



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 55702

Seq: 1

Gender: F

Reported: 26/08/1988

Weight: 72.00

Hospitalisation:

Age: 33Y

Onset Date: 26/08/1988

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Affect lability Dyspnoea			Phenergan 12.5mg given, complete resolution within 3 minutes. reaction occurred 5mins after droperidol given and 50mins after omnopon.
Tongue discolouration Tongue oedema			

Medicine Details:

DROPERIDOL (Suspected)	Reason:		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 26/08/1988	Stopped:	
OMNOPON (Suspected)	Reason: Premedication		
Injection	20.0 Milligram	1 time	Intramuscular
Batch:	Started: 26/08/1988	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 56929

Seq: 1

Gender: F

Reported: 13/10/1988

Weight: 57.00

Hospitalisation:

Age: 52Y

Onset Date: 15/08/1988

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Neuroleptic malignant syndrome			

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection, intravenous infusion	10.0 Milligram	Total	Intravenous
Batch:	Started: 14/08/1988	Stopped:	15/08/1988

PETHIDINE HYDROCHLORIDE (Suspected)		Reason:	
Injection, intravenous infusion	500.0 Milligram	Total	Intravenous
Batch:	Started: 14/08/1988	Stopped:	15/08/1988

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 58196

Seq: 1

Gender: F

Reported: 25/01/1989

Weight: 57.00

Hospitalisation:

Age: 34Y

Onset Date: 23/01/1989

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Cogentin & valium ivi

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 23/01/1989	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 59253 **Seq:** 1 **Gender:** F
Reported: 17/03/1989 **Weight:**
Hospitalisation: **Age:** 49Y
Onset Date: 14/03/1989 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dysarthria Dystonia			2mg cogentin

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	75.0 Milligram Daily Intravenous
Batch:	Started: 14/03/1989 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 59524 **Seq:** 1 **Gender:** F
Reported: 29/03/1989 **Weight:**
Hospitalisation: **Age:** 29Y
Onset Date: 13/02/1989 **DOB:**
Outcome: Recovered **Causality:** Causality certain

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Anaphylactoid reaction			Adrenaline 1:1000 1ml sc, hydrocortisone 100mg im, promethazine 25mg im

Medicine Details:

DIPRIVAN (Other drug)	Reason: Other disturbance of sensation
Injection 220.0 Milligram Total Intravenous	
Batch: Started: 13/02/1989 Stopped:	
FENTANYL CITRATE (Other drug)	Reason: Other disturbance of sensation
Injection 50.0 Microgram Total Intravenous	
Batch: Started: 13/02/1989 Stopped:	
SYNTOCINON (Other drug)	Reason:
Injection 10.0 International Unit Total Intravenous	
Batch: Started: 13/02/1989 Stopped:	
DROLEPTAN (Suspected)	Reason: Other disturbance of sensation
Injection 2.5 Milligram Total Intravenous	
Batch: Started: 13/02/1989 Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Immunology				Skin testing performed on 29/3/89 positive for droperidol (droleptan)

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 61376

Seq: 1

Gender: F

Reported: 13/07/1989

Weight: 50.00

Hospitalisation:

Age: 57Y

Onset Date: 05/07/1989

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dry mouth Dystonia Speech disorder			l.m.i. cogentin

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	5.0 Milligram	Daily	Intramuscular
Batch:	Started: 04/07/1989	Stopped: 05/07/1989	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 62150	Seq: 1	Gender: M
Reported: 28/08/1989		Weight:
Hospitalisation:		Age: 53Y
Onset Date: 23/08/1989		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Cogentin iv/valium

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Tablet	30.0 Milligram	Daily	Oral
Batch:	Started: 22/08/1989	Stopped:	23/08/1989

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



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Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 62472 **Seq:** 1 **Gender:** M
Reported: 31/08/1989 **Weight:**
Hospitalisation: **Age:** 53Y
Onset Date: 23/08/1989 **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			

Medicine Details:

DROPERIDOL (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: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 64103	Seq: 1	Gender: M
Reported: 25/10/1989		Weight:
Hospitalisation:		Age: 58Y
Onset Date: 26/09/1989		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Abdominal pain			
Blood amylase increased			
Jaundice			

Medicine Details:

DIPRIVAN (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started: 26/09/1989	Stopped:
FENTANYL CITRATE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started: 26/09/1989	Stopped:
DROPERIDOL (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started: 26/09/1989	Stopped:
ISOFLURANE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started: 26/09/1989	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry				30/8/89 26/9/89 27/9/89 28/9/89 16/10/89 ref range bili: 23 90 32 28 15 (0-20 umol/l) alk phos: 75 68 56 60 48 (35-115 u/l) ld: 176 718 572 467 252 (120-250 u/l) ast: 21 61 41 31 29 (0-40 u/l)
	Misclassified at				Amylase: - 283 - 77 101 (20-100 u/l)

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 64316

Seq: 1

Gender: F

Reported: 08/02/1990

Weight: 51.00

Hospitalisation:

Age: 21Y

Onset Date: 07/02/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine 2 mg iv stat.

Medicine Details:

DROPERIDOL (Suspected)	Reason: Otr intestne&peritoneum dsease
Injection	15.0 Milligram Total Intramuscular
Batch:	Started: 06/02/1990 Stopped: 07/02/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 64696

Seq: 1

Gender: F

Reported: 02/03/1990

Weight: 56.00

Hospitalisation:

Age: 34Y

Onset Date: 02/03/1990

DOB:

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	10.0 Milligram	As necessary	Intramuscular
Batch:	Started: 01/03/1990	Stopped: 02/03/1990	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65043	Seq: 1	Gender: F
Reported: 04/04/1990		Weight:
Hospitalisation:		Age: 46Y
Onset Date: 03/04/1990		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Pruritus			Ceased omnopon infusion. pethidine 100 mg im was used.
Rash			

Medicine Details:

OMNOPON (Suspected)		Reason: Other disturbance of sensation	
Injection, intravenous infusion	100.0 Millilitre	Total	Intravenous
Batch:	Started: 03/04/1990	Stopped:	04/04/1990
THIOPENTONE SODIUM (Suspected)		Reason:	
	300.0 Milligram	Daily	
Batch:	Started:	Stopped:	
VECURONIUM BROMIDE (Suspected)		Reason:	
	6.0 Milligram	Daily	
Batch:	Started:	Stopped:	
MORPHINE NOS (Suspected)		Reason:	
	5.0 Milligram	Daily	
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65043

Seq: 1

Gender: F

Reported: 04/04/1990

Weight:

Hospitalisation:

Age: 46Y

Onset Date: 03/04/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason:

4.0 Milligram

Daily

Batch:

Started:

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65534

Seq: 1

Gender: M

Reported: 17/05/1990

Weight: 60.00

Hospitalisation:

Age: 18Y

Onset Date: 16/05/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			lv benztropine 2mg

Medicine Details:

PANADEINE (Other drug)	Reason: Pain
	2.0 Dose Unspecified As necessary
Batch:	Started: 14/05/1990 Stopped:
FLOXAPEN (Other drug)	Reason: Otr&nos infec¶sit diseases
Injection	4.0 Gram Daily Intravenous
Batch:	Started: 14/05/1990 Stopped:
PENICILLIN NOS (Other drug)	Reason: Otr&nos infec¶sit diseases
Injection	4.0 Dose Unspecified Daily Intravenous
Batch:	Started: 14/05/1990 Stopped:
DROLEPTAN (Suspected)	Reason: Nausea and vomiting
Injection	10.0 Milligram Daily Intravenous
Batch:	Started: 15/05/1990 Stopped: 16/05/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65534

Seq: 1

Gender: M

Reported: 17/05/1990

Weight: 60.00

Hospitalisation:

Age: 18Y

Onset Date: 16/05/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

STEMETIL (Suspected)		Reason: Nausea and vomiting	
Injection	50.0 Milligram	Daily	Intramuscular
Batch:	Started: 15/05/1990	Stopped:	

MAXOLON (Suspected)		Reason: Nausea and vomiting	
Injection	40.0 Milligram	Daily	Intramuscular
Batch:	Started: 15/05/1990	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65964	Seq: 1	Gender: F
Reported: 07/06/1990		Weight: 59.00
Hospitalisation:		Age: 35Y
Onset Date:		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hot flush			
Palpitations			
Tremor			

Medicine Details:

TEMAZEPAM (Suspected)	Reason: Anxiety neurosis
1.0 Dose Unspecified As necessary	
Batch:	Started: 04/06/1990 Stopped:
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation
Injection 10.0 Milligram As necessary Intramuscular	
Batch:	Started: 04/06/1990 Stopped:
PANADEINE (Suspected)	Reason: Other disturbance of sensation
2.0 Dose Unspecified As necessary	
Batch:	Started: 04/06/1990 Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection 1.0 Milligram As necessary Intravenous	
Batch:	Started: 05/06/1990 Stopped: 05/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 65964

Seq: 1

Gender: F

Reported: 07/06/1990

Weight: 59.00

Hospitalisation:

Age: 35Y

Onset Date:

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

HEPARIN SODIUM (Suspected)

Reason: Other coagulation defects

Injection

10.0 Thousand Internal Daily

Subcutaneous

Batch:

Started: 04/06/1990

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 66411	Seq: 1	Gender: F
Reported: 27/06/1990		Weight: 57.00
Hospitalisation:		Age: 24Y
Onset Date: 26/06/1990		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Myalgia			
Oculogyration			Benztropine img iv

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	4.0 Milligram	Total	Intravenous
Batch:	Started: 25/06/1990	Stopped:	26/06/1990

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 67715	Seq: 1	Gender: F
Reported: 02/10/1990		Weight: 60.00
Hospitalisation:		Age: 45Y
Onset Date: 25/06/1990		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Injection site reaction			Dmsso, vitamin e cream.

Medicine Details:

EPIRUBICIN HYDROCHLORIDE (Suspected)		Reason: Malignant neoplasm of ovary
Injection	100.0 Milligram	1 time Intravenous
Batch:	Started: 25/06/1990	Stopped:
CYCLOPHOSPHAMIDE (Suspected)		Reason: Malignant neoplasm of ovary
Injection	1.2 Gram	1 time Intravenous
Batch:	Started: 25/06/1990	Stopped:
DEXAMETHASONE (Suspected)		Reason: Malignant neoplasm of ovary
Injection	10.0 Milligram	1 time Intravenous
Batch:	Started: 25/06/1990	Stopped:
PROCHLORPERAZINE MALEATE (Suspected)		Reason: Malignant neoplasm of ovary
Injection	12.0 Milligram	1 time Intravenous
Batch:	Started: 25/06/1990	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 67715

Seq: 1

Gender: F

Reported: 02/10/1990

Weight: 60.00

Hospitalisation:

Age: 45Y

Onset Date: 25/06/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason: Malignant neoplasm of ovary

Injection

10.0 Milligram

1 time

Intravenous

Batch:

Started: 25/06/1990

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69071 **Seq:** 1 **Gender:** M
Reported: 09/01/1991 **Weight:**
Hospitalisation: **Age:** 6M
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Cogentin 0.02mg/kg 1gm

Medicine Details:

DROLEPTAN (Suspected)	Reason: Premedication		
Injection	1.0 Milligram	Daily	Intramuscular
Batch:	Started:	Stopped:	
MORPHINE NOS (Suspected)	Reason: Premedication		
Injection	1.0 Milligram	Daily	Intramuscular
Batch:	Started:	Stopped:	
ATROPINE (Suspected)	Reason: Premedication		
Injection	0.5 Milligram	Daily	Intramuscular
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69072 **Seq:** 1 **Gender:** M
Reported: 09/01/1991 **Weight:** 8.00
Hospitalisation: **Age:** 7M
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Cogentin 0.02mg/kg, 1gm

Medicine Details:

DROLEPTAN (Suspected)	Reason: Premedication
Injection 1.0 Milligram Daily Intramuscular	
Batch: Started: Stopped:	
MORPHINE NOS (Suspected)	Reason: Premedication
Injection 1.0 Milligram Daily Intramuscular	
Batch: Started: Stopped:	
ATROPINE (Suspected)	Reason: Premedication
Injection 0.1 Milligram Daily Intramuscular	
Batch: Started: Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69738	Seq: 1	Gender: F
Reported: 27/02/1991		Weight: 40.00
Hospitalisation:		Age: 28Y
Onset Date: 21/02/1991		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hallucination			
Thinking abnormal			
Visual disturbance			Imi cogentin 2mg

Medicine Details:

DROPERIDOL (Suspected)		Reason: Other disturbance of sensation	
Injection	2.0 Milligram	Total	Intravenous
Batch:	Started: 21/02/1991	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69739	Seq: 1	Gender: M
Reported: 26/02/1991		Weight:
Hospitalisation:		Age: 19Y
Onset Date: 26/02/1991		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Opisthotonus			Imi cogentin 2mg.

Medicine Details:

DROPERIDOL (Suspected)		Reason: Other disturbance of sensation
Injection	2.5 Milligram	Total Intravenous
Batch:	Started: 26/02/1991	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69935	Seq: 1	Gender: F
Reported: 08/02/1991		Weight:
Hospitalisation:		Age: 26Y
Onset Date: 31/12/1990		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Grand mal convulsion			
Muscle twitching			
Nausea			

Medicine Details:

LORAZEPAM (Suspected)	Reason:
Tablet	0.0
Batch:	Started:
	Stopped:
DIAZEPAM (Suspected)	Reason:
Tablet	0.0
Batch:	Started:
	Stopped:
OMNOPON (Suspected)	Reason: Other disturbance of sensation
	0.0
Batch:	Started:
	Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
	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.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 69935

Seq: 1

Gender: F

Reported: 08/02/1991

Weight:

Hospitalisation:

Age: 26Y

Onset Date: 31/12/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

MIDAZOLAM (Suspected)	Reason: Other disturbance of sensation	
0.0		
Batch:	Started:	Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation	
120.0 Milligram	Daily	
Batch:	Started:	Stopped:

Laboratory Investigations:

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: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 70325	Seq: 1	Gender: F
Reported: 10/12/1990		Weight:
Hospitalisation:		Age: 17
Onset Date: 08/07/1988		DOB: 19/09/1970
Outcome: Unknown		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash			Promethazine commenced, cephalothin ceased.

Medicine Details:

ENFLURANE (Suspected)		Reason: Surgery
Inhalation	0.0	Inhalation
Batch:	Started: 08/07/1988	Stopped:
THIOPENTONE SODIUM (Suspected)		Reason: Surgery
Injection	0.0	Intravenous
Batch:	Started: 08/07/1988	Stopped:
SUXAMETHONIUM BROMIDE (Suspected)		Reason: Surgery
Injection	100.0 Milligram	Total Intravenous
Batch:	Started: 08/07/1988	Stopped:
PANCURONIUM BROMIDE (Suspected)		Reason: Surgery
Injection	6.0 Milligram	Total Intravenous
Batch:	Started: 08/07/1988	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 70325

Seq: 1

Gender: F

Reported: 10/12/1990

Weight:

Hospitalisation:

Age: 17

Onset Date: 08/07/1988

DOB: 19/09/1970

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

MORPHINE NOS (Suspected)	Reason: Surgery		
Injection	15.0 Milligram	Total	Intravenous
Batch:	Started: 08/07/1988	Stopped:	
DROPERIDOL (Suspected)	Reason: Surgery		
Injection	10.0 Milligram	Total	Intravenous
Batch:	Started: 08/07/1988	Stopped:	
CEPHALOTHIN SODIUM (Suspected)	Reason:		
Injection, intravenous infusion	4.0 Gram	Daily	Intravenous
Batch:	Started: 08/07/1988	Stopped: 08/07/1988	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 71230	Seq: 1	Gender: F
Reported: 15/05/1991		Weight:
Hospitalisation:		Age: 36Y
Onset Date: 07/11/1990		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Apnoea Paralysis			

Medicine Details:

PROPOFOL (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
SUXAMETHONIUM NOS (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
VECURONIUM BROMIDE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
NITROUS OXIDE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 71230

Seq: 1

Gender: F

Reported: 15/05/1991

Weight:

Hospitalisation:

Age: 36Y

Onset Date: 07/11/1990

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation	
0.0		
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation	
0.0		
Batch:	Started:	Stopped:
ENFLURANE (Suspected)	Reason: Other disturbance of sensation	
0.0		
Batch:	Started:	Stopped:
TEMAZEPAM (Suspected)	Reason: Premedication	
20.0 Milligram	1 time	
Batch:	Started: 07/11/1990	Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 72219 **Seq:** 1 **Gender:** M
Reported: 09/08/1991 **Weight:** 63.00
Hospitalisation: **Age:** 19Y
Onset Date: 29/07/1991 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Agitation			
Fatigue			
Insomnia			

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection	2.5 Milligram 1 time Intravenous
Batch:	Started: 29/07/1991 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 72472	Seq: 1	Gender: M
Reported: 07/08/1991		Weight: 65.00
Hospitalisation:		Age:
Onset Date: 16/06/1991		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Tremor	Severe	Dystonic reaction affecting mouth/tongue Whole body tremors	Settled with 6 mg cogentin

Medicine Details:

NILSTAT (Other drug)	Reason:
Mouthwash	40.0 Millilitre
Batch:	Started:
	Stopped:
ZANTAC (Other drug)	Reason: Other diseases of esophagus
Tablet	300.0 Milligram
Batch:	Started: 12/06/1991
	Stopped:
LASIX (Other drug)	Reason:
Injection	20.0 Milligram
Batch:	Started: 11/06/1991
	Stopped: 17/06/1991
ZOFRAN (Other drug)	Reason: Nausea and vomiting
Oral application	24.0 Milligram
Batch:	Started: 12/06/1991
	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 72472

Seq: 1

Gender: M

Reported: 07/08/1991

Weight: 65.00

Hospitalisation:

Age:

Onset Date: 16/06/1991

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROLEPTAN (Suspected)	Reason: Nausea and vomiting
	1.0 Milligram Per hour
Batch:	Started: 15/06/1991 Stopped: 16/06/1991

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 74464	Seq: 1	Gender: M
Reported: 27/11/1991		Weight: 88.00
Hospitalisation:		Age: 31Y
Onset Date: 22/11/1991		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Torticollis			Cogentin

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	15.0 Milligram	3 times	Intramuscular
Batch:	Started: 21/11/1991	Stopped: 22/11/1991	
OMNOPON (Suspected)		Reason:	
Injection	20.0 Milligram	1 time	Intramuscular
Batch:	Started: 21/11/1991	Stopped: 22/11/1991	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 74501	Seq: 1	Gender: F
Reported: 02/12/1991		Weight: 80.00
Hospitalisation:		Age: 46Y
Onset Date: 29/11/1991		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Cardiac arrest		Cardiac asystole.	

Medicine Details:

DROPERIDOL (Suspected)		Reason: Premedication	
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 29/11/1991	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient history: diazepam - dystonic reaction clindamycin - pseudomembranous colitis brufen - vomiting.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 75462	Seq: 1	Gender: M
Reported: 06/02/1992		Weight:
Hospitalisation:		Age: 21Y
Onset Date:		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Abdominal pain			
Renal tubular necrosis			

Medicine Details:

NITROUS OXIDE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
ISOFLURANE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:
FENTANYL CITRATE (Suspected)	0.0	Reason: Other disturbance of sensation
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatinine		03/12/1991	0.22	Acute tubular necrosis.
	Creatinine		05/12/1991	0.29	
	Creatinine		07/12/1991	0.24	
	Renal biopsy =				

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 75462

Seq: 1

Gender: M

Reported: 06/02/1992

Weight:

Hospitalisation:

Age: 21Y

Onset Date:

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PROPOFOL (Suspected)

Reason: Other disturbance of sensation

0.0

Batch:

Started:

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 75751	Seq: 1	Gender: F
Reported: 27/02/1992		Weight:
Hospitalisation:		Age: 33Y
Onset Date: 26/02/1992		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			

Medicine Details:

MORPHINE NOS (Other drug)		Reason:	
Injection	5.0 Milligram	As necessary	Subcutaneous
Batch:	Started:	Stopped:	
MIANSERIN HYDROCHLORIDE (Other drug)		Reason:	
	20.0 Milligram	Daily	
Batch:	Started:	Stopped:	
DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	2.5 Milligram	As necessary	Intravenous
Batch:	Started: 25/02/1992	Stopped: 26/02/1992	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 76088	Seq: 1	Gender: M
Reported: 20/03/1992		Weight: 80.00
Hospitalisation:		Age: 42Y
Onset Date: 20/03/1992		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Convulsion		Tonic clonic reaction	

Medicine Details:

PROPOFOL (Suspected)	Reason:
0.0	
Batch:	Started:
	Stopped:
TEMAZEPAM (Suspected)	Reason: Premedication
20.0 Milligram	Daily
Batch:	Started: 20/03/1992
	Stopped:
MAXOLON (Suspected)	Reason: Premedication
10.0 Milligram	Daily
Batch:	Started: 20/03/1992
	Stopped:
FENTANYL CITRATE (Suspected)	Reason: Premedication
50.0 Microgram	Daily
Batch:	Started: 20/03/1992
	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 76088

Seq: 1

Gender: M

Reported: 20/03/1992

Weight: 80.00

Hospitalisation:

Age: 42Y

Onset Date: 20/03/1992

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason: Premedication

1.2 Milligram

Daily

Batch:

Started: 20/03/1992

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 77057	Seq: 1	Gender: F
Reported: 18/05/1992		Weight:
Hospitalisation:		Age: 36Y
Onset Date: 04/05/1992		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Convulsion		Post operative convulsions	

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection	2.5 Milligram
	Intravenous
Batch:	Started:
Stopped:	
NITROUS OXIDE (Suspected)	Reason: Other disturbance of sensation
	0.0
Batch:	Started:
Stopped:	
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
Injection	150.0 Milligram
	Daily
	Intravenous
Batch:	Started:
Stopped:	
MIDAZOLAM (Suspected)	Reason: Other disturbance of sensation
Injection	5.0 Milligram
	Daily
	Intravenous
Batch:	Started:
Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 77459 **Seq:** 1 **Gender:** F
Reported: 07/05/1992 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: 07/03/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Cogentin 2mg imi to reverse droperidol.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	5.0 Milligram 5 times Intramuscular
Batch:	Started: 04/03/1992 Stopped: 07/03/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 77471 **Seq:** 1 **Gender:** M
Reported: 05/06/1992 **Weight:**
Hospitalisation: **Age:** 32Y
Onset Date: 05/06/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash		Red blotchy rash on trunk and arms	

Medicine Details:

MORPHINE NOS (Suspected)	Reason: Pain
Injection, intravenous infusion 50.0 Milligram Daily Intravenous	
Batch:	Started: 05/06/1992 Stopped: 05/06/1992
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection, intravenous infusion 5.0 Milligram Daily Intravenous	
Batch:	Started: 05/06/1992 Stopped: 05/06/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 78254

Seq: 1

Gender: F

Reported: 03/10/1991

Weight: 75.00

Hospitalisation:

Onset Date: 30/08/1991

Age: 45Y

Outcome: Recovered

DOB:

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dyskinesia			

Medicine Details:

SODIUM VALPROATE (Other drug)			Reason: Other&unspecified epilepsy
Oral application	600.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	CONTIN
THYROXINE SODIUM (Other drug)			Reason: Myxedema
Tablet	150.0 Microgram	Daily	Oral
Batch:	Started:	Stopped:	CONTIN
OXAZEPAM (Other drug)			Reason:
Tablet	90.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	CONTIN
DROPERIDOL (Suspected)			Reason: Nausea and vomiting
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 29/08/1991	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Oculogyric crisis previously experienced with maxolon and stemetil (phenothiazines) 29/10/82.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 78661

Seq: 1

Gender: F

Reported: 10/08/1992

Weight: 55.00

Hospitalisation:

Age: 22Y

Onset Date: 06/08/1992

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Cogentin

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	2.5 Milligram	Daily	Intramuscular
Batch:	Started: 05/08/1992	Stopped: 06/08/1992	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 79494	Seq: 1	Gender: M
Reported: 16/09/1992		Weight:
Hospitalisation: Admitted to hospital		Age: 15Y
Onset Date: 11/09/1992		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Tremor		Tremor in lower, then upper limbs.	
Dystonia			
Extrapyramidal disorder			Congentin 1mg im.

Medicine Details:

MAXOLON (Suspected)		Reason: Surgery
Injection	10.0 Milligram	1 time Intravenous
Batch:	Started: 11/09/1992	Stopped:
DROPERIDOL (Suspected)		Reason:
Injection	2.5 Milligram	1 time Intravenous
Batch:	Started: 11/09/1992	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 80557 **Seq:** 1 **Gender:** F
Reported: 16/10/1992 **Weight:** 57.00
Hospitalisation: **Age:** 32Y
Onset Date: 14/10/1992 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine 2mg iv

Medicine Details:

RANITIDINE (Other drug) Oral application Batch:	300.0 Milligram Started:	Daily L TERM	Oral Stopped: CONTIN	Reason:
OMNOPON (Suspected) Injection Batch:	20.0 Milligram Started: 14/10/1992	1 time	Intravenous Stopped: 14/10/1992	Reason: Other disturbance of sensation
STEMETIL (Suspected) Injection Batch:	12.5 Milligram Started: 14/10/1992	1 time	Intravenous Stopped: 14/10/1992	Reason:
DROLEPTAN (Suspected) Injection Batch:	2.5 Milligram Started: 14/10/1992	1 time	Intravenous Stopped: 14/10/1992	Reason:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 80557

Seq: 1

Gender: F

Reported: 16/10/1992

Weight: 57.00

Hospitalisation:

Age: 32Y

Onset Date: 14/10/1992

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)		Reason:	
Injection	50.0 Milligram	1 time	Intramuscular
Batch:	Started: 13/10/1992	Stopped:	13/10/1992
MAXOLON (Suspected)		Reason:	
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 13/10/1992	Stopped:	13/10/1992
HEPARIN SODIUM (Suspected)		Reason:	
Injection, intravenous infusion	1.0 Thousand Internal	Per hour	Intravenous
Batch:	Started: 14/10/1992	Stopped:	CONTIN

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 80923	Seq: 1	Gender: F
Reported: 12/11/1992		Weight: 70.00
Hospitalisation:		Age: 33
Onset Date: 11/11/1992		DOB: 27/01/1959
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Tremor			Cogentin iv 2mg.

Medicine Details:

MAXOLON (Suspected)	Reason: Hyperemes gravidar w/o neurits
Injection	20.0 Milligram 2 times Intravenous
Batch:	Started: 11/11/1992 Stopped: 11/11/1992
STEMETIL (Suspected)	Reason: Hyperemes gravidar w/o neurits
Suppository	25.0 Milligram As necessary Rectal
Batch:	Started: 03/11/1992 Stopped: 10/11/1992
PYRIDOXINE HYDROCHLORIDE (Suspected)	Reason: Hyperemes gravidar w/o neurits
Injection	50.0 Milligram Daily Intramuscular
Batch:	Started: 11/11/1992 Stopped:
DROPERIDOL (Suspected)	Reason: Hyperemes gravidar w/o neurits
Injection	7.5 Milligram Daily Intramuscular
Batch:	Started: 10/11/1992 Stopped: 11/11/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 81751	Seq: 1	Gender: M
Reported: 24/12/1992		Weight:
Hospitalisation:		Age: 12
Onset Date: 19/12/1992		DOB: 10/09/1980
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			8mg cogentin iv.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	3.0 Milligram 4 times Intramuscular
Batch:	Started: 18/12/1992 Stopped: 19/12/1992
PANADOL (Suspected)	Reason: Pain
	750.0 Milligram As necessary
Batch:	Started: 19/12/1992 Stopped: 21/12/1992
CODEINE (Suspected)	Reason: Pain
	20.0 Milligram As necessary
Batch:	Started: 19/12/1992 Stopped: 19/12/1992
BRICANYL (Suspected)	Reason: Bronchitis,unqualified
Inhalation	2.0 Dose Unspecified As necessary Inhalation
Batch:	Started: 16/12/1992 Stopped: 21/12/1992

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient appears to be sensitive to narcotics.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 81751

Seq: 1

Gender: M

Reported: 24/12/1992

Weight:

Hospitalisation:

Age: 12

Onset Date: 19/12/1992

DOB: 10/09/1980

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)

Reason: Pain

Injection, intravenous infusion

6.5 Milligram

Per hour

Intravenous

Batch:

Started: 16/12/1992

Stopped: 19/12/1992

Laboratory Investigations:

Additional Information:

Patient appears to be sensitive to narcotics.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 82184

Seq: 1

Gender: F

Reported: 28/01/1993

Weight: 60.00

Hospitalisation:

Onset Date: 27/01/1993

Age: 30Y

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Bronchospasm		Bronchoconstriction	Salbutamol.

Medicine Details:

ATRACURIUM BESYLATE (Suspected)		Reason:	
Injection	50.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	
SUXAMETHONIUM BROMIDE (Suspected)		Reason:	
Injection	100.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	
PROPOFOL (Suspected)		Reason:	
Injection	200.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	
TORADOL (Suspected)		Reason:	
Injection	30.0 Milligram	Daily	Intramuscular
Batch:	Started: 27/01/1993	Stopped: 27/01/1993	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 82184

Seq: 1

Gender: F

Reported: 28/01/1993

Weight: 60.00

Hospitalisation:

Age: 30Y

Onset Date: 27/01/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROLEPTAN (Suspected)		Reason:	
Injection	5.0 Milligram	Daily	Intravenous
Batch:	Started: 27/01/1993	Stopped:	27/01/1993
MAXOLON (Suspected)		Reason:	
Injection	10.0 Milligram	Daily	Intramuscular
Batch:	Started: 27/01/1993	Stopped:	27/01/1993
MEFOXIN (Suspected)		Reason:	
Injection	1.0 Gram	Daily	Intravenous
Batch:	Started: 27/01/1993	Stopped:	27/01/1993
FENTANYL CITRATE (Suspected)		Reason:	
Injection	100.0 Microgram	Daily	Intravenous
Batch:	Started: 27/01/1993	Stopped:	27/01/1993

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 82184

Seq: 1

Gender: F

Reported: 28/01/1993

Weight: 60.00

Hospitalisation:

Age: 30Y

Onset Date: 27/01/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

ATROPINE (Suspected)	Reason:
Injection	600.0 Microgram Daily Intravenous
Batch:	Started: 27/01/1993 Stopped: 27/01/1993

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 83315	Seq: 1	Gender: F
Reported: 19/03/1993		Weight: 55.00
Hospitalisation:		Age: 41Y
Onset Date: 23/02/1993		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Agitation Hyperkinesia Paraesthesia Thinking abnormal			Cogentin

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
5.0 Microgram	1 time
Batch:	Started: 23/02/1993 Stopped:
MAXOLON (Suspected)	Reason: Other disturbance of sensation
10.0 Milligram	1 time
Batch:	Started: 23/02/1993 Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
80.0 Milligram	1 time
Batch:	Started: 23/02/1993 Stopped:
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation
50.0 Milligram	1 time
Batch:	Started: 23/02/1993 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 83315

Seq: 1

Gender: F

Reported: 19/03/1993

Weight: 55.00

Hospitalisation:

Age: 41Y

Onset Date: 23/02/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

MIDAZOLAM (Suspected)	Reason: Other disturbance of sensation
	3.5 Milligram 1 time
Batch:	Started: 23/02/1993 Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 84248

Seq: 1

Gender: F

Reported: 27/04/1993

Weight: 72.00

Hospitalisation:

Age: 38

Onset Date: 19/03/1993

DOB: 31/10/1954

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Valium 10mg, cogentin 2mg im, observation for 3 hours.

Medicine Details:

PANADEINE (Other drug)	Reason: Pain
Oral application	2.0 Dose Unspecified Daily Oral
Batch:	Started: 17/03/1993 Stopped: 20/03/1993
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	1.2 Milligram Daily Intravenous
Batch:	Started: 18/03/1993 Stopped: 19/03/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 84781	Seq: 1	Gender: F
Reported: 31/05/1993		Weight:
Hospitalisation:		Age: 18Y
Onset Date: 09/04/1993		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			

Medicine Details:

TEMAZEPAM (Other drug)	Reason: Specific disorders of sleep
Capsule	20.0 Milligram Daily Oral
Batch:	Started: 06/04/1993 Stopped: 12/04/1993
PANADEINE (Other drug)	Reason: Pain
Oral application	12.0 Dose Unspecified Daily Oral
Batch:	Started: 06/04/1993 Stopped: 13/04/1993
VALIUM (Suspected)	Reason: Abnormal involuntary movement
	5.0 Milligram 1 time
Batch:	Started: 09/04/1993 Stopped:
MAXOLON (Suspected)	Reason: Nausea and vomiting
Injection	40.0 Milligram Daily Intramuscular
Batch:	Started: 06/04/1993 Stopped: 08/04/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 84781

Seq: 1

Gender: F

Reported: 31/05/1993

Weight:

Hospitalisation:

Age: 18Y

Onset Date: 09/04/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason: Nausea and vomiting

20.0 Milligram

Daily

Batch:

Started: 08/04/1993

Stopped: 09/04/1993

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87046 **Seq:** 1 **Gender:** F
Reported: 25/08/1993 **Weight:** 57.00
Hospitalisation: **Age:** 45Y
Onset Date: 12/08/1993 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dermatitis bullous Pruritus Rash		Blue/black blisters on r thumb & finger. Itchy at nape, breasts & extremities.	

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation
Injection 75.0 Milligram As necessary Intramuscular	
Batch: Started: 11/08/1993 Stopped: 13/08/1993	
PANADEINE (Suspected)	Reason:
Oral application 2.0 Dose Unspecified Daily Oral	
Batch: Started: 10/08/1993 Stopped: 14/08/1993	
MONOPRIL (Suspected)	Reason: Essential benign hypertension
Tablet 10.0 Milligram Daily Oral	
Batch: Started: 10/08/1993 Stopped: 14/08/1993	
ADALAT (Suspected)	Reason: Essential benign hypertension
Tablet 10.0 Milligram Daily Oral	
Batch: Started: 10/08/1993 Stopped: 13/08/1993	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had a previous reaction in penicillin.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87046

Seq: 1

Gender: F

Reported: 25/08/1993

Weight: 57.00

Hospitalisation:

Age: 45Y

Onset Date: 12/08/1993

DOB:

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	2.0 Milligram	Daily	Intramuscular
Batch:	Started: 11/08/1993	Stopped: 13/08/1993	
HEPARIN SODIUM (Suspected)	Reason:		
Injection	10.0 Thousand International	Daily	Subcutaneous
Batch:	Started: 11/08/1993	Stopped: 13/08/1993	
MAXOLON (Suspected)	Reason: Nausea and vomiting		
Injection	40.0 Milligram	Daily	Intramuscular
Batch:	Started: 11/08/1993	Stopped: 14/08/1993	

Laboratory Investigations:

Additional Information:

Patient has had a previous reaction in penicillin.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87490	Seq: 1	Gender: F
Reported: 06/09/1993		Weight: 59.00
Hospitalisation:		Age: 36Y
Onset Date: 11/06/1993		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Injection site reaction			
Urticaria			

Medicine Details:

ISOFLURANE (Other drug)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: Stopped:
ATROPINE (Suspected)	Reason:
0.0	
Batch:	Started: Stopped:
NEOSTIGMINE NOS (Suspected)	Reason:
0.0	
Batch:	Started: Stopped:
MORPHINE NOS (Suspected)	Reason: Other disturbance of sensation
Injection	15.0 Milligram Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had no known previous allergies.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87490

Seq: 1

Gender: F

Reported: 06/09/1993

Weight: 59.00

Hospitalisation:

Age: 36Y

Onset Date: 11/06/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection 15.0 Milligram	Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:
GLYCOPYRROLATE (Suspected)	Reason:
Injection 200.0 Microgram	Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
Injection 150.0 Milligram	Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:
ATRACURIUM BESYLATE (Suspected)	Reason:
Injection 35.0 Milligram	Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:

Laboratory Investigations:

Additional Information:

Patient has had no known previous allergies.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87490

Seq: 1

Gender: F

Reported: 06/09/1993

Weight: 59.00

Hospitalisation:

Age: 36Y

Onset Date: 11/06/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

Keflin Neutral (Suspected)

Reason: Prophylaxis

Injection

1.0 Gram

Daily

Intravenous

Batch:

Started: 11/06/1993

Stopped:

Laboratory Investigations:

Additional Information:

Patient has had no known previous allergies.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87961	Seq: 1	Gender: F
Reported: 28/09/1993		Weight: 68.00
Hospitalisation:		Age: 39Y
Onset Date: 01/06/1993		DOB:
Outcome: Not yet recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea Vomiting			

Medicine Details:

MORPHINE NOS (Suspected)	Reason: Pain
Injection, intravenous infusion	0.0
Batch:	Started: 31/05/1993
	Stopped: 01/06/1993
MAXOLON (Suspected)	Reason: Nausea and vomiting
Injection, intravenous infusion	0.0
Batch:	Started: 31/05/1993
	Stopped: 01/06/1993
OMNOPON-SCOPOLAMINE (Suspected)	Reason:
Injection	0.0
Batch:	Started: 31/05/1993
	Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	2.5 Milligram
Batch:	Started: 31/05/1993
	Stopped: 01/06/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Infusion ceased and patient commenced on toradol im.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 87961

Seq: 1

Gender: F

Reported: 28/09/1993

Weight: 68.00

Hospitalisation:

Age: 39Y

Onset Date: 01/06/1993

DOB:

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

HEPARIN CALCIUM (Suspected)	Reason:
Injection	15.0 Thousand Internal Daily Subcutaneous
Batch:	Started: 31/05/1993 Stopped: 01/06/1993

Laboratory Investigations:

Additional Information:

Infusion ceased and patient commenced on toradol im.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 88398	Seq: 1	Gender: U
Reported: 11/10/1993		Weight: 83.00
Hospitalisation:		Age:
Onset Date: 30/09/1993		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Neck stiff.	Cogentin 1gm iv.

Medicine Details:

MORPHINE HYDROCHLORIDE (Suspected)		Reason: Premedication	
Injection	15.0 Milligram	1 time	Intramuscular
Batch:	Started:	Stopped:	
DROPERIDOL (Suspected)		Reason: Premedication	
Injection	1.0 Milligram	1 time	Intramuscular
Batch:	Started:	Stopped:	
Keflin Neutral (Suspected)		Reason:	
Injection	4.0 Gram	Daily	Intravenous
Batch:	Started:	Stopped:	
FLAGYL (Suspected)		Reason:	
Injection	1.0 Gram	Daily	Intravenous
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 88502

Seq: 1

Gender: M

Reported: 15/10/1993

Weight: 68.00

Hospitalisation:

Age: 16Y

Onset Date: 30/09/1993

DOB:

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			lv cogentin 2mg.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection, intravenous infusion	4.0 Milligram Total Intravenous
Batch:	Started: 29/09/1993 Stopped: 30/09/1993

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has had no known previous allergies.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 89459

Seq: 1

Gender: F

Reported: 30/11/1993

Weight: 44.00

Hospitalisation:

Age: 38Y

Onset Date:

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypertonia			Benztropine 2mg im stat.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection 10.0 Milligram	1 time Intramuscular
Batch: Started: 24/11/1993	Stopped:
PETHIDINE HYDROCHLORIDE (Suspected)	Reason:
Injection 20.0 Milligram	1 time Intramuscular
Batch: Started:	Stopped:
PROMETHAZINE HYDROCHLORIDE (Suspected)	Reason:
Injection 50.0 Milligram	As necessary Intramuscular
Batch: Started:	Stopped:
DIAZEPAM (Suspected)	Reason:
Injection 10.0 Milligram	As necessary Intramuscular
Batch: Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 89459
Reported: 30/11/1993

Seq: 1

Gender: F
Weight: 44.00

Hospitalisation:

Onset Date:
Outcome: Recovered

Age: 38Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

PROTHIADEN (Suspected)	Reason:	
150.0 Milligram	Daily	
Batch:	Started:	Stopped:
ROHYPNOL (Suspected)	Reason:	
1.0 Dose Unspecified	Daily	
Batch:	Started:	Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 90557 **Seq:** 1 **Gender:** F
Reported: 19/01/1994 **Weight:**
Hospitalisation: **Age:** 35Y
Onset Date: 12/01/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Cogentin 5mg iv.

Medicine Details:

DROPERIDOL (Suspected)	Reason: Premedication
5.0 Milligram	1 time
Batch:	Started: 12/01/1994 Stopped: 12/01/1994
MORPHINE NOS (Suspected)	Reason: Premedication
10.0 Milligram	1 time
Batch:	Started: 12/01/1994 Stopped: 12/01/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 90876	Seq: 1	Gender: F
Reported: 03/02/1994		Weight: 60.00
Hospitalisation:		Age: 18Y
Onset Date: 21/01/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Benztropine 2mg

Medicine Details:

AMPICILLIN (Other drug)	Reason:
0.0	
Batch:	Started: Stopped:
FLAGYL (Other drug)	Reason:
Injection 1.5 Gram Daily Intravenous	
Batch:	Started: Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 3.0 Millilitre 1 time Intravenous	
Batch:	Started: 21/01/1994 Stopped: 21/01/1994
DROPERIDOL (Suspected)	Reason:
Injection 2.5 Milligram 1 time Intravenous	
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergic to morphine, maxalon, stemetil



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91229	Seq: 1	Gender: F
Reported: 23/02/1994		Weight: 68.00
Hospitalisation:		Age: 34
Onset Date: 14/06/1993		DOB: 12/08/1958
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Clenched teeth, locked jaw.	Cogentin imi 1mg stat than repeat in 30 mins.
Oculogyration			

Medicine Details:

CALCIPARINE (Other drug)	Reason:
Injection	10.0 Thousand Internal Daily Subcutaneous
Batch:	Started: 11/06/1993 Stopped: 14/06/1993
TORADOL (Other drug)	Reason: Other disturbance of sensation
Injection	30.0 Milligram Daily Intramuscular
Batch:	Started: 11/06/1993 Stopped: 16/06/1993
PETHIDINE HYDROCHLORIDE (Other drug)	Reason: Pain
Injection	800.0 Milligram Daily Intramuscular
Batch:	Started: 11/06/1993 Stopped: 14/06/1993
VALIUM (Other drug)	Reason:
Injection	20.0 Milligram Daily Intravenous
Batch:	Started: 11/06/1993 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient allergic to voltaren, adhesive plaster, pethidine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91229

Seq: 1

Gender: F

Reported: 23/02/1994

Weight: 68.00

Hospitalisation:

Age: 34

Onset Date: 14/06/1993

DOB: 12/08/1958

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting
Injection	30.0 Milligram	Daily Intramuscular
Batch:	Started: 10/06/1993	Stopped: 14/06/1993
STEMETIL (Suspected)		Reason: Nausea and vomiting
Injection	75.0 Milligram	Daily Intramuscular
Batch:	Started: 10/06/1993	Stopped: 14/06/1993
TEMAZEPAM (Suspected)		Reason: Pain
Capsule	1.0 Dose Unspecified	Daily Oral
Batch:	Started: 10/06/1993	Stopped: 14/06/1993

Laboratory Investigations:

Additional Information:

Patient allergic to voltaren, adhesive plaster, pethidine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91605	Seq: 1	Gender: F
Reported: 14/03/1994		Weight: 56.00
Hospitalisation:		Age: 16Y
Onset Date: 02/02/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Opisthotonus Dystonia		Arching back.	Iv cogentin 2mg.

Medicine Details:

MAXOLON (Suspected)		Reason:	
Injection	10.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
PETHIDINE HYDROCHLORIDE (Suspected)		Reason:	
Injection	50.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
ATROPINE (Suspected)		Reason:	
Injection	300.0 Microgram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
PROPOFOL (Suspected)		Reason:	
Injection	160.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91605

Seq: 1

Gender: F

Reported: 14/03/1994

Weight: 56.00

Hospitalisation:

Age: 16Y

Onset Date: 02/02/1994

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

LIGNOCAINE (Suspected)		Reason:	
Injection	100.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
DROPERIDOL (Suspected)		Reason:	
Injection	4.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
NARCAN (Suspected)		Reason:	
Injection	400.0 Microgram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	
TORADOL (Suspected)		Reason:	
Injection	30.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91605

Seq: 1

Gender: F

Reported: 14/03/1994

Weight: 56.00

Hospitalisation:

Age: 16Y

Onset Date: 02/02/1994

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

TACRINE HYDROCHLORIDE (Suspected)	Reason:		
Injection	30.0 Milligram	1 time	Intravenous
Batch:	Started: 02/02/1994	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91626	Seq: 1	Gender: M
Reported: 15/03/1994		Weight:
Hospitalisation:		Age: 20Y
Onset Date: 03/02/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dyspnoea		Difficulty with respiration	Cogentin 2mg, admission to icu.
Dysphagia			

Medicine Details:

Keflin Neutral (Other drug)		Reason: Prophylaxis	
Injection	4.0 Gram	Daily	Intravenous
Batch:	Started: 02/02/1994	Stopped: 04/02/1994	
FLAGYL (Other drug)		Reason: Prophylaxis	
Injection	1.0 Gram	Daily	Intravenous
Batch:	Started: 02/02/1994	Stopped: 04/02/1994	
DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	1.0 Milligram	As necessary	Intramuscular
Batch:	Started: 02/02/1994	Stopped: 03/02/1994	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91630

Seq: 1

Gender: M

Reported: 15/03/1994

Weight: 97.00

Hospitalisation:

Age: 32Y

Onset Date: 07/07/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Dyspnoea Laryngospasm		Lock-jaw.	Diazepam iv 2mg, cogentin 2mg imi.

Medicine Details:

PROPOFOL (Suspected) Injection Batch: 300.0 Milligram Started: 07/07/1993	Reason: Other disturbance of sensation 1 time Intravenous Stopped:
LIGNOCAINE (Suspected) Injection Batch: 60.0 Milligram Started: 07/07/1993	Reason: Other disturbance of sensation 1 time Intravenous Stopped:
ATROPINE (Suspected) Injection Batch: 400.0 Microgram Started: 07/07/1993	Reason: Other disturbance of sensation 1 time Intravenous Stopped:
DROPERIDOL (Suspected) Injection Batch: 4.0 Milligram Started: 07/07/1993	Reason: Other disturbance of sensation 1 time Intravenous Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No previous problems with anaesthetics.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 91630

Seq: 1

Gender: M

Reported: 15/03/1994

Weight: 97.00

Hospitalisation:

Age: 32Y

Onset Date: 07/07/1993

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

MAXOLON (Suspected)	Reason: Other disturbance of sensation		
Injection	10.0 Milligram	1 time	Intravenous
Batch:	Started: 07/07/1993	Stopped:	
THA (Suspected)	Reason: Other disturbance of sensation		
Injection	30.0 Milligram	1 time	Intravenous
Batch:	Started: 07/07/1993	Stopped:	
NARCAN (Suspected)	Reason: Other disturbance of sensation		
Injection	600.0 Microgram	1 time	Intravenous
Batch:	Started: 07/07/1993	Stopped:	

Laboratory Investigations:

Additional Information:

No previous problems with anaesthetics.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93067	Seq: 1	Gender: F
Reported: 17/05/1994		Weight:
Hospitalisation:		Age: 30
Onset Date: 11/08/1993		DOB: 03/07/1963
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypoaesthesia		Numbness in fingers and toes.	
Malaise			
Nausea			Stemetil iv.

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	1.2 Milligram	1 time	Intravenous
Batch:	Started: 11/08/1993	Stopped:	
VALIUM (Suspected)		Reason:	
Injection	2.5 Milligram	1 time	Intravenous
Batch:	Started: 11/08/1993	Stopped:	
FENTANYL CITRATE (Suspected)		Reason:	
Injection	15.0 Microgram	1 time	Intravenous
Batch:	Started: 11/08/1993	Stopped:	
DIAZEPAM (Suspected)		Reason:	
Tablet	10.0 Milligram	1 time	Oral
Batch:	Started: 11/08/1993	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous reactions to pethidine=vomiting, panadeine forte=vomiting.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93487	Seq: 1	Gender: F
Reported: 06/06/1994		Weight: 57.00
Hospitalisation:		Age: 85Y
Onset Date: 30/03/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Constipation			Nulax given

Medicine Details:

ZOFRAN (Suspected)	Reason: Nausea and vomiting
Injection	4.0 Milligram 1 time Intravenous
Batch:	Started: 28/03/1994 Stopped: 28/03/1994
PANADEINE FORTE (Suspected)	Reason:
Capsule	1.0 Dose Unspecified As necessary Oral
Batch:	Started: Stopped:
DROPERIDOL (Suspected)	Reason:
Injection	1.2 Milligram As necessary Intramuscular
Batch:	Started: Stopped:
STEMETIL (Suspected)	Reason:
Injection	12.0 Milligram Daily Intramuscular
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93490	Seq: 1	Gender: M
Reported: 06/06/1994		Weight: 73.00
Hospitalisation:		Age: 72
Onset Date: 26/05/1994		DOB: 12/06/1921
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Arrhythmia Dizziness Nausea Pallor			

Medicine Details:

CLEXANE (Other drug)		Reason: Other prophylactic procedures	
Injection	40.0 Milligram	Daily	Subcutaneous
Batch:	Started: 25/05/1994	Stopped:	CONTIN
PETHIDINE HYDROCHLORIDE (Other drug)		Reason: Pain	
Injection	800.0 Milligram	Daily	Intramuscular
Batch:	Started: 26/05/1994	Stopped:	
DROPERIDOL (Suspected)		Reason:	
Injection	2.5 Milligram	Daily	Intramuscular
Batch:	Started: 26/05/1994	Stopped:	26/05/1994
Keflin Neutral (Suspected)		Reason:	
Injection	4.0 Gram	Daily	Intravenous
Batch:	Started: 25/05/1994	Stopped:	27/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergies to penicillin and garlic.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93663	Seq: 1	Gender: F
Reported: 10/06/1994		Weight: 72.00
Hospitalisation:		Age: 71
Onset Date: 26/04/1994		DOB: 26/01/1923
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Coma			
Dystonia			

Medicine Details:

FLUCLOXACILLIN SODIUM (Other drug)	Reason:	
1.5 Gram	Daily	
Batch:	Started: 25/04/1994	Stopped: 26/04/1994
LASIX (Other drug)	Reason: Essential benign hypertension	
40.0 Milligram	Daily	
Batch:	Started: 26/04/1994	Stopped:
OROXINE (Other drug)	Reason:	
Tablet	50.0 Microgram	Daily Oral
Batch:	Started:	Stopped: CONTIN
ISOPTIN (Other drug)	Reason: Essential benign hypertension	
Tablet, modified release	240.0 Milligram	Daily Oral
Batch:	Started:	Stopped: 03/05/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93663

Seq: 1

Gender: F

Reported: 10/06/1994

Weight: 72.00

Hospitalisation:

Age: 71

Onset Date: 26/04/1994

DOB: 26/01/1923

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

ZANTAC (Other drug)	Reason: Diaph hern of abd cav w/o obst		
Tablet	300.0 Milligram	Daily	Oral
Batch:	Started:	Stopped: 05/05/1994	
FRAGMIN (Other drug)	Reason: Pulmonary embolism&infarction		
	10.0 Thousand Internal	Daily	
Batch:	Started:	Stopped:	
NORDIOL NOS (Other drug)	Reason:		
	15.0 Milligram	Daily	
Batch:	Started:	Stopped:	
SPAN-K (Other drug)	Reason:		
	6.0 Dose Unspecified	Daily	
Batch:	Started:	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 93663

Seq: 1

Gender: F

Reported: 10/06/1994

Weight: 72.00

Hospitalisation:

Age: 71

Onset Date: 26/04/1994

DOB: 26/01/1923

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	2.5 Milligram 1 time Intramuscular
Batch:	Started: 26/04/1994 Stopped: 26/04/1994
KEFLEX (Suspected)	Reason:
	1.5 Gram Daily
Batch:	Started: 25/04/1994 Stopped: 26/04/1994
LOSEC (Suspected)	Reason: Diaph hern of abd cav w/o obst
	40.0 Milligram Daily
Batch:	Started: 26/04/1994 Stopped:
ROHYPNOL (Suspected)	Reason:
	1.0 Dose Unspecified Daily
Batch:	Started: 26/04/1994 Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 95095	Seq: 1	Gender: U
Reported: 16/08/1994		Weight:
Hospitalisation:		Age:
Onset Date: 28/07/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Opisthotonus Dystonia	Required Specialist Consultation	Tightness in neck and jaw	Cogentin 1mg iv

Medicine Details:

PANADOL (Other drug)		Reason: Pain
Oral application	500.0 Milligram	Total Oral
Batch:	Started: 28/07/1994	Stopped:
PANADEINE FORTE (Suspected)		Reason: Pain
Tablet	0.0	Total Oral
Batch:	Started: 28/07/1994	Stopped:
PETHIDINE HYDROCHLORIDE (Suspected)		Reason: Pain
	300.0 Milligram	Total
Batch:	Started: 27/07/1994	Stopped: 28/07/1994
MAXOLON (Suspected)		Reason: Nausea and vomiting
Injection	10.0 Milligram	Total Intravenous
Batch:	Started: 27/07/1994	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 95095

Seq: 1

Gender: U

Reported: 16/08/1994

Weight:

Hospitalisation:

Age:

Onset Date: 28/07/1994

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	2.5 Milligram	Total	Intramuscular
Batch:	Started: 27/07/1994	Stopped:	
Keflin Neutral (Suspected)		Reason: Prophylaxis	
Injection	1.0 Gram	Daily	Intravenous
Batch:	Started: 27/07/1994	Stopped: 28/07/1994	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 96405	Seq: 1	Gender: F
Reported: 10/10/1994		Weight:
Hospitalisation:		Age: 66
Onset Date:		DOB: 08/10/1928
Outcome: Not yet recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Anxiety	Severe		Xanax
Hallucination	Severe		

Medicine Details:

DROLEPTAN (Suspected)	Reason:
2.0 Milligram	1 time
Batch:	Started: Stopped:
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 96932	Seq: 1	Gender: F
Reported: 07/11/1994		Weight: 56.00
Hospitalisation:		Age: 20Y
Onset Date: 22/10/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Mydriasis Opisthotonus			Benztropine iv 2mg.

Medicine Details:

METRONIDAZOLE (Other drug)		Reason: Prophylaxis	
Injection	1.5 Gram	Total	Intravenous
Batch:	Started: 21/10/1994	Stopped:	
AMOXYCILLIN SODIUM (Other drug)		Reason: Prophylaxis	
Injection	3.0 Gram	Total	Intravenous
Batch:	Started: 21/10/1994	Stopped:	
PETHIDINE HYDROCHLORIDE (Other drug)		Reason: Pain	
Injection	25.0 Milligram	As necessary	Intravenous
Batch:	Started: 20/10/1994	Stopped: 21/10/1994	
PANADEINE (Other drug)		Reason: Pain	
Oral application	1.0 Dose Unspecified	As necessary	Oral
Batch:	Started: 20/10/1994	Stopped:	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 96932

Seq: 1

Gender: F

Reported: 07/11/1994

Weight: 56.00

Hospitalisation:

Age: 20Y

Onset Date: 22/10/1994

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

BECOTIDE (Other drug)	Reason: Asthma
Inhalation	2.0 Dose Unspecified Daily Inhalation
Batch:	Started: Stopped: CONTIN
METOCLOPRAMIDE HYDROCHLORIDE (Suspected)	Reason: Nausea and vomiting
Injection	10.0 Milligram 2 times Intravenous
Batch:	Started: 21/10/1994 Stopped: 22/10/1994
DROLEPTAN (Suspected)	Reason: Nausea and vomiting
Injection	2.0 Milligram 1 time Intravenous
Batch:	Started: 22/10/1994 Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 96971 **Seq:** 1 **Gender:** M
Reported: 08/11/1994 **Weight:**
Hospitalisation: **Age:** 14Y
Onset Date: 17/08/1994 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Meningism			Cogentin

Medicine Details:

DROLEPTAN (Suspected)	Reason: Nausea and vomiting
	1.0 Dose Unspecified 1 time
Batch:	Started: 17/08/1994 Stopped: 17/08/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 98098	Seq: 1	Gender: F
Reported: 03/01/1995		Weight:
Hospitalisation:		Age: 29Y
Onset Date: 21/07/1993		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder Opisthotonus			

Medicine Details:

TEMAZE (Other drug)	Reason: Specific disorders of sleep
Capsule	10.0 Milligram
	As necessary Oral
Batch:	Started: 18/07/1993 Stopped:
PANADOL (Other drug)	Reason:
	0.0
	As necessary
Batch:	Started: 18/07/1993 Stopped:
PETHIDINE HYDROCHLORIDE (Other drug)	Reason:
	75.0 Milligram
	As necessary
Batch:	Started: 19/07/1993 Stopped:
DRIXINE (Other drug)	Reason:
	3.0 Dose Unspecified Daily
Batch:	Started: 20/07/1993 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergic to metoclopramide - extrapyramidal reaction.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 98098

Seq: 1

Gender: F

Reported: 03/01/1995

Weight:

Hospitalisation:

Age: 29Y

Onset Date: 21/07/1993

DOB:

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason: Nausea and vomiting

10.0 Milligram

As necessary

Batch:

Started: 20/07/1993

Stopped: 21/07/1993

Laboratory Investigations:

Additional Information:

Allergic to metoclopramide - extrapyramidal reaction.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 98105	Seq: 1	Gender: F
Reported: 03/01/1995		Weight:
Hospitalisation:		Age: 51Y
Onset Date: 17/08/1994		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder			Cogentin

Medicine Details:

ROHYPNOL (Other drug)	Reason:
Batch:	1.0 Dose Unspecified As necessary
Started:	Stopped:
DIHYDERGOT (Other drug)	Reason: Migraine
Injection	4.0 Milligram Daily Intravenous
Batch:	Started:
Stopped:	
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
	15.0 Milligram Daily
Batch:	Started: 16/08/1994
	Stopped: 17/08/1994

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 99245	Seq: 1	Gender: F
Reported: 28/02/1995		Weight:
Hospitalisation:		Age: 93
Onset Date: 03/01/1995		DOB: 10/01/1901
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypoglycaemia			Actrapid insulin

Medicine Details:

ISOTARD MC (Other drug)		Reason: Diabetes mellitus	
Injection	4.0 Unit	Daily	Subcutaneous
Batch:	Started:	L TERM	Stopped:
ASPIRIN (Other drug)		Reason:	
Oral application	150.0 Milligram	Daily	Oral
Batch:	Started:	L TERM	Stopped:
NICOTINIC ACID (Other drug)		Reason:	
Tablet	187.0 Milligram	Daily	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:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 99245

Seq: 1

Gender: F

Reported: 28/02/1995

Weight:

Hospitalisation:

Age: 93

Onset Date: 03/01/1995

DOB: 10/01/1901

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)

Reason: Premedication

Injection

2.5 Milligram

1 time

Intravenous

Batch:

Started: 03/01/1995

Stopped:

DIAZEPAM (Suspected)

Reason: Premedication

Injection

10.0 Milligram

1 time

Intravenous

Batch:

Started: 03/01/1995

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 99895	Seq: 1	Gender: M
Reported: 31/03/1995		Weight:
Hospitalisation:		Age: 27Y
Onset Date: 09/03/1995		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Haematuria	Severe	Gross haematuria	Iv fluids and panadol prn
Renal pain		Loin pain both sides.	
Renal impairment		Renal impairment	
Myoglobinuria			
Myopathy			

Medicine Details:

NITROUS OXIDE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 09/03/1995 Stopped:
ISOFLURANE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 09/03/1995 Stopped:
SUXAMETHONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation
Injection	100.0 Milligram 1 time Intravenous
Batch:	Started: 09/03/1995 Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection	125.0 Microgram 1 time Intravenous
Batch:	Started: 09/03/1995 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Biochemistry	<10			Urine myoglobin 10/3/95 42500 ug/l
	Creatine	30 - 210	09/03/1995	74	
	Creatine	30 - 210	10/03/1995	160	
	Creatine	30 - 210	12/03/1995	281	
	Creatine	30 - 210	11/03/1995	102	
	Creatine	30 - 210	13/03/1995	-	
	Creatinine	0.06 - 0.12	09/03/1995	0.13	

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 99895
Reported: 31/03/1995
Hospitalisation:
Onset Date: 09/03/1995
Outcome: Recovered

Seq: 1

Gender: M
Weight:
Age: 27Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

TORADOL (Suspected)	Reason: Pain
Injection 30.0 Milligram	1 time Intramuscular
Batch:	Started: 09/03/1995 Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
Injection 40.0 Milligram	1 time Intravenous
Batch:	Started: 09/03/1995 Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Milligram	1 time Intravenous
Batch:	Started: 09/03/1995 Stopped:
HYPNOVEL (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Milligram	1 time Intravenous
Batch:	Started: 09/03/1995 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatinine	0.06 - 0.12	10/03/1995	0.13	
	Creatinine	0.06 - 0.12	11/03/1995	0.13	
	Creatinine	0.06 - 0.12	13/03/1995	0.12	
	Creatinine	0.06 - 0.12	12/03/1995	0.14	
	Urea	3 - 7.5	09/03/1995	8.0	
	Urea	3 - 7.5	10/03/1995	8.4	
	Urea	3 - 7.5	11/03/1995	9.2	

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 99895

Seq: 1

Gender: M

Reported: 31/03/1995

Weight:

Hospitalisation:

Age: 27Y

Onset Date: 09/03/1995

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

AMOXIL (Suspected)	Reason: Prophylaxis		
Injection	1.0 Gram	1 time	Intravenous
Batch:	Started: 09/03/1995	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Urea	3 - 7.5	12/03/1995	8.3	
	Urea	3 - 7.5	13/03/1995	8.0	

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 102085 **Seq:** 1 **Gender:** F
Reported: 10/07/1995 **Weight:** 74.00
Hospitalisation: **Age:** 27
Onset Date: **DOB:** 14/03/1968
Outcome: Death, maybe drug **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Brain oedema Hyponatraemia Nausea Vomiting			

Medicine Details:

LARGACTIL (Suspected) Injection 25.0 Milligram Batch: Started: Stopped:	Reason: Nausea and vomiting As necessary Intravenous
DROPERIDOL (Suspected) Injection 2.0 Milligram Batch: Started: Stopped:	Reason: Nausea and vomiting As necessary Intravenous
STEMETIL (Suspected) Injection 12.5 Milligram Batch: Started: Stopped:	Reason: As necessary Intramuscular
ZOFRAN (Suspected) Injection 4.0 Milligram Batch: Started: Stopped:	Reason: Daily Intravenous

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was not well controlled epileptic, who after surgery was hyponatraemic and vomiting, died of cerebral oedema.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 102085 **Seq:** 1
Reported: 10/07/1995

Gender: F
Weight: 74.00
Age: 27
DOB: 14/03/1968
Causality: Causality possible

Hospitalisation:

Onset Date:
Outcome: Death, maybe drug

Reaction Details:

Medicine Details:

DILANTIN (Suspected)		Reason: Other&unspecified epilepsy	
Tablet	300.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	
PETHIDINE HYDROCHLORIDE (Suspected)		Reason:	
Injection	400.0 Milligram	Daily	Intramuscular
Batch:	Started:	Stopped:	
PANADEINE FORTE (Suspected)		Reason:	
Tablet	12.0 Dose Unspecified	Daily	Oral
Batch:	Started:	Stopped:	

Laboratory Investigations:

Additional Information:

Patient was not well controlled epileptic, who after surgery was hyponatraemic and vomiting, died of cerebral oedema.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 103115 **Seq:** 1 **Gender:** F
Reported: 28/08/1995 **Weight:**
Hospitalisation: **Age:** 23Y
Onset Date: 17/07/1995 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Tremor			Cogentin

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason:	
	1.0 Dose Unspecified 1 time	
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting	
	2.5 Milligram Daily	
Batch:	Started: 17/07/1995	Stopped: 17/07/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105062

Seq: 1

Gender: F

Reported: 04/12/1995

Weight: 69.00

Hospitalisation:

Age: 61Y

Onset Date:

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Mouth appeared tight,lips pursed,can't communicate	Cogentin 2mg iv, midazolam 10mg

Medicine Details:

DROPERIDOL (Suspected)	Reason:
0.0	
Batch:	Started: 28/11/1995 Stopped: 29/11/1995
MAXOLON (Suspected)	Reason: Nausea and vomiting
Injection	10.0 Milligram Daily Intramuscular
Batch:	Started: 29/11/1995 Stopped: 29/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105289 **Seq:** 1 **Gender:** U
Reported: 11/12/1995 **Weight:** 63.00
Hospitalisation: **Age:**
Onset Date: 16/11/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash maculo-papular		Over trunk and arms	Phenergan 25mg imi once

Medicine Details:

DIPRIVAN (Suspected)	Reason: Surgery
Injection 0.0 Intravenous	
Batch:	Started: 15/11/1995 Stopped: 15/11/1995
MAXOLON (Suspected)	Reason: Surgery
Injection 0.0 Intravenous	
Batch:	Started: 15/11/1995 Stopped: 15/11/1995
DROPERIDOL (Suspected)	Reason: Surgery
Injection 0.0 Intravenous	
Batch:	Started: 15/11/1995 Stopped:
CLAFORAN (Suspected)	Reason: Surgery
2.0 Gram 1 time	
Batch:	Started: 15/11/1995 Stopped: 15/11/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drugs



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105289 **Seq:** 1
Reported: 11/12/1995

Gender: U
Weight: 63.00

Hospitalisation:

Onset Date: 16/11/1995
Outcome: Not yet recovered

Age:
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)		Reason: Pain	
Injection	15.0 Milligram	Daily	Intravenous
Batch:	Started: 15/11/1995	Stopped: 17/11/1995	
TEMAZEPAM (Suspected)		Reason:	
Oral application	20.0 Milligram	Daily	Oral
Batch:	Started: 15/11/1995	Stopped: 15/11/1995	
RANITIDINE (Suspected)		Reason:	
Oral application	150.0 Milligram	Daily	Oral
Batch:	Started: 15/11/1995	Stopped: 15/11/1995	

Laboratory Investigations:

Additional Information:

See original report for other drugs



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105628	Seq: 1	Gender: F
Reported: 02/01/1996		Weight: 55.00
Hospitalisation:		Age: 22Y
Onset Date: 18/12/1995		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypertonia	Severe	Mainly in back, neck, legs and abdomen	7mg diazepam orally

Medicine Details:

BECONASE (Other drug)	Reason: Hay fever
Inhalation	4.0 Dose Unspecified Daily Inhalation
Batch:	Started: 01/01/1980 Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
	10.0 Milligram As necessary
Batch:	Started: 17/12/1995 Stopped: 18/12/1995
STEMETIL (Suspected)	Reason: Nausea and vomiting
Injection	12.5 Milligram As necessary Intramuscular
Batch:	Started: 17/12/1995 Stopped: 18/12/1995
Keflin Neutral (Suspected)	Reason: Othr diseases of urinary tract
	4.0 Gram Daily
Batch:	Started: 17/12/1995 Stopped: 18/12/1995

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: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105693 **Seq:** 1 **Gender:** F
Reported: 04/01/1996 **Weight:**
Hospitalisation: **Age:** 0
Onset Date: 03/12/1995 **DOB:** 11/08/1995
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Bentztropine.

Medicine Details:

PETHIDINE HYDROCHLORIDE (Other drug)	Reason: Pain
Injection 200.0 Milligram Daily Intravenous	
Batch: Started: 30/11/1995 Stopped: 06/12/1995	
DOXYCYCLINE HYDROCHLORIDE (Other drug)	Reason: Female pelvic inflammatory dis
Oral application 200.0 Milligram Daily Oral	
Batch: Started: 30/11/1995 Stopped:	
METRONIDAZOLE (Other drug)	Reason: Female pelvic inflammatory dis
Injection 1.5 Gram Daily Intravenous	
Batch: Started: 30/11/1995 Stopped:	
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection 10.0 Milligram Daily Intravenous	
Batch: Started: 02/12/1995 Stopped: 03/12/1995	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105693

Seq: 1

Gender: F

Reported: 04/01/1996

Weight:

Hospitalisation:

Age: 0

Onset Date: 03/12/1995

DOB: 11/08/1995

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

MAXOLON (Suspected)		Reason: Nausea and vomiting
	40.0 Milligram	Daily
Batch:	Started: 30/11/1995	Stopped: 01/12/1995
STEMETIL (Suspected)		Reason: Nausea and vomiting
Injection	50.0 Milligram	Daily Intramuscular
Batch:	Started: 01/12/1995	Stopped: 02/12/1995

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105768	Seq: 1	Gender: M
Reported: 03/01/1996		Weight:
Hospitalisation:		Age: 36Y
Onset Date:		DOB:
Outcome: Recovered with sequelae		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypertonia Pyrexia Tachycardia			Treated with 8 unilateral ect sessions

Medicine Details:

DIAZEPAM (Suspected)	Reason:
Injection	0.0 Intravenous
Batch:	Started: Stopped:
DROPERIDOL (Suspected)	Reason:
	0.0
Batch:	Started: Stopped:
TRIFLUOPERAZINE HYDROCHLORIDE (Suspected)	Reason: Unspecified schizophrenia
	50.0 Milligram Total
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Creatine	25-125 UNITS/LITER			Ck level in plasma- 1180 on 2nd day of admission; peaked at 2574 on 3rd day

Additional Information:

Overdose with 50mg trifluoperazine



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 105919 **Seq:** 1 **Gender:** F
Reported: 18/01/1996 **Weight:**
Hospitalisation: **Age:** 22
Onset Date: 30/10/1995 **DOB:** 25/01/1973
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Agitation Anxiety Palpitations			Midazolam iv 2mg

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection 2.5 Milligram	1 time Intramuscular
Batch:	Started: 30/10/1995 Stopped: 30/10/1995
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation
Injection 100.0 Milligram	1 time Intramuscular
Batch:	Started: 30/10/1995 Stopped: 30/10/1995
BETAMETHASONE (Suspected)	Reason:
Injection 11.0 Milligram	1 time Intramuscular
Batch:	Started: 30/10/1995 Stopped: 30/10/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 108250
Reported: 03/05/1996

Seq: 1

Gender: F

Weight:

Hospitalisation:

Onset Date: 20/03/1996

Age:

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Extrapyramidal disorder			Cogent

Medicine Details:

PANCREAZE (Other drug)	Reason: Acute pancreatitis
20.0 Milligram Daily	
Batch:	Started: L TERM Stopped:
TEMAZE (Other drug)	Reason:
10.0 Milligram Daily	
Batch:	Started: 19/03/1996 Stopped:
PANADEINE FORTE (Other drug)	Reason: Pain
20.0 Milligram Daily	
Batch:	Started: 19/03/1996 Stopped:
GENTAMICIN SULPHATE (Other drug)	Reason: Unspecified septicemia
Oral application 240.0 Milligram Daily Oral	
Batch:	Started: 19/03/1996 Stopped: 24/03/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 108250
Reported: 03/05/1996

Seq: 1

Gender: F

Weight:

Hospitalisation:

Age:

Onset Date: 20/03/1996

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

AMOXYCILLIN SODIUM (Other drug)	Reason: Unspecified septicemia		
Injection	3.0 Gram	Daily	Intravenous
Batch:	Started: 19/03/1996	Stopped: 25/03/1996	
METRONIDAZOLE (Other drug)	Reason: Unspecified septicemia		
Injection	1.5 Gram	Daily	Intravenous
Batch:	Started: 19/03/1996	Stopped: 25/03/1996	
MORPHINE NOS (Other drug)	Reason: Pain		
Injection	60.0 Milligram	Daily	Intramuscular
Batch:	Started: 19/03/1996	Stopped: 27/03/1996	
DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	15.0 Milligram	Daily	Intravenous
Batch:	Started: 20/03/1996	Stopped: 21/03/1996	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 108350 **Seq:** 1 **Gender:** F
Reported: 08/05/1996 **Weight:**
Hospitalisation: **Age:** 48Y
Onset Date: 12/04/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Bronchospasm Flushing Lacrimal disorder Periorbital oedema			

Medicine Details:

HEPARIN SODIUM (Other drug) Injection Batch:	0.0 Thousand Internal Started: 10/04/1996	Reason: 5 times Stopped:	Subcutaneous
ZANTAC (Other drug) Tablet Batch:	150.0 Milligram Started: 08/04/1996	Reason: 1 time Stopped:	Oral
ATROPINE (Other drug) Batch:	600.0 Microgram Started: 09/04/1996	Reason: Otr&nos disord of heart rhythm 1 time Stopped:	
MEFOXIN (Other drug) Injection Batch:	4.0 Gram Started: 09/04/1996	Reason: Total Stopped:	Intravenous

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 108350
Reported: 08/05/1996
Hospitalisation:
Onset Date: 12/04/1996
Outcome: Recovered

Seq: 1

Gender: F
Weight:
Age: 48Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

EPHEDRINE (Other drug)	Reason: Hypotension		
Injection	12.0 Milligram	Total	Intravenous
Batch:	Started: 10/04/1996	Stopped:	
MORPHINE NOS (Suspected)	Reason:		
	2.0 Milligram	1 time	
Batch:	Started: 12/04/1996	Stopped:	
DROPERIDOL (Suspected)	Reason:		
	83.0 Microgram	1 time	
Batch:	Started: 12/04/1996	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 108934 **Seq:** 1 **Gender:** F
Reported: 03/06/1996 **Weight:**
Hospitalisation: **Age:** 23Y
Onset Date: **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration Hypertonia Abdominal pain Bronchospasm Dyspnoea Dystonia Extrapyramidal disorder Neuroleptic malignant syndrome Paraesthesia Tachycardia		Eyes rolling up Rigidity and spasms in leg	

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Pain
200.0 Milligram	Total
Batch:	Started: 29/11/1995 Stopped: 30/11/1995
DROLEPTAN (Suspected)	Reason: Nausea and vomiting
10.0 Milligram	Daily
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient known as 'vomiter'.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109165	Seq: 1	Gender: F
Reported: 17/06/1996		Weight: 97.00
Hospitalisation:		Age: 43
Onset Date: 28/05/1996		DOB: 15/03/1953
Outcome: Unknown		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Drug ineffective Nausea Vomiting			Ondansetron

Medicine Details:

ISOFLURANE (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
FENTANYL CITRATE (Suspected)	100.0 Milligram	1 time
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	500.0 Microgram	1 time
Batch:	Started:	Stopped:
PETHIDINE HYDROCHLORIDE (Suspected)	75.0 Milligram	1 time
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109165

Seq: 1

Gender: F

Reported: 17/06/1996

Weight: 97.00

Hospitalisation:

Age: 43

Onset Date: 28/05/1996

DOB: 15/03/1953

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

MAXOLON (Suspected)	Reason:
10.0 Milligram	1 time
Batch:	Started: Stopped:
DIPRIVAN (Suspected)	Reason:
200.0 Milligram	1 time
Batch:	Started: Stopped:
VECURONIUM BROMIDE (Suspected)	Reason:
5.0 Milligram	1 time
Batch:	Started: Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109297

Seq: 1

Gender: F

Reported: 17/06/1996

Weight: 58.00

Hospitalisation:

Age: 36

Onset Date: 23/05/1996

DOB: 28/10/1959

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea			Maxolon, stemetil

Medicine Details:

FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 23/05/1996 Stopped: 23/05/1996
VECURONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 23/05/1996 Stopped: 23/05/1996
ISOFLURANE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 23/05/1996 Stopped: 23/05/1996
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
2.5 Milligram 1 time	
Batch:	Started: 23/05/1996 Stopped: 23/05/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109297

Seq: 1

Gender: F

Reported: 17/06/1996

Weight: 58.00

Hospitalisation:

Age: 36

Onset Date: 23/05/1996

DOB: 28/10/1959

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)

Reason: Pain

50.0 Milligram

1 time

Batch:

Started: 23/05/1996

Stopped: 23/05/1996

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109309	Seq: 1	Gender: F
Reported: 17/06/1996		Weight: 81.00
Hospitalisation:		Age: 58
Onset Date: 20/05/1996		DOB: 02/04/1938
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Headache			
Nausea			Maxolon

Medicine Details:

ISOFLURANE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 20/05/1996 Stopped: 20/05/1996
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection	100.0 Milligram 1 time Intravenous
Batch:	Started: 20/05/1996 Stopped: 20/05/1996
OMNOPON (Suspected)	Reason: Other disturbance of sensation
Injection	20.0 Milligram 1 time Intramuscular
Batch:	Started: 20/05/1996 Stopped: 20/05/1996
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation
Injection	200.0 Milligram 1 time Intravenous
Batch:	Started: 20/05/1996 Stopped: 20/05/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109309

Seq: 1

Gender: F

Reported: 17/06/1996

Weight: 81.00

Hospitalisation:

Age: 58

Onset Date: 20/05/1996

DOB: 02/04/1938

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation		
Injection	5.0 Milligram	Daily	Intravenous
Batch:	Started: 20/05/1996	Stopped: 20/05/1996	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109310 **Seq:** 1 **Gender:** F
Reported: 17/06/1996 **Weight:** 71.00
Hospitalisation: **Age:** 64
Onset Date: 20/05/1996 **DOB:** 14/10/1931
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea			Maxolon

Medicine Details:

ISOFLURANE (Suspected)		Reason: Other disturbance of sensation
	0.0	
Batch:	Started: 20/05/1996	Stopped: 20/05/1996
FENTANYL CITRATE (Suspected)		Reason: Other disturbance of sensation
Injection	100.0 Milligram	1 time Intravenous
Batch:	Started: 20/05/1996	Stopped:
DROPERIDOL (Suspected)		Reason: Other disturbance of sensation
Injection	2.5 Milligram	1 time Intravenous
Batch:	Started: 20/05/1996	Stopped: 20/05/1996
DIPRIVAN (Suspected)		Reason: Other disturbance of sensation
Injection	200.0 Milligram	1 time Intravenous
Batch:	Started: 20/05/1996	Stopped: 20/05/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109310

Seq: 1

Gender: F

Reported: 17/06/1996

Weight: 71.00

Hospitalisation:

Age: 64

Onset Date: 20/05/1996

DOB: 14/10/1931

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

SUXAMETHONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation		
Injection	50.0 Milligram	1 time	Intravenous
Batch:	Started: 20/05/1996	Stopped: 20/05/1996	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109720	Seq: 1	Gender: M
Reported: 04/07/1996		Weight:
Hospitalisation:		Age: 12
Onset Date: 14/06/1996		DOB: 24/04/1984
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Jaw tightness	Benztropine 0.7mg ivi stat only
Hypertonia		Neck stiffness	

Medicine Details:

BISACODYL (Other drug)	Reason:
Suppository	5.0 Milligram 1 time Rectal
Batch:	Started: 11/06/1996 Stopped:
IBUPROFEN (Other drug)	Reason: Pain
	200.0 Milligram As necessary
Batch:	Started: 11/06/1996 Stopped: CONTIN
CEFACLOR MONOHYDRATE (Other drug)	Reason:
	750.0 Milligram Daily
Batch:	Started: 11/06/1996 Stopped: 14/06/1996
DROPERIDOL (Suspected)	Reason: Pain
Injection, intravenous infusion	0.0 As necessary Intravenous
Batch:	Started: 13/06/1996 Stopped: 14/06/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Droperidol 5mg into pca with morphine 60mg 1mg/ml. see original report for other drugs.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109720

Seq: 1

Gender: M

Reported: 04/07/1996

Weight:

Hospitalisation:

Age: 12

Onset Date: 14/06/1996

DOB: 24/04/1984

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

MORPHINE NOS (Suspected)	Reason:	
Injection, intravenous infusion	As necessary Intravenous	
Batch:	Started:	Stopped:
CEPHAZOLIN SODIUM (Suspected)	Reason:	
Injection	1.5 Gram Daily Intravenous	
Batch:	Started: 13/06/1996	Stopped: 2D

Laboratory Investigations:

Additional Information:

Droperidol 5mg into pca with morphine 60mg 1mg/ml. see original report for other drugs.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109798	Seq: 1	Gender: U
Reported: 05/07/1996		Weight: 72.00
Hospitalisation:		Age:
Onset Date: 16/12/1995		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dizziness			
Visual disturbance			

Medicine Details:

PANADOL (Other drug)	Reason: Other disturbance of sensation
Oral application	2.0 Dose Unspecified Daily Oral
Batch:	Started: 14/12/1995 Stopped:
PANADEINE FORTE (Suspected)	Reason: Other disturbance of sensation
Tablet	2.0 Dose Unspecified As necessary Oral
Batch:	Started: 15/12/1995 Stopped: 16/12/1995
MAXOLON (Suspected)	Reason:
Injection	10.0 Milligram As necessary Intramuscular
Batch:	Started: Stopped:
PREMARIN (Suspected)	Reason: Menopausal symptoms
Tablet	625.0 Microgram Daily Oral
Batch:	Started: 16/12/1995 Stopped: 17/12/1995

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Premarin recommenced on 18/12/95 without any further dizziness or blurred vision.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 109798

Seq: 1

Gender: U

Reported: 05/07/1996

Weight: 72.00

Hospitalisation:

Onset Date: 16/12/1995

Age:

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)		Reason:	
	20.0 Milligram	Daily	
Batch:	Started: 13/12/1995	Stopped:	15/12/1995
DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	3.0 Milligram	Daily	Intramuscular
Batch:	Started: 14/12/1995	Stopped:	15/12/1995

Laboratory Investigations:

Additional Information:

Premarin recommenced on 18/12/95 without any further dizziness or blurred vision.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110509 **Seq:** 1 **Gender:** F
Reported: 06/08/1996 **Weight:** 100.00
Hospitalisation: **Age:** 38Y
Onset Date: 30/07/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Pruritus			Promethazine 25mg imi six hourly prn.
Rash erythematous			

Medicine Details:

DROLEPTAN (Suspected)	Reason: Nausea and vomiting
Injection	2.5 Milligram As necessary Intramuscular
Batch:	Started: 30/07/1996 Stopped:
PANADEINE (Suspected)	Reason: Pain
	2.0 Dose Unspecified As necessary
Batch:	Started: 30/07/1996 Stopped:
FORTRAL (Suspected)	Reason: Pain
	45.0 Milligram As necessary
Batch:	Started: 30/07/1996 Stopped:
PAPAVERETUM (Suspected)	Reason: Pain
Injection, intravenous infusion	4.0 Milligram Per hour Intravenous
Batch:	Started: 30/07/1996 Stopped: 30/07/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has previous allergy to pethidine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110509

Seq: 1

Gender: F

Reported: 06/08/1996

Weight: 100.00

Hospitalisation:

Age: 38Y

Onset Date: 30/07/1996

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

NAPROSYN (Suspected)	Reason: Pain		
Suppository	1.0 Gram	Daily	Rectal
Batch:	Started: 30/07/1996	Stopped:	
ESTIGYN (Suspected)	Reason: Menopausal symptoms		
	20.0 Microgram	Daily	
Batch:	Started: 30/07/1996	Stopped:	

Laboratory Investigations:

Additional Information:

Patient has previous allergy to pethidine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110815
Reported: 13/08/1996

Seq: 1

Gender: M
Weight: 80.00
Age: 57
DOB: 31/10/1938
Causality: Causality possible

Hospitalisation:

Onset Date: 29/07/1996
Outcome: Recovered

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Vomiting		Post-op emesis	Maxolon 10mg im at 5:10pm

Medicine Details:

OMNOPON (Suspected)	Reason: Pain		
Injection	20.0 Milligram	1 time	Intramuscular
Batch:	Started: 29/07/1996	Stopped:	
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation		
Injection	200.0 Milligram	1 time	Intravenous
Batch:	Started: 29/07/1996	Stopped:	
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation		
Injection	100.0 Microgram	1 time	Intravenous
Batch:	Started: 29/07/1996	Stopped:	
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 29/07/1996	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110818 **Seq:** 1 **Gender:** F
Reported: 13/08/1996 **Weight:** 90.00
Hospitalisation: Admitted to hospital **Age:** 53
Onset Date: 24/07/1996 **DOB:** 04/10/1942
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea Pain Vomiting		Post-op nausea	

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason:
Injection 100.0 Milligram 2 times Intramuscular	
Batch: Started: 24/07/1996 Stopped:	
MAXOLON (Suspected)	Reason:
Injection 10.0 Milligram 2 times Intramuscular	
Batch: Started: 24/07/1996 Stopped:	
THIOPENTONE SODIUM (Suspected)	Reason: Other disturbance of sensation
Injection 300.0 Milligram 1 time Intravenous	
Batch: Started: 24/07/1996 Stopped:	
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 100.0 Microgram 1 time Intravenous	
Batch: Started: 24/07/1996 Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drugs



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110818 **Seq:** 1 **Gender:** F
Reported: 13/08/1996 **Weight:** 90.00
Hospitalisation: Admitted to hospital **Age:** 53
Onset Date: 24/07/1996 **DOB:** 04/10/1942
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection 1.5 Milligram 1 time Intravenous	
Batch:	Started: 24/07/1996 Stopped:
ATROPINE (Suspected)	Reason: Other disturbance of sensation
Injection 1.2 Milligram 1 time Intravenous	
Batch:	Started: 24/07/1996 Stopped:

Laboratory Investigations:

Additional Information:

See original report for other drugs



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110827 **Seq:** 1 **Gender:** F
Reported: 13/07/1996 **Weight:** 63.00
Hospitalisation: **Age:** 36
Onset Date: 10/07/1996 **DOB:** 14/08/1959
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea		Post-op nausea	Maxolon 10mg imi 10:20am

Medicine Details:

NEOSTIGMINE NOS (Other drug)	Reason: Other disturbance of sensation
2.5 Milligram	1 time
Batch:	Started: 10/07/1996 Stopped:
ATROPINE (Other drug)	Reason: Other disturbance of sensation
1.0 Milligram	1 time
Batch:	Started: 10/07/1996 Stopped:
NITROUS OXIDE (Other drug)	Reason:
1.0 Dose Unspecified	1 time
Batch:	Started: Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection	100.0 Microgram 1 time Intravenous
Batch:	Started: 10/07/1996 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110827
Reported: 13/07/1996
Hospitalisation:
Onset Date: 10/07/1996
Outcome: Recovered

Seq: 1

Gender: F
Weight: 63.00
Age: 36
DOB: 14/08/1959
Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation		
Injection	500.0 Microgram	1 time	Intravenous
Batch:	Started: 10/07/1996	Stopped:	
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation		
	150.0 Milligram	1 time	
Batch:	Started: 10/07/1996	Stopped:	
VECURONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation		
	4.0 Milligram	1 time	
Batch:	Started: 10/07/1996	Stopped:	
ISOFLURANE (Suspected)	Reason: Other disturbance of sensation		
	0.0	1 time	
Batch:	Started: 10/07/1996	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110827

Seq: 1

Gender: F

Reported: 13/07/1996

Weight: 63.00

Hospitalisation:

Age: 36

Onset Date: 10/07/1996

DOB: 14/08/1959

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)		Reason: Pain
Injection	100.0 Milligram	Total Intramuscular
Batch:	Started: 10/07/1996	Stopped: 10/07/1996

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110832

Seq: 1

Gender: F

Reported: 13/08/1996

Weight: 54.00

Hospitalisation:

Onset Date:

Age: 36Y

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dizziness			
Nausea			
Somnolence			

Medicine Details:

FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
100.0 Microgram	1 time
Batch:	Started: 01/09/1995 Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
200.0 Milligram	1 time
Batch:	Started: 01/09/1995 Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
2.0 Milligram	1 time
Batch:	Started: 01/09/1995 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Immunology				Skin tests done and show reactivity to droperidol.

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: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110914 **Seq:** 1 **Gender:** U
Reported: 16/08/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: 13/08/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Unable to speak, jaw "locked"	iv cogentin 1.5ml, iv midazolam 3mg,
Torticollis			

Medicine Details:

PETHIDINE HYDROCHLORIDE (Other drug)	Reason: Pain
Injection 75.0 Milligram As necessary Intramuscular	
Batch: Started: 09/08/1996 Stopped: CONTIN	
PANADEINE FORTE (Other drug)	Reason: Pain
1.0 Dose Unspecified As necessary	
Batch: Started: 11/08/1996 Stopped: CONTIN	
FLAGYL (Other drug)	Reason: Gastritis&duodenitis
Suppository 2.0 Gram Daily Rectal	
Batch: Started: 12/08/1996 Stopped: 13/08/1996	
BUSCOPAN (Other drug)	Reason: Pain
Injection 60.0 Milligram Daily Intramuscular	
Batch: Started: 09/08/1996 Stopped: CONTIN	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

One previous dose of droperidol 12/8/96 with no reaction observed.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110914
Reported: 16/08/1996
Hospitalisation:
Onset Date: 13/08/1996
Outcome: Recovered

Seq: 1

Gender: U
Weight:
Age:
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

MAXOLON (Other drug)	Reason: Nausea and vomiting		
Injection	40.0 Milligram	Daily	Intramuscular
Batch:	Started: 09/08/1996	Stopped:	CONTIN
DROPERIDOL (Suspected)	Reason:		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 13/08/1996	Stopped:	

Laboratory Investigations:

Additional Information:

One previous dose of droperidol 12/8/96 with no reaction observed.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110992 **Seq:** 1 **Gender:** F
Reported: 20/08/1996 **Weight:** 65.00
Hospitalisation: **Age:** 26Y
Onset Date: 13/08/1996 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Locked jaw unable to open mouth + neck stiffness.	Cogentin, midazolam.
Extrapyramidal disorder			

Medicine Details:

BUSCOPAN (Other drug)	Reason: Abdominal pain
80.0 Milligram Total	
Batch:	Started: 09/08/1996 Stopped: 12/08/1996
STEMETIL (Suspected)	Reason: Nausea and vomiting
1.0 Dose Unspecified 1 time	
Batch:	Started: Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
10.0 Milligram Total	
Batch:	Started: 12/08/1996 Stopped: 13/08/1996
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Pain
75.0 Milligram As necessary	
Batch:	Started: 09/08/1996 Stopped: 13/08/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 110992
Reported: 20/08/1996

Seq: 1

Gender: F
Weight: 65.00

Hospitalisation:

Onset Date: 13/08/1996
Outcome: Recovered

Age: 26Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

PANADEINE (Suspected)	Reason: Pain
	1.0 Dose Unspecified As necessary
Batch:	Started: 11/08/1996 Stopped: 13/08/1996
PANADEINE FORTE (Suspected)	Reason: Pain
	2.0 Dose Unspecified As necessary
Batch:	Started: 12/08/1996 Stopped: 15/08/1996
MAXOLON (Suspected)	Reason: Nausea and vomiting
	40.0 Milligram Daily
Batch:	Started: 09/08/1996 Stopped: 12/08/1996

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112232	Seq: 1	Gender: F
Reported: 14/10/1996		Weight:
Hospitalisation:		Age: 44Y
Onset Date: 01/10/1996		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Bentztropine.

Medicine Details:

TEMAZEPAM (Other drug)	Reason:
Oral application	10.0 Milligram As necessary Oral
Batch:	Started: 26/09/1996 Stopped: 02/10/1996
SEREVENT (Other drug)	Reason: Asthma
Inhalation	2.0 Dose Unspecified As necessary Inhalation
Batch:	Started: Stopped:
VENTOLIN (Other drug)	Reason: Asthma
	20.0 Milligram Daily
Batch:	Started: 26/09/1996 Stopped: 02/10/1996
ONDANSETRON HYDROCHLORIDE DIHYDRATE (Other drug)	Reason: Nausea and vomiting
Injection	16.0 Milligram Daily Intravenous
Batch:	Started: 26/09/1996 Stopped: 02/10/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112232
Reported: 14/10/1996
Hospitalisation:
Onset Date: 01/10/1996
Outcome: Recovered

Seq: 1

Gender: F
Weight:
Age: 44Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

HEPARIN SODIUM (Other drug)	Reason: Other prophylactic procedures
Injection	10.0 Thousand Internal Daily Subcutaneous
Batch:	Started: 26/09/1996 Stopped: 02/10/1996
THEO-DUR (Other drug)	Reason: Asthma
Tablet	600.0 Milligram Daily Oral
Batch:	Started: Stopped:
PREDNISONE (Other drug)	Reason: Asthma
Oral application	10.0 Milligram Daily Oral
Batch:	Started: Stopped:
ROCALTROL (Other drug)	Reason:
Capsule	250.0 Microgram Daily Oral
Batch:	Started: Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112232
Reported: 14/10/1996
Hospitalisation:
Onset Date: 01/10/1996
Outcome: Recovered

Seq: 1

Gender: F
Weight:
Age: 44Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

FLIXOTIDE (Other drug)	Reason: Asthma
Inhalation	2.0 Dose Unspecified Daily Inhalation
Batch:	Started: Stopped:
CAPADEX (Other drug)	Reason:
Capsule	8.0 Dose Unspecified Daily Oral
Batch:	Started: Stopped:
DROPERIDOL (Suspected)	Reason:
Injection	2.5 Milligram 1 time Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112341 **Seq:** 1 **Gender:** F
Reported: 17/10/1996 **Weight:**
Hospitalisation: **Age:** 45Y
Onset Date: 09/10/1995 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Paraesthesia		Tingles in hands and toes.	

Medicine Details:

MIDAZOLAM (Suspected)	Reason:
0.0	
Batch:	Started: 09/10/1995 Stopped:
FENTANYL CITRATE (Suspected)	Reason:
0.0	
Batch:	Started: 09/10/1995 Stopped:
DROPERIDOL (Suspected)	Reason:
0.0	
Batch:	Started: 09/10/1995 Stopped:
METOCLOPRAMIDE HYDROCHLORIDE (Suspected)	Reason:
0.0	
Batch:	Started: 09/10/1995 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				All tests and mri scan normal

Additional Information:

Reaction started when patient woke from anaesthetic.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112341

Seq: 1

Gender: F

Reported: 17/10/1996

Weight:

Hospitalisation:

Age: 45Y

Onset Date: 09/10/1995

DOB:

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)		Reason:
	0.0	
Batch:	Started: 09/10/1995	Stopped:
TRACRIUM INJECTABLE (Suspected)		Reason:
	35.0 Milligram	Total
Batch:	Started: 09/10/1995	Stopped:

Laboratory Investigations:

Additional Information:

Reaction started when patient woke from anaesthetic.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112803	Seq: 1	Gender: M
Reported: 01/11/1996		Weight:
Hospitalisation:		Age: 8
Onset Date: 05/10/1994		DOB: 06/11/1985
Outcome: Unknown		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash erythematous		Spreading red areas chest/neck/face	Cepahazolin ceased

Medicine Details:

MIDAZOLAM (Suspected)	0.0	Reason: Premedication
Batch:	Started:	Stopped:
THIOPENTONE SODIUM (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
ATRACURIUM BESYLATE (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
MORPHINE NOS (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 112803

Seq: 1

Gender: M

Reported: 01/11/1996

Weight:

Hospitalisation:

Age: 8

Onset Date: 05/10/1994

DOB: 06/11/1985

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason:		
0.0			
Batch:	Started:	Stopped:	
CEPHAZOLIN SODIUM (Suspected)	Reason:		
Injection	500.0 Milligram	2 times	Intravenous
Batch:	Started: 05/10/1994	Stopped: 06/10/1994	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 113748	Seq: 1	Gender: F
Reported: 06/12/1996		Weight: 50.00
Hospitalisation:		Age: 62Y
Onset Date:		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Coma			
Grand mal convulsion			

Medicine Details:

NAROPIN (Suspected)	Reason:	
400.0 Milligram	1 time	
Batch:	Started:	Stopped:
MIDAZOLAM (Suspected)	Reason:	
Injection 1.0 Milligram	1 time	Intravenous
Batch:	Started:	Stopped:
FENTANYL CITRATE (Suspected)	Reason:	
Injection 50.0 Microgram	1 time	Intravenous
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	Reason:	
Injection 5.0 Milligram	1 time	Intravenous
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 113947 **Seq:** 1 **Gender:** M
Reported: 16/12/1996 **Weight:**
Hospitalisation: **Age:**
Onset Date: **DOB:**
Outcome: Unknown **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			

Medicine Details:

DROPERIDOL (Suspected)	Reason:		
0.0			
Batch:	Started:	Stopped:	5D

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 113968	Seq: 1	Gender: F
Reported: 16/12/1996		Weight: 62.00
Hospitalisation:		Age: 22Y
Onset Date: 04/09/1996		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypoxia Shock		Oxygen saturation dropped to about 60 Peripheral circulatory failure	Ceased minomycin; resuscitation; iv fluids (haemaccel); iv ephedrine 10mg; iv adrenaline 1:1000 diluted to 10ml - 5ml given; iv dexamethasone 4mg; iv lignocaine 50mg
Extrasystoles Ventricular tachycardia			

Medicine Details:

INDOCID (Suspected)	Reason:	
Suppository	0.0	Rectal
Batch:	Started:	Stopped:
PROPOFOL (Suspected)	Reason: Surgery	
	130.0 Milligram	1 time
Batch:	Started:	Stopped:
FENTANYL CITRATE (Suspected)	Reason: Surgery	
	100.0 Microgram	1 time
Batch:	Started: 04/09/1996	Stopped:
DROPERIDOL (Suspected)	Reason: Surgery	
	2.5 Milligram	1 time
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Minomycin 100mg in 500mls of normal saline over 1/2 hour.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 113968
Reported: 16/12/1996

Seq: 1

Gender: F
Weight: 62.00

Hospitalisation:

Onset Date: 04/09/1996
Outcome: Recovered

Age: 22Y
DOB:
Causality: Causality possible

Reaction Details:

Medicine Details:

ATRACURIUM BESYLATE (Suspected)		Reason: Surgery	
	30.0 Milligram	1 time	
Batch:	Started: 04/09/1996	Stopped:	
MINOMYCIN (Suspected)		Reason:	
Injection, intravenous infusion	100.0 Milligram	Total	Intravenous
Batch:	Started: 04/09/1996	Stopped:	

Laboratory Investigations:

Additional Information:

Minomycin 100mg in 500mls of normal saline over 1/2 hour.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 113992

Seq: 1

Gender: M

Reported: 18/12/1996

Weight: 71.00

Hospitalisation:

Age: 16Y

Onset Date: 21/11/1996

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration Diplopia Tremor		Upper rotation of eyes	

Medicine Details:

ZOFRAN (Suspected)	Reason:
Injection	4.0 Milligram 1 time Intravenous
Batch:	Started: 21/11/1996 Stopped:
DROLEPTAN (Suspected)	Reason: Nausea and vomiting
Injection, intravenous infusion	5.0 Milligram Total Intravenous
Batch:	Started: 20/11/1996 Stopped: 21/11/1996

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 114424	Seq: 1	Gender: F
Reported: 08/01/1997		Weight: 65.00
Hospitalisation:		Age: 19Y
Onset Date: 25/08/1996		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder		Extrapyramidal disorder	Benztropine 2mg 9v 2230 25/8/96.
Rash maculo-papular		Maculo-papular rash on neck and back	

Medicine Details:

PANADOL (Other drug)	Reason:
0.0	
Batch:	Started: Stopped:
ATIVAN (Other drug)	Reason:
Tablet	1.0 Milligram As necessary Sublingual
Batch:	Started: Stopped:
ZYCLIR (Other drug)	Reason:
Injection	750.0 Milligram Daily Intravenous
Batch:	Started: 16/08/1996 Stopped:
DIFLUCAN (Other drug)	Reason: Prophylaxis
Injection	400.0 Milligram Daily Intravenous
Batch:	Started: 16/08/1996 Stopped: CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 114424

Seq: 1

Gender: F

Reported: 08/01/1997

Weight: 65.00

Hospitalisation:

Age: 19Y

Onset Date: 25/08/1996

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

STEMETIL (Suspected)		Reason:	
Injection	12.5 Milligram	2 times	Intravenous
Batch:	Started: 24/08/1996	Stopped:	25/08/1996
DROLEPTAN (Suspected)		Reason:	
Injection	2.5 Milligram	1 time	Intravenous
Batch:	Started: 19/08/1996	Stopped:	25/08/1996

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 115265 **Seq:** 1 **Gender:** F
Reported: 14/02/1997 **Weight:**
Hospitalisation: **Age:** 26
Onset Date: 25/07/1996 **DOB:** 07/02/1970
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Angioedema			Adrenaline, hydrocortisone, ventilated, intubated, paralysed & o/night icu.

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection 2.5 Milligram 1 time Intramuscular	
Batch: Started: Stopped:	
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Pain
Injection 75.0 Milligram 1 time Intramuscular	
Batch: Started: Stopped:	
TRIMETHOPRIM (Suspected)	Reason:
Oral application 300.0 Milligram 1 time Oral	
Batch: Started: 25/07/1996 Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Previous history of dystonic reactions to metaclopramide, promethazine and prochlorperizine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 115266	Seq: 1	Gender: F
Reported: 14/02/1997		Weight:
Hospitalisation:		Age: 24
Onset Date: 07/03/1995		DOB: 24/10/1970
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine 2mg im, droperidol ceased.

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting
Injection	2.0 Milligram	Total Intravenous
Batch:	Started: 07/03/1995	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred 165 minutes after receiving second dose. has had similar reactions to metoclopramide and prochlorperazine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 116853

Seq: 1

Gender: F

Reported: 18/04/1997

Weight: 12.00

Hospitalisation:

Age: 3Y

Onset Date: 10/04/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine 0.25mg im bd

Medicine Details:

DROPERIDOL (Suspected)	Reason: Pain		
Injection	3.6 Milligram	Total	Intramuscular
Batch:	Started: 02/04/1997	Stopped:	10/04/1997
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation		
Injection	12.5 Milligram	Daily	Intravenous
Batch:	Started: 10/04/1997	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 116915

Seq: 1

Gender: M

Reported: 18/04/1997

Weight: 31.00

Hospitalisation:

Age: 8Y

Onset Date: 21/03/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Diazepam

Medicine Details:

DROPERIDOL (Suspected)	Reason:
3.0 Milligram	Daily
Batch:	Started: 21/03/1997
	Stopped: 21/03/1997
PETHIDINE HYDROCHLORIDE (Suspected)	Reason:
30.0 Milligram	Daily
Batch:	Started: 21/03/1997
	Stopped: 21/03/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 116918

Seq: 1

Gender: F

Reported: 18/04/1997

Weight: 16.00

Hospitalisation:

Onset Date: 02/04/1997

Age: 3Y

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			

Medicine Details:

DROPERIDOL (Suspected)	Reason:
1.5 Milligram	1 time
Batch:	Started: 02/04/1997 Stopped: 02/04/1997
PETHIDINE HYDROCHLORIDE (Suspected)	Reason:
1.5 Milligram	1 time
Batch:	Started: 02/04/1997 Stopped: 02/04/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 117716 **Seq:** 1 **Gender:** M
Reported: 20/05/1997 **Weight:**
Hospitalisation: **Age:** 31Y
Onset Date: 15/05/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Anxiety Dystonia Hypertonia	Severe		Benztropine

Medicine Details:

LIGNOCAINE (Suspected)	Reason:
0.0	
Batch:	Started: 15/05/1997 Stopped: 15/05/1997
ETHRANE (Suspected)	Reason:
0.0	
Batch:	Started: 15/05/1997 Stopped: 15/05/1997
DIPRIVAN (Suspected)	Reason:
200.0 Milligram 1 time	
Batch:	Started: 15/05/1997 Stopped: 15/05/1997
DROLEPTAN (Suspected)	Reason:
1.5 Milligram 1 time	
Batch:	Started: 15/05/1997 Stopped: 15/05/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 117716
Reported: 20/05/1997

Seq: 1

Gender: M

Weight:

Hospitalisation:

Age: 31Y

Onset Date: 15/05/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

ATROPINE (Suspected)	Reason:
600.0 Microgram	1 time
Batch:	Started: 15/05/1997 Stopped: 15/05/1997
TORADOL (Suspected)	Reason:
30.0 Milligram	1 time
Batch:	Started: 15/05/1997 Stopped: 15/05/1997

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 117857

Seq: 1

Gender: M

Reported: 26/05/1997

Weight:

Hospitalisation:

Onset Date: 19/02/1997

Age: 31

DOB: 23/04/1965

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypertonia Anxiety Dyspnoea Tongue oedema		Muscle spasm in neck and face.	Oxygen, diazepam.

Medicine Details:

EPOETIN ALFA (Other drug) Injection Batch:	8.0 Thousand Internal Weekly Started: L TERM	Reason: Chronic nephritis	Subcutaneous Stopped:
COLCHICINE (Other drug) Batch:	500.0 Microgram Started: 17/02/1997	Reason: Gout	Daily Stopped:
ZANTAC (Other drug) Tablet Batch:	1.0 Gram Started: L TERM	Reason:	Daily Oral Stopped:
NORVASC (Other drug) Tablet Batch:	40.0 Milligram Started: L TERM	Reason: Essential benign hypertension	Daily Oral Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking tritace.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 117857
Reported: 26/05/1997

Seq: 1

Gender: M

Weight:

Hospitalisation:

Onset Date: 19/02/1997

Age: 31

Outcome: Recovered

DOB: 23/04/1965

Causality: Causality probable

Reaction Details:

Medicine Details:

TRITACE (Other drug)	Reason: Essential benign hypertension
Capsule	5.0 Milligram Daily Oral
Batch:	Started: L TERM Stopped:
CALTRATE (Other drug)	Reason: Other specified symptoms nec
Tablet	1.0 Dose Unspecified Daily Oral
Batch:	Started: L TERM Stopped:
VITAMIN B COMPLEX (Other drug)	Reason:
Oral application	2.0 Dose Unspecified Daily Oral
Batch:	Started: L TERM Stopped:
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	7.5 Milligram Daily Subcutaneous
Batch:	Started: 18/02/1997 Stopped: 19/02/1997

Laboratory Investigations:

Additional Information:

Patient also taking tritace.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 118506	Seq: 1	Gender: F
Reported: 24/06/1997		Weight: 61.00
Hospitalisation:		Age: 13Y
Onset Date: 07/06/1997		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Acute dystonia	Benztropine 1mg im given orally
Oculogyration			
Opisthotonus			

Medicine Details:

DROPERIDOL (Suspected)		Reason: Depression	
Injection	0.0	As necessary	Intramuscular
Batch:	Started: 06/06/1997	Stopped:	07/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 118725	Seq: 1	Gender: F
Reported: 01/07/1997		Weight:
Hospitalisation:		Age: 47Y
Onset Date: 03/06/1997		DOB:
Outcome: Not yet recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration			Benztropine.

Medicine Details:

DOMPERIDONE (Other drug)		Reason: Vertigo	
Tablet	10.0 Milligram	As necessary	Oral
Batch:	Started: 01/06/1997	Stopped: 02/06/1997	
PROMETHAZINE HYDROCHLORIDE (Other drug)		Reason:	
Oral application	20.0 Milligram	As necessary	Oral
Batch:	Started: 31/05/1997	Stopped:	
PETHIDINE HYDROCHLORIDE (Other drug)		Reason: Pain	
Injection	100.0 Milligram	As necessary	Intramuscular
Batch:	Started: 31/05/1997	Stopped:	
PANADEINE FORTE (Other drug)		Reason: Pain	
Tablet	1.0 Dose Unspecified	As necessary	Oral
Batch:	Started: 31/05/1997	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 118725

Seq: 1

Gender: F

Reported: 01/07/1997

Weight:

Hospitalisation:

Age: 47Y

Onset Date: 03/06/1997

DOB:

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DUPHALAC (Other drug)		Reason:	
Oral Liquid	30.0 Millilitre	Daily	Oral
Batch:	Started:	Stopped:	
PREMARIN (Other drug)		Reason:	
Tablet	1.2 Milligram	Daily	Oral
Batch:	Started:	Stopped:	
CHLOROMYCETIN (Other drug)		Reason:	
	3.0 Dose Unspecified	Daily	
Batch:	Started: 03/05/1997	Stopped:	
CEFTAZIDIME (Other drug)		Reason: Other cellulitis and abscess	
Injection	4.0 Gram	Daily	Intravenous
Batch:	Started: 31/05/1997	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119047	Seq: 1	Gender: F
Reported: 14/07/1997		Weight:
Hospitalisation:		Age: 30
Onset Date: 07/07/1997		DOB: 01/04/1967
Outcome: Not yet recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea	Severe	Severe nausea postoperatively	
Vomiting	Severe	Severe vomiting postoperatively	

Medicine Details:

PROPOFOL (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
ATRACURIUM BESYLATE (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
DROPERIDOL (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:
KEFZOL (Suspected)	0.0	Reason:
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119047 **Seq:** 1
Reported: 14/07/1997

Gender: F

Weight:

Hospitalisation:

Onset Date: 07/07/1997
Outcome: Not yet recovered

Age: 30

DOB: 01/04/1967

Causality: Causality possible

Reaction Details:

Medicine Details:

FLAGYL (Suspected)	Reason:
0.0	
Batch:	Started: Stopped:
NITROUS OXIDE (Suspected)	Reason:
0.0	
Batch:	Started: Stopped:
ISOFLURANE (Suspected)	Reason:
0.0	
Batch:	Started: Stopped:
MORPHINE NOS (Suspected)	Reason: Prophylaxis
Injection, intravenous infusion	48.0 Milligram Daily Intravenous
Batch:	Started: 07/07/1997 Stopped: 08/07/1997

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119204 **Seq:** 1 **Gender:** F
Reported: 17/07/1997 **Weight:** 83.00
Hospitalisation: Hospitalisation prolonged **Age:** 32
Onset Date: 24/06/1997 **DOB:** 23/08/1964
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea			Sleep.

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection 0.0 Intravenous	
Batch: Started: 24/06/1997 Stopped: 24/06/1997	
SEVORANE (Suspected)	Reason: Other disturbance of sensation
Injection 0.0 Intravenous	
Batch: Started: 24/06/1997 Stopped: 24/06/1997	
PANADEINE FORTE (Suspected)	Reason: Pain
Tablet 2.0 Dose Unspecified 1 time Oral	
Batch: Started: 24/06/1997 Stopped: 24/06/1997	
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 100.0 Milligram 1 time Intravenous	
Batch: Started: 24/06/1997 Stopped: 24/06/1997	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Oral analgesia given on an empty stomach



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119204 **Seq:** 1

Reported: 17/07/1997

Hospitalisation: Hospitalisation prolonged

Onset Date: 24/06/1997

Outcome: Recovered

Gender: F

Weight: 83.00

Age: 32

DOB: 23/08/1964

Causality: Causality possible

Reaction Details:

Medicine Details:

DIPRIVAN (Suspected)	Reason: Other disturbance of sensation		
Injection	180.0 Milligram	1 time	Intravenous
Batch:	Started: 24/06/1997	Stopped: 24/06/1997	

Laboratory Investigations:

Additional Information:

Oral analgesia given on an empty stomach



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119210 **Seq:** 1 **Gender:** F
Reported: 17/07/1997 **Weight:** 97.00
Hospitalisation: Hospitalisation prolonged **Age:** 46
Onset Date: 12/06/1997 **DOB:** 25/11/1950
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Vomiting		Post op emesis	Maxolon, ondansetron

Medicine Details:

SEVOFLURANE (Suspected)		Reason: Other disturbance of sensation
	0.0	
Batch:	Started: 12/06/1997	Stopped: 12/06/1997
NEOSTIGMINE NOS (Suspected)		Reason: Other disturbance of sensation
Injection	0.0	Intravenous
Batch:	Started: 12/06/1997	Stopped: 12/06/1997
FENTANYL CITRATE (Suspected)		Reason: Other disturbance of sensation
Injection	100.0 Milligram	1 time Intravenous
Batch:	Started: 12/06/1997	Stopped: 12/06/1997
DROPERIDOL (Suspected)		Reason: Other disturbance of sensation
Injection	500.0 Microgram	1 time Intravenous
Batch:	Started: 12/06/1997	Stopped: 12/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119210 **Seq:** 1 **Gender:** F
Reported: 17/07/1997 **Weight:** 97.00
Hospitalisation: Hospitalisation prolonged **Age:** 46
Onset Date: 12/06/1997 **DOB:** 25/11/1950
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Medicine Details:

DIPRIVAN (Suspected)	Reason: Other disturbance of sensation
Injection 120.0 Milligram 1 time Intravenous	
Batch:	Started: 12/06/1997 Stopped: 12/06/1997
ATROPINE (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Dose Unspecified 1 time Intravenous	
Batch:	Started: 12/06/1997 Stopped: 12/06/1997
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Pain
Injection 75.0 Milligram Total Intravenous	
Batch:	Started: 12/06/1997 Stopped: 12/06/1997

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119222 **Seq:** 1 **Gender:** F
Reported: 17/07/1997 **Weight:**
Hospitalisation: Hospitalisation prolonged **Age:** 31
Onset Date: 16/06/1997 **DOB:** 02/06/1966
Outcome: Unknown **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Nausea		Post op nausea	Maxolon

Medicine Details:

ISOFLURANE (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 16/06/1997 Stopped: 16/06/1997
PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Pain
Injection 75.0 Milligram 1 time Intramuscular	
Batch:	Started: 16/01/1997 Stopped: 16/06/1997
ROCURONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation
Injection 40.0 Milligram 1 time Intravenous	
Batch:	Started: 16/06/1997 Stopped: 16/06/1997
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 75.0 Milligram 1 time Intravenous	
Batch:	Started: 16/06/1997 Stopped: 16/06/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 119222 **Seq:** 1 **Gender:** F
Reported: 17/07/1997 **Weight:**
Hospitalisation: Hospitalisation prolonged **Age:** 31
Onset Date: 16/06/1997 **DOB:** 02/06/1966
Outcome: Unknown **Causality:** Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection 1.5 Milligram 1 time Intravenous	
Batch: Started: 16/06/1997 Stopped: 16/06/1997	
ATROPINE (Suspected)	Reason: Other disturbance of sensation
Injection 1.6 Milligram 1 time Intravenous	
Batch: Started: 16/06/1997 Stopped: 16/06/1997	
NEOSTIGMINE NOS (Suspected)	Reason: Other disturbance of sensation
Injection 3.5 Milligram 1 time Intravenous	
Batch: Started: 16/06/1997 Stopped: 16/06/1997	
THIOPENTONE SODIUM (Suspected)	Reason: Other disturbance of sensation
Injection 425.0 Milligram Daily Intravenous	
Batch: Started: 16/06/1997 Stopped: 16/06/1997	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 120877

Seq: 1

Gender: F

Reported: 16/09/1997

Weight: 68.00

Hospitalisation:

Age: 24Y

Onset Date: 24/12/1996

DOB:

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dyspnoea		Couldnt breathe properly	

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	2.5 Milligram	1 time	Intramuscular
Batch:	Started: 24/12/1996	Stopped: 24/12/1996	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 121866

Seq: 1

Gender: M

Reported: 27/10/1997

Weight: 61.00

Hospitalisation:

Age: 67Y

Onset Date: 22/10/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Pyrexia Confusional state Hypertonia Neuroleptic malignant syndrome		Temp 38 oc	Iv dantrolene and bromocryptine

Medicine Details:

ISOFLURANE (Suspected)	Reason:		
0.0			
Batch:	Started:	Stopped:	
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation		
Injection	1.5 Milligram	1 time	Intravenous
Batch:	Started: 21/10/1997	Stopped:	
PROPOFOL (Suspected)	Reason: Other disturbance of sensation		
Injection	100.0 Milligram	1 time	Intravenous
Batch:	Started:	Stopped:	
DICLOXACILLIN SODIUM (Suspected)	Reason:		
Injection	2.0 Gram	1 time	Intravenous
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

History of previous anaesthetics without problems, has history of hypertension



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 121866

Seq: 1

Gender: M

Reported: 27/10/1997

Weight: 61.00

Hospitalisation:

Age: 67Y

Onset Date: 22/10/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

GENTAMICIN SULPHATE (Suspected)	Reason:
Injection 180.0 Milligram	1 time Intravenous
Batch:	Started: Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 162.5 Milligram	Total Intravenous
Batch:	Started: Stopped:
ATRACURIUM BESYLATE (Suspected)	Reason: Other disturbance of sensation
Injection, intravenous infusion 10.0 Milligram	Per hour Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Additional Information:

History of previous anaesthetics without problems, has history of hypertension



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 122742 **Seq:** 1 **Gender:** F
Reported: 24/11/1997 **Weight:**
Hospitalisation: **Age:** 60Y
Onset Date: 15/10/1997 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Convulsion			

Medicine Details:

MIDAZOLAM (Suspected)	Reason: Other disturbance of sensation
2.5 Milligram	2 times
Batch:	Started: Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
100.0 Milligram	1 time
Batch:	Started: Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
15.0 Milligram	1 time
Batch:	Started: Stopped:
BUPIVACAINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation
75.0 Milligram	1 time
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 122742

Seq: 1

Gender: F

Reported: 24/11/1997

Weight:

Hospitalisation:

Age: 60Y

Onset Date: 15/10/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)

Reason: Other disturbance of sensation

50.0 Milligram

1 time

Batch:

Started:

Stopped:

ISOFLURANE (Suspected)

Reason: Other disturbance of sensation

1.0 Percent

1 time

Batch:

Started:

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 122744	Seq: 1	Gender: F
Reported: 25/11/1997		Weight:
Hospitalisation:		Age: 45Y
Onset Date: 14/11/1997		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Tongue disorder		Ocomotor spasm Tongue rolled back	Benztropine 4mg iv

Medicine Details:

MORPHINE NOS (Suspected)	Reason:
0.0	
Batch:	Started:
	Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
5.0 Milligram	Daily
Batch:	Started: 13/11/1997
	Stopped: 14/11/1997

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient has previous history to maxolon, stemetil and other drugs.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 129669	Seq: 1	Gender: M
Reported: 28/07/1998		Weight:
Hospitalisation:		Age: 21
Onset Date: 06/04/1998		DOB: 17/06/1976
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Hypertonia Opisthotonus Tachycardia			Benztropine iv 1mg.

Medicine Details:

PANADOL (Other drug)	Reason: Pain
0.0	
Batch:	Started: 04/04/1998 Stopped:
PANADEINE (Other drug)	Reason: Pain
0.0	
Batch:	Started: 04/04/1998 Stopped:
ONDANSETRON HYDROCHLORIDE DIHYDRATE (Other drug)	Reason: Nausea and vomiting
Tablet	8.0 Milligram 1 time Oral
Batch:	Started: 06/04/1998 Stopped:
VENTOLIN (Other drug)	Reason:
Inhalation	0.0 As necessary Inhalation
Batch:	Started: 05/04/1998 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 129669
Reported: 28/07/1998

Seq: 1

Gender: M

Weight:

Hospitalisation:

Onset Date: 06/04/1998
Outcome: Recovered

Age: 21

DOB: 17/06/1976

Causality: Causality possible

Reaction Details:

Medicine Details:

CLINDAMYCIN HYDROCHLORIDE (Other drug)	Reason: Otr&nos infec¶sit diseases		
Oral application	1.2 Gram	Daily	Oral
Batch:	Started: 04/04/1998	Stopped:	
DICLOXACILLIN SODIUM (Other drug)	Reason: Otr&nos infec¶sit diseases		
Injection	4.0 Gram	Daily	Intravenous
Batch:	Started: 04/04/1998	Stopped:	
METOCLOPRAMIDE HYDROCHLORIDE (Other drug)	Reason: Nausea and vomiting		
Injection	40.0 Milligram	Daily	Intravenous
Batch:	Started: 04/04/1998	Stopped: 05/04/1998	
DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
	0.0		
Batch:	Started: 03/04/1998	Stopped: 06/04/1998	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 130780 **Seq:** 1 **Gender:** F
Reported: 03/09/1998 **Weight:**
Hospitalisation: **Age:** 49
Onset Date: 17/08/1998 **DOB:** 05/09/1948
Outcome: Recovered **Causality:** Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Tongue oedema		Pulling and swelling at back of tongue.	

Medicine Details:

DIHYDROERGOTAMINE (Other drug)	Reason: Migraine		
Injection	1.0 Milligram	1 time	Intravenous
Batch:	Started: 17/08/1998	Stopped:	
DROPERIDOL (Suspected)	Reason: Nausea and vomiting		
Injection	500.0 Microgram	1 time	Intravenous
Batch:	Started: 17/08/1998	Stopped: 17/08/1998	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 130789	Seq: 1	Gender: M
Reported: 03/09/1998		Weight:
Hospitalisation:		Age: 2Y
Onset Date: 30/07/1998		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Opisthotonus Dystonia		Hyperextension of neck	Cogentin iv 2 doses.

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
0.0	
Batch:	Started: 30/07/1998 Stopped: 30/07/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 131602	Seq: 1	Gender: F
Reported: 30/09/1998		Weight: 98.00
Hospitalisation:		Age: 49Y
Onset Date: 09/07/1998		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypotension Cyanosis Flushing		Extreme hypotention	Pt was resuscitated using adrenaline 0.5mg x oxygen and iv fluids.

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason: Other disturbance of sensation
50.0 Milligram	1 time
Batch:	Started: 09/07/1998 Stopped: 09/07/1998
DROLEPTAN (Suspected)	Reason:
1.0 Milligram	1 time
Batch:	Started: 09/07/1998 Stopped: 09/07/1998
HYPNOVEL (Suspected)	Reason: Other disturbance of sensation
2.0 Milligram	1 time
Batch:	Started: 09/07/1998 Stopped: 09/07/1998
ESMERON (Suspected)	Reason:
60.0 Milligram	1 time
Batch:	Started: 09/07/1998 Stopped: 09/07/1998

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Pt has allergies to penicillin, amoxil, morphine & furadantin.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 131602

Seq: 1

Gender: F

Reported: 30/09/1998

Weight: 98.00

Hospitalisation:

Age: 49Y

Onset Date: 09/07/1998

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

ZOFRAN (Suspected)

Reason:

200.0 Milligram

Daily

Batch:

Started: 09/07/1998

Stopped: 09/07/1998

Laboratory Investigations:

Additional Information:

Pt has allergies to penicillin, amoxil, morphine & furadantin.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 134787	Seq: 1	Gender: U
Reported: 04/01/1999		Weight: 40.00
Hospitalisation:		Age: 13
Onset Date: 01/07/1997		DOB: 03/02/1984
Outcome: Unknown		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration Visual disturbance			

Medicine Details:

PANADEINE (Other drug)	Reason:
0.0	
Batch:	Started: 28/06/1997 Stopped:
AMOXYCILLIN TRIHYDRATE (Other drug)	Reason:
Injection	4.0 Gram
	Daily
	Intravenous
Batch:	Started: 28/06/1997 Stopped:
METRONIDAZOLE (Other drug)	Reason:
Injection	900.0 Milligram
	Daily
	Intravenous
Batch:	Started: 25/06/1997 Stopped:
INDOCID (Other drug)	Reason:
Suppository	200.0 Milligram
	Daily
	Rectal
Batch:	Started: 30/06/1997 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 134787

Seq: 1

Gender: U

Reported: 04/01/1999

Weight: 40.00

Hospitalisation:

Age: 13

Onset Date: 01/07/1997

DOB: 03/02/1984

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	1.0 Milligram	1 time	Intravenous
Batch:	Started: 01/07/1997	Stopped:	01/07/1997
PROCHLORPERAZINE MALEATE (Suspected)		Reason:	
Injection	15.0 Milligram	Daily	Intravenous
Batch:	Started: 29/06/1997	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 134855	Seq: 1	Gender: F
Reported: 04/01/1999		Weight: 18.00
Hospitalisation:		Age: 8
Onset Date: 22/05/1998		DOB: 30/01/1990
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hallucination Depersonalisation Urinary retention		Stating she was another person.	

Medicine Details:

MORPHINE NOS (Other drug)		Reason: Cong anom of spine
Injection	10.0 Milligram	Daily Intravenous
Batch:	Started: 20/05/1998	Stopped:
DIAZEPAM (Suspected)		Reason: Cong anom of spine
Tablet	2.0 Milligram	1 time Oral
Batch:	Started: 22/05/1998	Stopped:
GENTAMICIN SULPHATE (Suspected)		Reason:
Injection	50.0 Milligram	1 time Intravenous
Batch:	Started: 22/05/1998	Stopped:
DROPERIDOL (Suspected)		Reason: Cong anom of spine
Injection	1.2 Milligram	Daily Intravenous
Batch:	Started: 21/05/1998	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 134855

Seq: 1

Gender: F

Reported: 04/01/1999

Weight: 18.00

Hospitalisation:

Age: 8

Onset Date: 22/05/1998

DOB: 30/01/1990

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

PANADOL (Suspected)

Reason: Cong anom of spine

Suppository

1.5 Gram

Daily

Rectal

Batch:

Started: 20/05/1998

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 135175 **Seq:** 1 **Gender:** F
Reported: 14/01/1999 **Weight:**
Hospitalisation: **Age:** 30Y
Onset Date: 01/10/1998 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash		Generalised rash over body	Oxygen given via hudson mask, im phenergan 25mg.
Hypoxia			

Medicine Details:

Keflin Neutral (Suspected)	Reason:
Injection 1.0 Gram 1 time Intravenous	
Batch:	Started: 01/10/1998 Stopped:
FENTANYL CITRATE (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Dose Unspecified 1 time Intravenous	
Batch:	Started: 01/10/1998 Stopped:
MIDAZOLAM (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Dose Unspecified 1 time Intravenous	
Batch:	Started: 01/10/1998 Stopped:
DIPRIVAN (Suspected)	Reason: Other disturbance of sensation
Injection 1.0 Dose Unspecified 1 time Intravenous	
Batch:	Started: 01/10/1998 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Other data				Saturation oxygen decreased to 89%.

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 135175

Seq: 1

Gender: F

Reported: 14/01/1999

Weight:

Hospitalisation:

Age: 30Y

Onset Date: 01/10/1998

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROLEPTAN (Suspected)

Reason: Other disturbance of sensation

Injection

1.0 Dose Unspecified 1 time

Intravenous

Batch:

Started: 01/10/1998

Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 137338

Seq: 1

Gender: M

Reported: 16/03/1999

Weight:

Hospitalisation:

Onset Date: 22/10/1998

Age: 23

DOB: 15/11/1974

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Leukopenia			
Neutropenia			

Medicine Details:

MIDAZOLAM (Suspected)	Reason:		
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 21/10/1998	Stopped:	
DROPERIDOL (Suspected)	Reason:		
Injection	10.0 Milligram	1 time	Intramuscular
Batch:	Started: 20/10/1998	Stopped:	
TEMAZEPAM (Suspected)	Reason:		
	10.0 Milligram	As necessary	
Batch:	Started:	Stopped:	
EFEXOR (Suspected)	Reason: Depression		
Tablet	75.0 Milligram	Daily	Oral
Batch:	Started: 21/10/1998	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Neutrophils		22/10/1998	0.99	
	White blood cells		22/10/1998	2.62	

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 137338

Seq: 1

Gender: M

Reported: 16/03/1999

Weight:

Hospitalisation:

Age: 23

Onset Date: 22/10/1998

DOB: 15/11/1974

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

DIAZEPAM (Suspected)	Reason: Anxiety neurosis		
Tablet	20.0 Milligram	Daily	Oral
Batch:	Started: 21/10/1998	Stopped:	
THIAMINE (Suspected)	Reason:		
Oral application	100.0 Milligram	Daily	Oral
Batch:	Started: 21/10/1998	Stopped:	
FLUOXETINE HYDROCHLORIDE (Suspected)	Reason:		
	40.0 Milligram	Daily	
Batch:	Started: 20/08/1998	Stopped: 20/10/1998	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 138864 **Seq:** 1 **Gender:** M
Reported: 29/04/1999 **Weight:**
Hospitalisation: Admitted to hospital **Age:** 2
Onset Date: 30/07/1998 **DOB:** 22/07/1996
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash Dystonia		Rash upper chest	Benztropine 0.24mg iv (0.02mg/kg).

Medicine Details:

DROPERIDOL (Suspected)	Reason: Premedication
Injection	3.0 Milligram 1 time Intramuscular
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No known allergies. see original report for details of other drug - paracetamol. date of recovery: 30/7/98.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 139532 **Seq:** 1 **Gender:** F
Reported: 14/05/1999 **Weight:**
Hospitalisation: **Age:** 28Y
Onset Date: 14/02/1999 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia			Ceased droperidol, given benztropine iv x 4.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	1.0 Milligram Daily Intramuscular
Batch:	Started: 14/02/1999 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

See original report for other drug details - methadone, dicloxacillin iv & dicloxacillin oral. previous allergies to metoclopramid & haloperidol. four doses of stemetil 12.5mg im given but proved ineffective.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 139883	Seq: 1	Gender: M
Reported: 24/05/1999		Weight:
Hospitalisation:		Age: 21Y
Onset Date: 12/05/1999		DOB:
Outcome: Recovered without treatment		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Delirium			
Vomiting			

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	10.0 Milligram	2 times	Intramuscular
Batch:	Started:	Stopped:	
CLOZARIL (Suspected)		Reason: Unspecified schizophrenia	
Tablet	350.0 Milligram	Daily	Oral
Batch:	Started: 14/04/1999	Stopped:	CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Please see original report for details of other drugs - thioridazine. patient has a history of aggressiveness



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 142893 **Seq:** 1 **Gender:** M
Reported: 23/08/1999 **Weight:**
Hospitalisation: **Age:** 26
Onset Date: 18/07/1999 **DOB:** 17/12/1972
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Dystonic reaction	Treated with cogentin 2mg orally.
Agitation		Motor restlessness	

Medicine Details:

ZUCLOPENTHIXOL DECANOATE (Suspected)	Reason:
200.0 Milligram	Weekly
Batch:	Started: 12/07/1999 Stopped: CONTIN
THIORIDAZINE HYDROCHLORIDE (Suspected)	Reason:
50.0 Milligram	As necessary
Batch:	Started: 16/07/1999 Stopped: CONTIN
DROPERIDOL (Suspected)	Reason:
5.0 Milligram	As necessary
Batch:	Started: 16/07/1999 Stopped: CONTIN
DIAZEPAM (Suspected)	Reason:
5.0 Milligram	As necessary
Batch:	Started: 16/07/1999 Stopped: CONTIN

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reduced thioridazine dose to maximum 600mg/d,



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 142893

Seq: 1

Gender: M

Reported: 23/08/1999

Weight:

Hospitalisation:

Age: 26

Onset Date: 18/07/1999

DOB: 17/12/1972

Outcome: Not yet recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

COGENTIN (Suspected)	Reason:
	0.5 Milligram As necessary
Batch:	Started: 16/07/1999
	Stopped: CONTIN

Laboratory Investigations:

Additional Information:

Reduced thioridazine dose to maximum 600mg/d,



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 150478

Seq: 1

Gender: F

Reported: 17/03/2000

Weight: 85.00

Hospitalisation:

Age: 56Y

Onset Date: 22/02/2000

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Angina pectoris Chest pain Electrocardiogram abnormal Myocardial infarction			

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	2.5 Milligram Daily Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Electrocardiograph Immunology Radiology				Marked st elevation inferior lead. Troponin i = 3.3ng/l. (<0.1) Subsequent normal coronary angiogram.

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 150535
Reported: 21/03/2000
Hospitalisation:
Onset Date: 25/11/1999
Outcome: Recovered

Seq: 1

Gender: F
Weight: 56.00
Age: 22
DOB: 27/11/1976
Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Flushing	Life threatening	Generalised skin flush.	Adrenaline and supportive measures. 8 hours icu.
Bronchospasm	Life threatening		
Hypotension	Life threatening		
Tachycardia	Life threatening		

Medicine Details:

MORPHINE NOS (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: 25/11/1999 Stopped:
DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: 25/11/1999 Stopped:
PROPOFOL (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: 25/11/1999 Stopped:
LIGNOCAINE (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: 25/11/1999 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
	Immunology				Mast cell tryptase (>200). skin tests positive to rocuronium. also positive to pancuronium, vecuronium and morphine (normal).

Additional Information:

All drugs given as single dose all within 2 minutes prior to reaction. date of recovery: 26/11/1999.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 150535

Seq: 1

Gender: F

Reported: 21/03/2000

Weight: 56.00

Hospitalisation:

Age: 22

Onset Date: 25/11/1999

DOB: 27/11/1976

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

ROCURONIUM BROMIDE (Suspected)

Reason: Other disturbance of sensation

Injection

1.0 Dose Unspecified 1 time

Intravenous

Batch:

Started: 25/11/1999

Stopped: 25/11/1999

Laboratory Investigations:

Additional Information:

All drugs given as single dose all within 2 minutes prior to reaction. date of recovery: 26/11/1999.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 150746

Seq: 1

Gender: M

Reported: 24/03/2000

Weight: 118.00

Hospitalisation:

Onset Date: 25/12/1999

Age: 37Y

Outcome: Recovered

DOB:

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Agitation			
Confusional state			
Hypertension			
Hypertonia			
Tachycardia			
Tremor			

Medicine Details:

DROPERIDOL (Suspected)	Reason:		
Injection	10.0 Milligram	As necessary	Intramuscular
Batch:	Started: 23/12/1999	Stopped: 26/12/1999	
TIMENTIN (Suspected)	Reason:		
Injection	18.6 Gram	Daily	Intravenous
Batch:	Started: 17/12/1999	Stopped: 27/12/1999	
RANITIDINE (Suspected)	Reason:		
Injection	150.0 Milligram	Daily	Intravenous
Batch:	Started: 23/12/1999	Stopped: 26/12/1999	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking paracetamol, heparin, nystatin, vancomycin, mianserin, metoclopramide, tramadol. p/h of reaction to stemetil - tardive dyskinesia.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 154154

Seq: 1

Gender: F

Reported: 23/06/2000

Weight: 70.00

Hospitalisation:

Age: 60

Onset Date: 16/06/2000

DOB: 29/01/1940

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Urticaria			Ceased pethidine and droperidol. given phenergan.

Medicine Details:

PETHIDINE HYDROCHLORIDE (Suspected)	Reason:
0.0	
Batch:	Started: 15/06/2000 Stopped: 16/06/2000
DROPERIDOL (Suspected)	Reason:
0.0	
Batch:	Started: 15/06/2000 Stopped: 16/06/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking voltaren, paracetamol. patient recovered 16/6/00.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 154883	Seq: 1	Gender: M
Reported: 17/07/2000		Weight:
Hospitalisation:		Age: 25Y
Onset Date: 22/01/2000		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypotension Cardiac arrest Respiratory depression	Life threatening	Bp dropped to 60/40.	Resuscitated. iv fluids, 5l normal saline, 500ml haemacel, 1.5 l albumin.

Medicine Details:

HYPNOVEL (Suspected)		Reason:	
Injection	7.5 Milligram	Daily	Intravenous
Batch:	Started: 22/01/2000	Stopped:	22/01/2000
DROPERIDOL (Suspected)		Reason:	
Injection	10.0 Milligram	Daily	Intravenous
Batch:	Started: 22/01/2000	Stopped:	22/01/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 161274

Seq: 1

Gender: U

Reported: 07/02/2001

Weight: 22.00

Hospitalisation:

Age: AC

Onset Date: 19/01/2001

DOB:

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Consciousness fluctuating		Less talkative, staring into space.	

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	3.0 Milligram	Total	Intravenous
Batch:	Started: 18/01/2001	Stopped:	19/01/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking fentanyl, paracetamol, maxolon, ondansetron.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 161794

Seq: 1

Gender: F

Reported: 21/02/2001

Weight:

Hospitalisation:

Age: 42Y

Onset Date: 18/12/2000

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Malaise Hyperkinesia		General feeling of unwell. Involuntary muscle restlessness.	Given iv cogentin 4mg and valium iv 2mg.

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Oral Liquid	30.0 Millilitre Total Oral
Batch:	Started: 18/12/2000 Stopped: 18/12/2000

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 165036 **Seq:** 1 **Gender:** F
Reported: 01/06/2001 **Weight:** 43.00
Hospitalisation: **Age:** 10
Onset Date: 25/05/2001 **DOB:** 08/03/1991
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Hed held in extorsion, jaw clenched shut	Cogentin. date of recovery 25/05/01

Medicine Details:

ONDANSETRON HYDROCHLORIDE DIHYDRATE (Other drug)	Reason: Nausea and vomiting
4.0 Milligram	As necessary
Batch:	Started: 25/05/2001 Stopped: CONTIN
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
0.2 Milligram	As necessary
Batch:	Started: 24/05/2001 Stopped: 25/05/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Reaction occurred 4.5 hrs post droperidol.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 169227	Seq: 1	Gender: F
Reported: 17/10/2001		Weight:
Hospitalisation:		Age: 48Y
Onset Date: 04/10/2001		DOB:
Outcome: Recovered		Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Rash erythematous		Face and chest.	
Tachycardia		Increasing tachycardia.	
Hypertension			

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	1.2 Milligram	Daily	Intravenous
Batch:	Started: 04/10/2001	Stopped: 04/10/2001	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 171713
Reported: 04/01/2002

Seq: 1

Gender: F
Weight: 18.00

Hospitalisation:

Onset Date: 23/12/2001
Outcome: Recovered

Age: 6
DOB: 08/11/1995
Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypertonia			Date of recovery 23.12.01. patient already in hospital.
Staring			

Medicine Details:

MIDAZOLAM (Other drug) Oral application Batch:	4.0 Milligram Started: 20/12/2001	Reason: Daily	Oral Stopped:
RANITIDINE (Other drug) Injection Batch:	54.0 Milligram Started: 22/12/2001	Reason: Daily	Intravenous Stopped: CONTIN
PARACETAMOL (Other drug) Oral application Batch:	1.6 Gram Started: 20/12/2001	Reason: Daily	Oral Stopped: CONTIN
DROPERIDOL (Suspected) Injection Batch:	2.0 Milligram Started: 22/12/2001	Reason: Nausea and vomiting As necessary	Intravenous Stopped: 23/12/2001

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient had had 2 previous doses of droperidol before reaction. reaction occurred when giving relatively close to a dose of metoclopramide.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 171713

Seq: 1

Gender: F

Reported: 04/01/2002

Weight: 18.00

Hospitalisation:

Age: 6

Onset Date: 23/12/2001

DOB: 08/11/1995

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

METOCLOPRAMIDE HYDROCHLORIDE (Suspected)

Reason: Nausea and vomiting

Injection

8.0 Milligram

Daily

Intravenous

Batch:

Started: 21/12/2001

Stopped: 23/12/2001

Laboratory Investigations:

Additional Information:

Patient had had 2 previous doses of droperidol before reaction. reaction occurred when giving relatively close to a dose of metoclopramide.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 172415	Seq: 1	Gender: F
Reported: 05/02/2002		Weight: 85.00
Hospitalisation:		Age: 43Y
Onset Date: 03/12/1997		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Flushing Hypotension Tachycardia		Facial flushing.	

Medicine Details:

ESMERON (Suspected)		Reason:	
Injection	50.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	
DROPERIDOL (Suspected)		Reason:	
Injection	0.5 Milligram	Daily	Intramuscular
Batch:	Started:	Stopped:	
THIOPENTONE SODIUM (Suspected)		Reason:	
Injection	350.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	
FENTANYL (Suspected)		Reason: Pain	
Injection	200.0 Milligram	Daily	Intravenous
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Recent ga with all same agents, except droperidol, 3 months ago uneventful. vecuron used instead of esmeron.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 172415

Seq: 1

Gender: F

Reported: 05/02/2002

Weight: 85.00

Hospitalisation:

Age: 43Y

Onset Date: 03/12/1997

DOB:

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

HYPNOVEL (Suspected)	Reason:
Injection	2.5 Milligram Daily Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Additional Information:

Recent ga with all same agents, except droperidol, 3 months ago uneventful. vecuron used instead of esmeron.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 173278	Seq: 1	Gender: F
Reported: 01/03/2002		Weight: 98.00
Hospitalisation: Admitted to hospital		Age: 43Y
Onset Date: 24/01/2002		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypotension Blood creatine phosphokinase incr Anuria Dyspnoea Generalised oedema Peripheral ischaemia		Decreased bp. Elevated ck. Not passing urine.	Patient was hospitalised. medication was reviewed and some withdrawn with polypharmacy being the major factor.

Medicine Details:

DERALIN (Other drug)		Reason: Anxiety neurosis	
Tablet	40.0 Milligram	Daily	Oral
Batch:	Started: 24/01/2002	Stopped:	CONTIN
DERALIN (Suspected)		Reason: Anxiety neurosis	
Tablet	80.0 Milligram	Daily	Oral
Batch:	Started: 17/01/2002	Stopped:	24/01/2002
CIPRAMIL (Suspected)		Reason: Depression	
Tablet	40.0 Milligram	Daily	Oral
Batch:	Started: 17/01/2002	Stopped:	24/01/2002
PAXAM (Suspected)		Reason: Other disturbance of sensation	
Tablet	4.0 Milligram	Daily	Oral
Batch:	Started: 17/01/2002	Stopped:	24/01/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 173278 **Seq:** 1 **Gender:** F
Reported: 01/03/2002 **Weight:** 98.00
Hospitalisation: Admitted to hospital **Age:** 43Y
Onset Date: 24/01/2002 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Medicine Details:

PAXAM (Suspected)		Reason: Other disturbance of sensation
Tablet	2.0 Milligram	Daily Oral
Batch:	Started: 24/01/2002	Stopped: 24/01/2002
NEURONTIN (Suspected)		Reason:
Capsule	1.8 Gram	Daily Oral
Batch:	Started: 21/01/2002	Stopped: 24/01/2002
NEURONTIN (Suspected)		Reason:
Capsule	300.0 Milligram	Daily Oral
Batch:	Started: 24/01/2002	Stopped: 28/01/2002
DROLEPTAN (Suspected)		Reason: Otr spec symp psychopathol nec
Injection	5.0 Milligram	Daily Intravenous
Batch:	Started: 17/01/2002	Stopped: 26/01/2002

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 173278 **Seq:** 1 **Gender:** F
Reported: 01/03/2002 **Weight:** 98.00
Hospitalisation: Admitted to hospital **Age:** 43Y
Onset Date: 24/01/2002 **DOB:**
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Medicine Details:

ZYPREXA (Suspected)	Reason: Unspecified schizophrenia
Oral application 20.0 Milligram Daily Oral	
Batch:	Started: 15/01/2002 Stopped: 24/01/2002
SEROQUEL (Suspected)	Reason: Unspecified schizophrenia
Tablet 400.0 Milligram Daily Oral	
Batch:	Started: 21/01/2002 Stopped: 24/01/2002
SERENACE (Suspected)	Reason: Unspecified psychosis
Tablet 10.0 Milligram Daily Oral	
Batch:	Started: 21/01/2002 Stopped: 24/01/2002

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 173703	Seq: 1	Gender: M
Reported: 19/03/2002		Weight:
Hospitalisation:		Age: 30
Onset Date: 06/01/2002		DOB: 22/01/1971
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Pyrexia	Required Specialist Consultation	Mild fever.	
Blood creatine phosphokinase incr			
Confusional state			
Dystonia			Treated with fluids. seroquel and droperidol ceased.
Hypertonia			

Medicine Details:

SEROQUEL (Suspected)	Reason:
400.0 Milligram	Daily
Batch:	Started: 15/12/2001 Stopped: 06/01/2002
DROPERIDOL (Suspected)	Reason: Otr spec symp psychopathol nec
40.0 Milligram	Daily
Batch:	Started: 05/01/2002 Stopped: 06/01/2002

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 175129 **Seq:** 1 **Gender:** M
Reported: 06/05/2002 **Weight:** 0.00
Hospitalisation: **Age:** 25
Onset Date: 22/03/2002 **DOB:** 18/01/1977
Outcome: Death 22/03/2002 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Arrhythmia	Life threatening		
Sudden death			

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	As necessary Intramuscular
Batch:	Started: Stopped:
CLOZARIL (Suspected)	Reason:
Tablet	450.0 Milligram Daily Oral
Batch:	Started: 03/08/1999 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
06/05/2002	Autopsy report				

Additional Information:

Patient died approximately 2 and half years after commencing Clozaril. He had been receiveing droperidol for some time and had received an IM injection half an hour prior to his death.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 175870

Seq: 1

Gender: F

Reported: 03/06/2002

Weight: 71.00

Hospitalisation:

Age: 39

Onset Date: 08/05/2002

DOB: 15/01/1963

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Bradycardia			
Cardiac arrest			

Medicine Details:

MORPHINE SULPHATE (Suspected)	Reason: Other disturbance of sensation		
Injection	0.0	Intravenous	
Batch:	Started:	Stopped:	
MIDAZOLAM (Suspected)	Reason: Premedication		
Injection	5.0 Milligram	1 time	Intramuscular
Batch:	Started: 08/05/2002	Stopped:	
PROPOFOL (Suspected)	Reason: Other disturbance of sensation		
Injection	1.0 Dose Unspecified	1 time	Intravenous
Batch:	Started: 08/05/2002	Stopped:	
VECURONIUM BROMIDE (Suspected)	Reason: Other disturbance of sensation		
Injection	1.0 Dose Unspecified	1 time	Intravenous
Batch:	Started: 08/05/2002	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

No reaction to previous anaesthetics.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 175870

Seq: 1

Gender: F

Reported: 03/06/2002

Weight: 71.00

Hospitalisation:

Age: 39

Onset Date: 08/05/2002

DOB: 15/01/1963

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation
Injection	1.0 Dose Unspecified 1 time Intramuscular
Batch:	Started: Stopped:
MAXOLON (Suspected)	Reason: Nausea and vomiting
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: Stopped:
DUCENE (Suspected)	Reason:
Tablet	15.0 Milligram Daily Oral
Batch:	Started: 08/05/2002 Stopped:

Laboratory Investigations:

Additional Information:

No reaction to previous anaesthetics.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 182338 **Seq:** 1 **Gender:** F
Reported: 04/02/2003 **Weight:** 0.00
Hospitalisation: **Age:** 30
Onset Date: 04/10/2002 **DOB:** 25/01/1972
Outcome: Recovered **Causality:** Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia Anxiety Tremor		Patient became shakey, anxious and developed dystonia	Droperidol ceased.

Medicine Details:

PARACETAMOL (Other drug)	Reason:		
4.0 Gram	Daily		
Batch:	Started:	Stopped:	contin
MS CONTIN (Other drug)	Reason:		
Tablet	20.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	Contin
DROPERIDOL (Suspected)	Reason:	Nausea and vomiting	
	250.0 Microgram	1 time	
Batch:	Started:	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Morphine.
Previous reactions:-24/8/02-dystonic reaction to metoclopramide, requiring benztropine iv prn.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 186485 **Seq:** 1 **Gender:** M
Reported: 02/06/2003 **Weight:** 0.00
Hospitalisation: **Age:** 27Y
Onset Date: 20/03/2003 **DOB:**
Outcome: Not yet recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Neutropenia	Caused or prolonged inpatient hospitalisation	Neutropenia	Ceased Seroquel and Chlorpromazine, no further Droperidol given.

Medicine Details:

SEROQUEL (Suspected) Tablet Batch:	Reason: Oral
DROPERIDOL (Suspected) Injection Batch:	Reason: Otr spec symp psychopathol nec 10.0 Milligram 1 time Intramuscular Started: Stopped:
CHLORPROMAZINE HYDROCHLORIDE (Suspected) Tablet Batch:	Reason: Otr spec symp psychopathol nec 100.0 Milligram Daily Oral Started: 18/03/2003 Stopped: 21/03/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details
02/06/2003	Creatinine				
02/06/2003	Neutrophils	(2-7.5)	03/04/2003	1.24	
02/06/2003	Neutrophils	(2-7.5)	28/03/2003	1.09	
02/06/2003	Neutrophils	(2-7.5)	20/03/2003	1.61	
02/06/2003	White blood cells		20/03/2003	3.5	
02/06/2003	White blood cells		28/03/2003	2.8	
02/06/2003	White blood cells		03/04/2003	2.7	

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 190616 **Seq:** 1 **Gender:** F
Reported: 18/09/2003 **Weight:** 0.00
Hospitalisation: **Age:** 31Y
Onset Date: 12/09/2003 **DOB:**
Outcome: Recovered 12/09/2003 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia		Dystonic reaction	Cogentin 1 mg IV

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Batch:	1.0 Dose Unspecified Daily
Started:	Stopped:
STEMETIL (Suspected)	Reason:
Batch:	1.0 Dose Unspecified Daily
Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Other medications taken: Zofran, Na Heparin.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 193274	Seq: 1	Gender: F
Reported: 17/12/2003		Weight: 0.00
Hospitalisation:		Age: 17
Onset Date: 05/12/2003		DOB: 15/01/1986
Outcome: Recovered	05/12/2003	Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Depressed level of consciousness		Consciousness impaired.	Required ICU admission, given Flumazenil 500mg iv with rapid response.

Medicine Details:

SEROQUEL (Interaction)	Reason: Otr spec symp psychopathol nec
Batch:	Started:
Batch:	Stopped:
MIDAZOLAM (Interaction)	Reason: Otr spec symp psychopathol nec
Injection	10.0 Milligram
	As necessary Intramuscular
Batch:	Started:
Batch:	Stopped:
DROPERIDOL (Interaction)	Reason: Otr spec symp psychopathol nec
Injection	5.0 Milligram
	As necessary Intramuscular
Batch:	Started:
Batch:	Stopped:
Abilify (Interaction)	Reason: Otr spec symp psychopathol nec
Tablet	15.0 Milligram
	Daily Oral
Batch:	Started: 26/11/2003
Batch:	Stopped: 05/12/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking Levlen, Ferro-Gradmet, Nicabate, Mylanta.
Aripiprazole commenced 1 week prior (26/11/03).
Multiple drug interactions many have pre this event.cipitated



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 193274

Seq: 1

Gender: F

Reported: 17/12/2003

Weight: 0.00

Hospitalisation:

Age: 17

Onset Date: 05/12/2003

DOB: 15/01/1986

Outcome: Recovered

05/12/2003

Causality: Causality possible

Reaction Details:

Medicine Details:

LAMICTAL (Interaction)		Reason:	
Tablet	137.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	
DIAZEPAM (Interaction)		Reason: Otr spec symp psychopathol nec	
Tablet	20.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	
EPILIM (Interaction)		Reason:	
Tablet	1000.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	

Laboratory Investigations:

Additional Information:

Patient was also taking Levlen, Ferro-Gradmet, Nicabate, Mylanta.
Aripiprazole commenced 1 week prior (26/11/03).
Multiple drug interactions many have pre this event.cipitated



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 193653 **Seq:** 1 **Gender:** F
Reported: 06/01/2004 **Weight:** 0.00
Hospitalisation: **Age:** 9
Onset Date: 13/11/2003 **DOB:** 30/03/1994
Outcome: Recovered 13/11/2003 **Causality:** Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration		Oculogyric crisis.	

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Batch:	Dose Unspecified
Started:	Stopped: 13/11/2003

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 196398 **Seq:** 1 **Gender:** F
Reported: 13/04/2004 **Weight:** 50.00
Hospitalisation: **Age:** 15
Onset Date: 24/03/2004 **DOB:** 06/01/1989
Outcome: Recovered 26/03/2004 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypotension		Hypotensive (66/26).	Aripiprazole withheld then ceased. IV fluids given but was hard to re-establish BP.
Drug interaction			

Medicine Details:

DROPERIDOL (Interaction)		Reason: Otr spec symp psychopathol nec	
Injection	10.0 Milligram	As necessary	Intramuscular
Batch:	Started:	Stopped:	
Abilify (Interaction)		Reason: Anxiety neurosis	
Tablet	15.0 Milligram	Daily	Oral
Batch:	Started: 04/03/2004	Stopped: 24/03/2004	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking Buspirone, Lamictal, Centrum and Olanzapine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 197256	Seq: 1	Gender: U
Reported: 11/05/2004		Weight: 0.00
Hospitalisation:		Age: 51
Onset Date: 12/02/2004		DOB: 04/10/1952
Outcome: Recovered	15/02/2004	Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Diarrhoea	Caused or prolonged inpatient hospitalisation	Severe diarrhoea.	Augmentin duo forte ceased. Lomotil 1 tds prn

Medicine Details:

DROPERIDOL (Suspected)		Reason: Nausea and vomiting	
Injection	1.2 Milligram	As necessary	Intravenous
Batch:	Started: 09/02/2004	Stopped: 11/02/2004	
AUGMENTIN DUO FORTE (Suspected)		Reason:	
	2.0 Dose Unspecified	Daily	
Batch:	Started: 10/02/2004	Stopped: 11/02/2004	
ZOFRAN (Suspected)		Reason: Nausea and vomiting	
Injection	16.0 Milligram	Daily	
Batch:	Started: 09/02/2004	Stopped: 11/02/2004	
KEFLEX (Suspected)		Reason:	
Capsule	2.0 Gram	Daily	Oral
Batch:	Started: 09/02/2004	Stopped: 11/02/2004	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient also taking panadol, maxolon,



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2009 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 197414
Reported: 17/05/2004

Seq: 1

Gender: F
Weight: 0.00

Hospitalisation:

Onset Date: 13/05/2004
Outcome: Recovered

Age: 15Y
DOB:
Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Extrapyramidal disorder		Extrapyramidal reaction	Benztropine

Medicine Details:

DROPERIDOL (Suspected)	Reason: Dose Unspecified		
Batch:	Started:	Stopped:	
MAXOLON (Suspected)	Reason: Nausea and vomiting		
Injection	40.0 Milligram	Daily	Intravenous
Batch:	Started: 13/05/2004	Stopped: 13/05/2004	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 198190	Seq: 1	Gender: M
Reported: 16/06/2004		Weight: 0.00
Hospitalisation:		Age: 42
Onset Date: 13/03/2004		DOB:
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Priapism		Priapism and agitation.	Nebulised 5mg salbutamol with no effect (twice). Transferred to A & E. Withdraw blood and fluid from penile area to resolve the situation. Injected adrenaline 1:10,000 diluted 1 in 10ml. Urologist injected aramine to subside.
Agitation			

Medicine Details:

DROPERIDOL (Suspected)		Reason:	
Injection	10.0 Milligram	Daily	
Batch:	Started: 13/03/2004	Stopped:	
QUETIAPINE (Suspected)		Reason:	
Tablet	300.0 Milligram	Daily	Oral
Batch:	Started:	Stopped:	
CHLORPROMAZINE HYDROCHLORIDE (Suspected)		Reason: Otr spec symp psychopathol nec	
Tablet	200.0 Milligram	Daily	Oral
Batch:	Started: 13/03/2004	Stopped:	
DIAZEPAM (Suspected)		Reason:	
Tablet	10.0 Milligram	Daily	Oral
Batch:	Started: 13/03/2004	Stopped:	

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Large amounts of alcohol injected previous night (18 schooners). Priapism resolved but was exacerbated by chlorpromazine.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 198674

Seq: 1

Gender: F

Reported: 02/07/2004

Weight: 0.00

Hospitalisation:

Age: 37

Onset Date: 30/05/2004

DOB: 25/05/1967

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Suicidal ideation		Sad and suicidal thoughts, pain, dysphoria.	Counselling and reassurance.
Dysphoria			
Pain			

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	625.0 Microgram 1 time Intravenous
Batch:	Started: 30/05/2004 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Patient was also taking tramadol, Paracetamol, Metronidazole, Omeprazole, Enoxaparin, Ondansetron, Nicotine patch, Promethazine.



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 200055 **Seq:** 1 **Gender:** F
Reported: 17/08/2004 **Weight:** 0.00
Hospitalisation: Admitted to hospital **Age:** 50
Onset Date: 12/08/2004 **DOB:** 26/07/1954
Outcome: Recovered 12/08/2004 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Muscle rigidity	Caused or prolonged inpatient hospitalisation	Patient experienced muscle rigidity and akathisia.	
Akathisia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

MAXOLON (Suspected)	Reason: Nausea and vomiting
30.0 Milligram	Daily
Batch:	Started: 11/08/2004 Stopped: 12/08/2004
DROPERIDOL (Suspected)	Reason: Nausea and vomiting
	Daily
Batch:	Started: 11/08/2004 Stopped: 12/08/2004

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 201682

Seq: 1

Gender: U

Reported: 13/10/2004

Weight: 0.00

Hospitalisation:

Age: 99U

Onset Date: 29/06/2004

DOB:

Outcome: Recovered

30/06/2004

Causality: Causality probable

Reaction Details:

Medicine Details:

ONDANSETRON HYDROCHLORIDE DIHYDRATE (Other drug)		Reason:	
Injection	12.0 Milligram	Daily	Intravenous
Batch:	Started: 22/06/2004	Stopped:	0
DROPERIDOL (Suspected)		Reason:	
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 28/06/2004	Stopped: 29/06/2004	0

Laboratory Investigations:

Additional Information:

Mother states had similar reaction x 1 before to Maxalon (Metoclopramide). Patient appeared to be hallucinating.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 203329	Seq: 1	Gender: M
Reported: 07/12/2004		Weight: 8.00
Hospitalisation: Admitted to hospital		Age: 0
Onset Date: 25/11/2004		DOB: 16/05/2004
Outcome: Recovered	25/11/2004	Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration	Caused or prolonged inpatient hospitalisation	Oculogyric dystonic reaction.	Required IV Benztropine
Dystonia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	1.0 Dose Unspecified 1 time Intravenous
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 203664	Seq: 1	Gender: F
Reported: 20/12/2004		Weight: 0.00
Hospitalisation:		Age: 38
Onset Date: 30/11/2004		DOB:
Outcome: Not yet recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Supraventricular tachycardia		Patient experienced a cardiac event twenty years ago. On the 30/11/04 was administered Ondansetron HCL. At the same time, she also received Droperidol injection, almost straight after she was administered both medications, she experienced supraventricular tachycardia.	

Medicine Details:

ZOFRAN (Suspected)	Reason:
Batch:	Started: 30/11/2004 Stopped:
DROLEPTAN (Suspected)	Reason:
Batch:	Started: Stopped: 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 207252	Seq: 1	Gender: F
Reported: 19/04/2005		Weight: 58.00
Hospitalisation:		Age: 16
Onset Date: 20/02/2005		DOB: 10/03/1988
Outcome: Recovered		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Muscle twitching	Caused or prolonged inpatient hospitalisation	Twitching legs and arms after 1 dose of droperidol, vomiting.	Benztropine 1mg im.
Vomiting	Caused or prolonged inpatient hospitalisation		

Medicine Details:

MORPHINE SULPHATE (Other drug)	Reason:
Batch:	Started: Stopped:
OXYCODONE HYDROCHLORIDE (Other drug)	Reason:
Batch:	Started: Stopped:
TROPISETRON HYDROCHLORIDE (Other drug)	Reason:
Batch:	Started: Stopped:
AMPICILLIN (Other drug)	Reason:
Batch:	Started: Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 207252

Seq: 1

Gender: F

Reported: 19/04/2005

Weight: 58.00

Hospitalisation:

Age: 16

Onset Date: 20/02/2005

DOB: 10/03/1988

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Medicine Details:

GENTAMICIN SULPHATE (Other drug)	Reason:		
Batch:	Started:	Stopped:	
METRONIDAZOLE (Other drug)	Reason:		
Batch:	Started:	Stopped:	
DROPERIDOL (Suspected)	Reason:		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 20/02/2005	Stopped:	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 210461	Seq: 1	Gender: M
Reported: 05/08/2005		Weight: 80.50
Hospitalisation:		
Onset Date: 18/05/2005		Age: 72
Outcome: Recovered	18/05/2005	DOB: 13/11/1932
		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia	Caused or prolonged inpatient hospitalisation	Dystonic reaction, restlessness, anxiety, felt out of control, could not focus.	Treated with Cogentin IV/1mg.
Anxiety	Caused or prolonged inpatient hospitalisation		
Feeling abnormal	Caused or prolonged inpatient hospitalisation		

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation As necessary
Batch:	Started:
Batch:	Stopped:
MORPHINE SULPHATE (Suspected)	Reason: Nausea and vomiting
	30.0 Milligram Daily
Batch:	Started:
	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2009 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 210461 **Seq:** 1 **Gender:** M
Reported: 05/08/2005 **Weight:** 80.50
Hospitalisation: **Age:** 72
Onset Date: 18/05/2005 **DOB:** 13/11/1932
Outcome: Recovered 18/05/2005 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Restlessness	Caused or prolonged inpatient hospitalisation		

Medicine Details:

DROPERIDOL (Suspected)	Reason: Other disturbance of sensation As necessary	
Batch:	Started:	Stopped:
MORPHINE SULPHATE (Suspected)	Reason: Nausea and vomiting	
	30.0 Milligram Daily	
Batch:	Started:	Stopped:

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 210462 **Seq:** 1 **Gender:** M
Reported: 05/08/2005 **Weight:** 72.00
Hospitalisation: **Age:** 13
Onset Date: 18/05/2005 **DOB:** 24/10/1991
Outcome: Recovered 18/05/2005 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Dystonia	Caused or prolonged inpatient hospitalisation	Dystonic reaction to Droperidol/Morphine in normal saline PCA.	Treated with Cogentin 1mg/IV.

Medicine Details:

MORPHINE SULPHATE (Suspected)	Reason: Pain	
1.0 Milligram	Per hour	
Batch:	Started:	Stopped: 0
DROPERIDOL (Suspected)	Reason: Nausea and vomiting	
1.0 Milligram	Per hour	
Batch:	Started:	Stopped: 18/05/2005

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 215059 **Seq:** 1 **Gender:** F
Reported: 18/01/2006 **Weight:** 46.00
Hospitalisation: **Age:** 12
Onset Date: 14/11/2005 **DOB:** 26/09/1993
Outcome: Recovered 14/11/2005 **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Gaze palsy		After the first dose of Droperidol, patient had difficulty closing eyes with an upward gaze.	

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Injection	0.5 Milligram 1 time Intravenous
Batch:	Started: 14/11/2005 Stopped: 0

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 215576 **Seq:** 1 **Gender:** M
Reported: 06/02/2006 **Weight:** 0.00
Hospitalisation: Required a visit to the doctor **Age:** 56
Onset Date: 27/01/2006 **DOB:** 22/12/1949
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Trismus	Caused or prolonged inpatient hospitalisation	The patient developed lock jaw, thick tongue, restlessness, anxiousness and dystonic movements with arching back.	Droperidol ceased. Given oral Valium 5mg.
Anxiety	Caused or prolonged inpatient hospitalisation		
Dystonia	Caused or prolonged inpatient hospitalisation		

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	2.5 Milligram Daily Intravenous
Batch:	Started: 26/01/2006 Stopped: 27/01/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 215576 **Seq:** 1 **Gender:** M
Reported: 06/02/2006 **Weight:** 0.00
Hospitalisation: Required a visit to the doctor **Age:** 56
Onset Date: 27/01/2006 **DOB:** 22/12/1949
Outcome: Recovered **Causality:** Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Restlessness	Caused or prolonged inpatient hospitalisation		
Tongue disorder	Caused or prolonged inpatient hospitalisation		

Medicine Details:

DROPERIDOL (Suspected)	Reason: Nausea and vomiting
Injection	2.5 Milligram Daily Intravenous
Batch:	Started: 26/01/2006 Stopped: 27/01/2006

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 219433

Seq: 1

Gender: M

Reported: 23/06/2006

Weight: 44.00

Hospitalisation:

Age: 10

Onset Date: 05/05/2006

DOB: 01/01/1996

Outcome: Recovered

Causality: Causality probable

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Muscle spasms Eye rolling Opisthotonus Pain		Patient developed spasm of legs, fingers and neck, arching of the back, eye rolled back.	Droperidol ceased and observed.

Medicine Details:

PANADOL (Other drug)	Reason:	
3.5 Gram	Oral	
Batch:	Started: 03/05/2006	Stopped:
TROPISETRON HYDROCHLORIDE (Other drug)	Reason:	
Injection 2.0 Milligram	Intravenous	
Batch:	Started: 02/05/2006	Stopped:
OXYCODONE HYDROCHLORIDE (Other drug)	Reason:	
5.0 Milligram	Oral	
Batch:	Started: 03/05/2006	Stopped:
DROPERIDOL (Suspected)	Reason:	
500.0 Microgram	As necessary	
Batch:	Started: 03/05/2006	Stopped: 05/05/2006

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 219441

Seq: 1

Gender: F

Reported: 23/06/2006

Weight: 50.00

Hospitalisation:

Age: 14

Onset Date: 23/12/2005

DOB: 20/07/1991

Outcome: Recovered

24/12/2006

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Anxiety		Patient became very anxious, developed spasm feeling over entire body, occasional dyskinesia movement of upper limbs.	Treated with 2 x Benztropine 1mg IV- no response 1 x Diazepam 5 mg IV-settled after 30 min.
Dyskinesia			
Muscle spasms			

Medicine Details:

OXYCODONE HYDROCHLORIDE (Other drug)	Reason:	
Batch:	Started:	Stopped:
METRONIDAZOLE (Other drug)	Reason:	
Batch:	Started:	Stopped:
GENTAMICIN SULPHATE (Other drug)	Reason:	
Batch:	Started:	Stopped:
DICLOFENAC SODIUM (Other drug)	Reason:	
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 219441
Reported: 23/06/2006

Seq: 1

Gender: F
Weight: 50.00

Hospitalisation:

Onset Date: 23/12/2005
Outcome: Recovered

24/12/2006

Age: 14
DOB: 20/07/1991
Causality: Causality possible

Reaction Details:

Medicine Details:

PARACETAMOL (Other drug)		Reason:	
Batch:	Started:	Stopped:	
TROPISETRON HYDROCHLORIDE (Other drug)		Reason:	
Batch:	Started:	Stopped:	
PROMETHAZINE HYDROCHLORIDE (Suspected)		Reason:	
Injection	25.0 Milligram	As necessary	Intravenous
Batch:	Started: 23/12/2005	Stopped: 23/12/2005	
DROPERIDOL (Suspected)		Reason:	
Injection	500.0 Microgram	As necessary	Intravenous
Batch:	Started: 20/12/2005	Stopped: 23/12/2005	

Laboratory Investigations:

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 219583	Seq: 1	Gender: F
Reported: 30/06/2006		Weight: 0.00
Hospitalisation:		Age: 48
Onset Date: 08/04/2006		DOB: 02/08/1957
Outcome: Recovered	08/04/2006	Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Hypotension		Profound hypotension (BP <30mmHg with artenal line), rash and tachycardia.	IV fluid, IV increments Aramine, IV Adrenaline total 0.8mg.
Rash			
Tachycardia			

Medicine Details:

PROPOFOL (Suspected)		Reason: Other disturbance of sensation
Injection	200.0 Milligram	1 time Intravenous
Batch:	Started: 08/04/2006	Stopped:
ROCURONIUM BROMIDE (Suspected)		Reason: Other disturbance of sensation
Injection	50.0 Milligram	1 time Intravenous
Batch:	Started: 08/04/2006	Stopped:
Keflin Neutral (Suspected)		Reason: Other disturbance of sensation
Injection	100.0 Milligram	1 time Intravenous
Batch:	Started: 08/04/2006	Stopped:
DROPERIDOL (Suspected)		Reason: Other disturbance of sensation
Injection	1.0 Milligram	1 time Intravenous
Batch:	Started: 08/04/2006	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:

Allergies: Sulphurs, Penicillin, Ceforoxamine, Erythromycin. Previous exposure to Cephalothin with OA in 2005- no problem.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 219583

Seq: 1

Gender: F

Reported: 30/06/2006

Weight: 0.00

Hospitalisation:

Age: 48

Onset Date: 08/04/2006

DOB: 02/08/1957

Outcome: Recovered

08/04/2006

Causality: Causality possible

Reaction Details:

Medicine Details:

DEXAMETHASONE (Suspected)	Reason: Other disturbance of sensation		
Injection	8.0 Milligram	1 time	Intravenous
Batch:	Started: 08/04/2006	Stopped:	
GRANISETRON (Suspected)	Reason: Other disturbance of sensation		
Injection	1.0 Milligram	1 time	Intravenous
Batch:	Started: 08/04/2006	Stopped:	
MORPHINE SULPHATE (Suspected)	Reason: Other disturbance of sensation		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 08/04/2006	Stopped:	
MIDAZOLAM (Suspected)	Reason:		
Injection	5.0 Milligram	1 time	Intravenous
Batch:	Started: 08/04/2006	Stopped:	

Laboratory Investigations:

Additional Information:

Allergies: Sulphurs, Penicillin, Ceforoxamine, Erythromycin. Previous exposure to Cephalothin with OA in 2005- no problem.



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 223236

Seq: 1

Gender: U

Reported: 16/11/2006

Weight: 37.00

Hospitalisation:

Age: 11

Onset Date: 10/09/2006

DOB: 19/05/1995

Outcome: Recovered

Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Oculogyration		Oculogyric crisis.	Patient required benztropine injection.

Medicine Details:

DROPERIDOL (Suspected)	Reason:	
	0.5 Milligram	
Batch:	Started:	Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames:
DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 224140	Seq: 1	Gender: F
Reported: 18/12/2006		Weight: 0.00
Hospitalisation:		Age: 57
Onset Date: 20/10/2006		DOB: 19/01/1949
Outcome: Unknown		Causality: Causality possible

Reaction Details:

Preferred Term	Severity	Report Description	Treatment
Anaphylactic reaction		Developed rash over body, acute hypotension, decreased saturation, bronchospasm.	Adrenaline, dexamethasone, fluids and promethazine.

Medicine Details:

Desflurane (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:
ROCURONIUM BROMIDE (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:
SUXAMETHONIUM BROMIDE (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:
THIOPENTONE SODIUM (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:

Laboratory Investigations:

Date	Type	Range	Date Tested	Result	Details

Additional Information:



THERAPEUTIC GOODS ADMINISTRATION

Public Case Detail

Date Range: 26 Aug 1988 To 31 Dec 2059 Unclear causality excluded GM medicines Only Tradenames: DROLEPTAN,DROPERIDOL

Report Details:

Case Number: 224140

Seq: 1

Gender: F

Reported: 18/12/2006

Weight: 0.00

Hospitalisation:

Age: 57

Onset Date: 20/10/2006

DOB: 19/01/1949

Outcome: Unknown

Causality: Causality possible

Reaction Details:

Medicine Details:

DROPERIDOL (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:
FENTANYL (Suspected)	Reason:
Batch:	Started: 20/10/2006 Stopped:
MORPHINE HYDROCHLORIDE (Suspected)	Reason:
Batch:	1 time Started: 20/10/2006 Stopped:

Laboratory Investigations:

Additional Information: