/07/2004 **Stopped:** 13/07/2004 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Moclobemide dose was 300mg daily for 4 days then increased to 300mg bd. Other drugs taken: Serenace, Haloperidol, Effexor XR, Simvastatin, Cogentin, Tritace, Dilatrend and Cartia. Medical history: Brain injury 10 years ago, MI, CABG x 4 on day of infarct, depression, psych = paranoya, IHD and severe Liver dysfunction. Inadvertant concomitant use of Venlafaxine SR + Moclobemide for 11 days. Venlafaxine was meant to be ceased by patient .

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 199111 **Seq:** 1
Reported: 19/07/2004
Hospitalisation: Admitted to hospital
Onset Date: 13/07/2004
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 57
DOB: 26/09/1946
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Lethargy	Caused or prolonged inpatient hospitalisation		
Muscle twitching	Caused or prolonged inpatient hospitalisation		
Serotonin syndrome	Caused or prolonged inpatient hospitalisation		

Medicine Details:

VENLAFAXINE HYDROCHLORIDE (Suspected) **Reason:**
Dose Unspecified
Batch: **Started:** **Stopped:** 11 days

MOCLOBEMIDE (Suspected) **Reason:**
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** 02/07/2004 **Stopped:** 13/07/2004 0

Laboratory Investigations:

Additional Information:

Moclobemide dose was 300mg daily for 4 days then increased to 300mg bd. Other drugs taken: Serenace, Haloperidol, Effexor XR, Simvastatin, Cogentin, Tritace, Dilatrend and Cartia. Medical history: Brain injury 10 years ago, MI, CABG x 4 on day of infarct, depression, psych = paranoya, IHD and severe Liver dysfunction. Inadvertant concomitant use of Venlafaxine SR + Moclobemide for 11 days. Venlafaxine was meant to be ceased by patient .

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 199589 **Seq:** 1 **Gender:** M
Reported: 02/08/2004 **Weight:** 0.00
Hospitalisation: **Age:** 42
Onset Date: 17/06/2004 **DOB:** 23/08/1961
Outcome: Recovered 09/07/2004 **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Haematuria	Required Visit to Doctor	Patient developed haematuria.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 300.0 Milligram Daily
Batch: **Started:** 17/09/2003 L TERM **Stopped:** 02/07/2004 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 200560 **Seq:** 1 **Gender:** F
Reported: 06/09/2004 **Weight:** 0.00
Hospitalisation: **Age:** 29
Onset Date: 20/08/2004 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Gravitational oedema		Swelling above and below the ankle bone and on top of the feet-on both legs.	

Medicine Details:

NORIMIN (Other drug) **Reason:** Disturbance of sleep
Tablet 1.0 Dose Unspecified Daily Oral
Batch: **Started:** 15/07/2003 L TERM **Stopped:**

XANAX (Other drug) **Reason:**
Tablet 200.0 Microgram Daily Oral
Batch: **Started:** 15/11/2003 S TERM **Stopped:**

AURORIX (Suspected) **Reason:** Depression
Tablet 375.0 Milligram Daily Oral
Batch: **Started:** 13/08/2004 S TERM **Stopped:** 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 200562 **Seq:** 1 **Gender:** M
Reported: 06/09/2004 **Weight:** 0.00
Hospitalisation: **Age:** 70Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Oedema peripheral Joint swelling Pitting oedema		Dependent bilateral pitting oedema.	Elevation and compression hosiery, withdrawal of Moclobemide

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 300.0 Milligram Daily Oral
Batch: **Started:** S TERM **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** S TERM **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Article from 'The Medical Journal of Australia, Vol 157, July 20, 1992'.

Report Detail

Case Number: 201037 **Seq:** 1 **Gender:** F
Reported: 22/09/2004 **Weight:** 54.00
Hospitalisation: **Age:** 82
Onset Date: 25/08/2004 **DOB:** 31/01/1922
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Myoclonus		Within 30 minutes developed myoclonic	Ceased Tramal.

Medicine Details:

MORPHINE SULPHATE (Interaction) **Reason:** Pain in chest
Injection 5.0 Milligram 1 time Intravenous
Batch: **Started:** **Stopped:**

ARIMA (Interaction) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

TRAMAL (Interaction) **Reason:** Pain
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 06/07/2004 **Stopped:** 25/08/2004

CARDIPRIN 100 (Other drug) **Reason:**
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previously had been exposed to Morphine shile also taking Meclobomide without problems. Combination of Moclobomide and Tramadol seemed OK but addition of Morphine started sequence of events.

Report Detail

Case Number: 201037 **Seq:** 1 **Gender:** F
Reported: 22/09/2004 **Weight:** 54.00
Hospitalisation: **Age:** 82
Onset Date: 25/08/2004 **DOB:** 31/01/1922
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

COVERSYL (Other drug)			**Reason:**	
Tablet	4.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**
DURIDE (Other drug)			**Reason:**	
Tablet	120.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**
OROXINE (Other drug)			**Reason:**	
Tablet	100.0	Microgram	Daily	Oral
Batch:	**Started:**			**Stopped:**
SOLONE (Other drug)			**Reason:**	
Tablet	5.0	Milligram	Daily	Oral
Batch:	**Started:**			**Stopped:**

Laboratory Investigations:

Additional Information:

Previously had been exposed to Morphine shile also taking Meclobomide without problems. Combination of Moclobomide and Tramadol seemed OK but addition of Morphine started sequence of events.

Report Detail

Case Number: 201037 **Seq:** 1 **Gender:** F
Reported: 22/09/2004 **Weight:** 54.00
Hospitalisation: **Age:** 82
Onset Date: 25/08/2004 **DOB:** 31/01/1922
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Medicine Details:

RULIDE (Other drug) **Reason:**

Tablet 300.0 Milligram Daily Oral

Batch: **Started:** 25/08/2004 **Stopped:**

Laboratory Investigations:

Additional Information:

Previously had been exposed to Morphine shile also taking Meclobomide without problems. Combination of Moclobomide and Tramadol seemed OK but addition of Morphine started sequence of events.

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 201683 **Seq:** 1 **Gender:** F
Reported: 13/10/2004 **Weight:** 0.00
Hospitalisation: **Age:** 62
Onset Date: 04/09/2004 **DOB:** 30/08/1942
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fatigue Somnolence		Two weeks after commencing oral Moclobemide (300mg twice a day), patient experienced fatigue and drowsiness, particularly in the afternoons. One afternoon patient hit a parked car whilst driving as a result of these events.	Patient reduced the dose to 150mg twice a day and the events subsequently resolved.

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 21/08/2004 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 204275 **Seq:** 1
Reported: 17/01/2005
Hospitalisation:
Onset Date: 15/10/2004
Outcome: Not yet recovered

Gender: F
Weight: 0.00
Age: 48
DOB: 27/01/1956
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Tinnitus		Tinnitus.	

Medicine Details:

AURORIX (Suspected) **Reason:** Depression

Tablet 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 204664 **Seq:** 1 **Gender:** F
Reported: 27/01/2005 **Weight:** 0.00
Hospitalisation: **Age:** 68
Onset Date: 23/11/2004 **DOB:** 11/10/1936
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hyponatraemia Dizziness		The patient experiencecd hyponatraemia, dizziness and insomnia.	Fluid restriction.

Medicine Details:

TIOTROPIUM (Other drug) **Reason:**
Inhalation 1.0 Dose Unspecified Inhalation
Batch: **Started:** **Stopped:**

ALENDRONATE SODIUM (Other drug) **Reason:**
Tablet 70.0 Milligram Oral
Batch: **Started:** **Stopped:**

Zolpidem Sandoz (Other drug) **Reason:** Specific disorders of sleep
Tablet 10.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

METOPROLOL TARTRATE (Other drug) **Reason:** Essential benign hypertension
Tablet 100.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
27/01/2005	Sodium				Na 116 on admission, Na 131 at discharge.

Additional Information:

Report Detail

Case Number: 204664 **Seq:** 1 **Gender:** F
Reported: 27/01/2005 **Weight:** 0.00
Hospitalisation: **Age:** 68
Onset Date: 23/11/2004 **DOB:** 11/10/1936
Outcome: Recovered **Causality:** Causality probable

Reaction Detail

Medicine Details:

GLICLAZIDE (Other drug)			**Reason:**	Diabetes mellitus	
Tablet	30.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	
PANCRELIPASE (Other drug)			**Reason:**	Chronic pancreatitis	
Injection	3.0	Dose Unspecified	Daily		Intravenous
Batch:	**Started:**			**Stopped:**	
MOCLOBEMIDE (Suspected)			**Reason:**	Depression	
Tablet	150.0	Milligram	Daily		Oral
Batch:	**Started:**			**Stopped:**	23/11/2004

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 204683 **Seq:** 1 **Gender:** M
Reported: 27/01/2005 **Weight:** 0.00
Hospitalisation: **Age:** 47
Onset Date: 09/01/2005 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Dizziness Nausea		The patient experienced dizziness and nausea approximately 30 minutes post dose	Mohexal ceased, replaced with Aurorix.

Medicine Details:

CELEBREX (Other drug) **Reason:**
Capsule 200.0 Milligram Oral
Batch: **Started:** **Stopped:**

Mohexal (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 05/01/2005 **Stopped:** 13/01/2005

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 204685 **Seq:** 1 **Gender:** F
Reported: 27/01/2005 **Weight:** 0.00
Hospitalisation: **Age:** 47
Onset Date: 09/01/2005 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Visual disturbance		The patient's vision occasionally blacks out and she has fluctuating moods from day to	Mohexal ceased, replaced by

Medicine Details:

Mohexal (Suspected) **Reason:** Depression
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 05/01/2005 **Stopped:** 13/01/2005

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 205920 **Seq:** 1 **Gender:** F
Reported: 10/03/2005 **Weight:** 0.00
Hospitalisation: **Age:** 6Y
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Breast mass		The child developed breast lumps.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

FRISIUM (Suspected) **Reason:**
Tablet Oral
Batch: **Started:** **Stopped:**

EPILIM (Suspected) **Reason:** Other&unspecified epilepsy
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 206380 **Seq:** 1
Reported: 31/03/2005
Hospitalisation:
Onset Date: 15/12/2004
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 34
DOB: 13/05/1970
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anaemia macrocytic Macrocytosis		The patient developed macrocytosis and macrocytic anaemia.	Moclobemide ceased.

Medicine Details:

AURORIX (Suspected) **Reason:**

Tablet 600.0 Milligram Daily Oral

Batch: **Started:** 12/11/2004 **Stopped:** 30/12/2004

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 206907 **Seq:** 1 **Gender:** M
Reported: 11/04/2005 **Weight:** 0.00
Hospitalisation: **Age:** 51
Onset Date: 07/04/2005 **DOB:** 13/10/1953
Outcome: Recovered with sequelae 11/04/2005 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Circulatory collapse	Life	Patient has been taking venlafaxine XR 300 mg mane, for 3 years. Inadvertently consumed moclobemide 300 mg tablets (2 or 3, according to patient). Noted to collapse and experience clonus approximately 20 minutes later. Taken to hospital, where required intubation and ICU	Intubation and ventilation in ICU for 24 hours, supportive care. Cessation of moclobemide. Withheld venlafaxine for four days.
Clonus	Life		
Hypoxia	Life		
Medication error	Life		

Medicine Details:

EFEXOR-XR (Interaction) **Reason:**
Capsule 300.0 Milligram 1 time Oral
Batch: **Started:** 01/04/2002 **Stopped:** 0

MOCLOBEMIDE (Interaction) **Reason:**
Tablet 600.0 Milligram 1 time Oral
Batch: **Started:** 07/04/2005 **Stopped:** 07/04/2005 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 206968 **Seq:** 1 **Gender:** F
Reported: 13/04/2005 **Weight:** 52.00
Hospitalisation: **Age:** 99u
Onset Date: 10/03/2005 **DOB:**
Outcome: Recovered 17/03/2005 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Fall	Life	The patient fell, striking her head on the toilet bowl, which resulted in a left orbital	Surgical repair 17/3/05.

Medicine Details:

STILNOX (Suspected) **Reason:** Specific disorders of sleep
Tablet 10.0 Milligram Oral
Batch: **Started:** **Stopped:** 0

ALPRAZOLAM (Suspected) **Reason:** Anxiety neurosis
Tablet 250.0 Microgram As necessary Oral
Batch: **Started:** **Stopped:**

ADALAT OROS (Suspected) **Reason:**
Tablet 60.0 Milligram Daily Oral
Batch: **Started:** 08/03/2005 **Stopped:**

MOCLOBEMIDE (Suspected) **Reason:** Depression
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** 07/03/2005 **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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 208856 **Seq:** 1 **Gender:** F
Reported: 21/06/2005 **Weight:** 0.00
Hospitalisation: **Age:** 34Y
Onset Date: 19/05/2005 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Convulsion Drug withdrawal syndrome		Patient experienced a seizure ten to eleven days after stopping Xanax therapy.	Xanax ceased.

Medicine Details:

XANAX (Suspected)		**Reason:**	
Tablet			Oral
Batch:	**Started:**	**Stopped:**	
AURORIX (Suspected)		**Reason:**	
Tablet			Oral
Batch:	**Started:**	**Stopped:**	
SEROQUEL (Suspected)		**Reason:**	
Tablet			Oral
Batch:	**Started:**	**Stopped:**	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

This was the second in three years under same circumstances. Both occured within 7-10 days of abrupt cessation of Xanax

Report Detail

Case Number: 211780 **Seq:** 1 **Gender:** M
Reported: 20/09/2005 **Weight:** 0.00
Hospitalisation: **Age:** 60
Onset Date: 15/06/2005 **DOB:** 20/03/1945
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Loss of consciousness Circulatory collapse		Patient started oral Moclobemide (600mg daily) in January 2004 and had been taking it for about 18 months when he "blacked out" whilst drinking at a party. When he collapsed he experienced a few lacerations to the arm and forehead and was "blacked out" for about an hour. He was unable to walk.	

Medicine Details:

AVAPRO (Other drug) **Reason:**
Tablet Oral
Batch: **Started:** 15/10/1999 L TERM **Stopped:**

LIPITOR (Other drug) **Reason:**
Tablet Oral
Batch: **Started:** 15/07/2003 L TERM **Stopped:**

LOSEC (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

AURORIX (Suspected) **Reason:**
Tablet 600.0 Milligram Daily Oral
Batch: **Started:** 15/01/2004 L TERM **Stopped:** Continuing.

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

The patient had received Moclobemide in the past and had an unknown adverse reaction.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 214228 **Seq:** 1 **Gender:** F
Reported: 12/12/2005 **Weight:** 80.00
Hospitalisation: **Age:** 99U
Onset Date: 15/11/2005 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Agitation Insomnia		Patient experienced agitation and	Ceased Moclobemide.

Medicine Details:

COVERSYL PLUS (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

GLUCOSAMINE SULPHATE (Other drug) **Reason:**
Batch: **Started:** **Stopped:**

SOMAC (Other drug) **Reason:** Gastritis&duodenitis
Batch: **Started:** **Stopped:** 0

TAMOXIFEN CITRATE (Other drug) **Reason:** Malignant neoplasm of breast
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 214228 **Seq:** 1 **Gender:** F
Reported: 12/12/2005 **Weight:** 80.00
Hospitalisation: **Age:** 99U
Onset Date: 15/11/2005 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Medicine Details:

MOCLOBEMIDE (Suspected) **Reason:** Malignant neoplasm of breast

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 215154 **Seq:** 1 **Gender:** F
Reported: 23/01/2006 **Weight:** 0.00
Hospitalisation: **Age:** 99u
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Urticaria		The patient developed minor itchy, dry skin approximately a month after starting Moclobemide 300mg. The dose was increased to 600mg and the patient developed an itchy hive rash.	Moclobemide ceased.

Medicine Details:

AURORIX (Suspected) **Reason:**
300.0 Milligram Daily 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 215968 **Seq:** 1 **Gender:** F
Reported: 17/02/2006 **Weight:** 65.00
Hospitalisation: **Age:** 45
Onset Date: 05/01/2006 **DOB:** 18/07/1960
Outcome: Recovered 12/01/2006 **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Psychotic disorder Drug interaction Hallucination		Within 12 days of first dose, onset of psychosis and hallucinations.	Ceased Aurorix and Somac.

Medicine Details:

AURORIX (Interaction) **Reason:** Unspecifid affective psychosis
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

SOMAC (Interaction) **Reason:** Other diseases of esophagus
Tablet 40.0 Milligram Daily Oral
Batch: **Started:** 04/01/2006 **Stopped:** 08/01/2006

LITHICARB (Other drug) **Reason:** Unspecifid affective psychosis
Batch: **Started:** **Stopped:**

COUMADIN (Other drug) **Reason:** Otr venous embolism&thrombosis
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Report Detail

Case Number: 215968 **Seq:** 1 **Gender:** F
Reported: 17/02/2006 **Weight:** 65.00
Hospitalisation: **Age:** 45
Onset Date: 05/01/2006 **DOB:** 18/07/1960
Outcome: Recovered 12/01/2006 **Causality:** Causality possible

Reaction Detail

Medicine Details:

CALTRATE (Other drug)		**Reason:**	
Batch:	**Started:**		**Stopped:**
PANADOL (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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 218430 **Seq:** 1
Reported: 17/05/2006
Hospitalisation: Required a visit to the doctor
Onset Date: 05/05/2006
Outcome: Not yet recovered

Gender: F
Weight: 108.00
Age: 31
DOB: 23/07/1974
Causality: Causality certain

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Hypoglycaemia		Patient uses Insulin pump with Novorapid, commenced Moclobemide and became severely hypoglycaemia.	Dose adjustment.

Medicine Details:

NOVORAPID (Interaction) **Reason:** Diabetes mellitus
Batch: **Started:** 19/04/2004 **Stopped:**

MOCLOBEMIDE (Interaction) **Reason:** Depression
Tablet 450.0 Milligram Daily Oral
Batch: **Started:** 05/05/2006 **Stopped:**

METFORMIN HYDROCHLORIDE (Interaction) **Reason:** Diabetes mellitus
Tablet 2.0 Gram Daily Oral
Batch: **Started:** **Stopped:**

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Suspect drug certain rechallenge.

THERAPEUTIC GOODS ADMINISTRATION
Public Case Detail

Date Range: 1 Jan 1989 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 220766 **Seq:** 1 **Gender:** U
Reported: 14/08/2006 **Weight:** 0.00
Hospitalisation: **Age:** 99u
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Renal impairment		Suspected. A patient who experienced	

Medicine Details:

AURORIX (Suspected) **Reason:**

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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 222095 **Seq:** 1
Reported: 29/09/2006
Hospitalisation:
Onset Date: 15/06/2002
Outcome: Not yet recovered

Gender: M
Weight: 0.00
Age: 53
DOB: 13/11/1948
Causality: Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Anosmia	Incapacity/disability	A patient experienced permanent bilateral anosmia after taking Aurorix.	

Medicine Details:

AURORIX (Suspected) **Reason:**
Tablet 60.0 Milligram Oral
Batch: **Started:** 15/01/2002 **Stopped:** 15/03/2002 0

VALIUM (Suspected) **Reason:**
5.0 Milligram
Batch: **Started:** 15/01/2002 **Stopped:** 15/03/2002 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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 224080 **Seq:** 1 **Gender:** F
Reported: 14/12/2006 **Weight:** 0.00
Hospitalisation: **Age:** 34
Onset Date: **DOB:** 03/04/1972
Outcome: Recovered **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Bradycardia Muscular weakness Nausea Somnolence Vomiting		10 minutes after taking Imigran for migraine, developed bradycardia, severe nausea/vomiting, generalised muscle weakness like anaesthetic, would then go to sleep, 2 to 3 hours later wake with symptom free - migraine gone.	

Medicine Details:

IMIGRAN (Interaction) **Reason:** Migraine
Tablet 50.0 Milligram As necessary Oral
Batch: **Started:** **Stopped:**

AURORIX (Interaction) **Reason:**
Tablet 150.0 Milligram Daily Oral
Batch: **Started:** **Stopped:**

DUPHASTON (Other drug) **Reason:**
2.0 Milligram
Batch: **Started:** **Stopped:**

PROGYNOVA (Other drug) **Reason:**
2.0 Milligram
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: ARIMA,AURORIX,MOCLOBEMIDE,Mohexal

Report Detail

Case Number: 227261 **Seq:** 1 **Gender:** M
Reported: 03/04/2007 **Weight:** 0.00
Hospitalisation: **Age:** 99u
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Detail

Preferred Term	Severity	Report Description	Treatment
Suicidal behaviour	Caused or prolonged inpatient hospitalisation		

Medicine Details:

AURORIX (Suspected) **Reason:**
1200.0 Milligram
Batch: **Started:** **Stopped:**

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

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information: