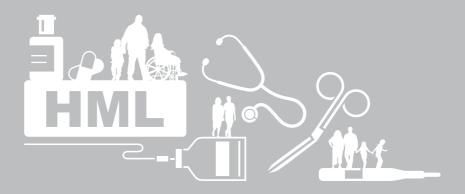
# The Hospital Medicines List (HML) Section H for Hospital Pharmaceuticals

**Update effective 1 May 2014**Cumulative for April and May 2014





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# Summary of decisions EFFECTIVE 1 MAY 2014

- Adrenaline (Hospira) inj 1 in 1,000, 1 ml ampoule and 1 in 10,000, 10 ml ampoule amendment to brand name
- Amoxycillin (Apo-Amoxi) cap 500 mg new listing and addition of HSS
- Amoxycillin (Alphamox) cap 500 mg delisting from 1 July 2014
- Atropine sulphate (Atropt) eye drops 1% addition of HSS
- Bupivacaine hydrochloride (Marcain) inj 2.5 mg per ml, 100 ml bag, and (Marcain Isobaric) inj 5 mg per ml, 4 ml ampoule addition of HSS
- Calcium gluconate (Hospira) inj 10%, 10 ml ampoule amendment to brand name
- Clonidine (Catapres-TTS-1, TTS-2 and TTS-3) patch 2.5 mg, 100 mcg per day;
   5 mg, 200 mcg per day and 7.5 mg, 300 mcg per day price reduction and addition of HSS
- Diazepam (Hospira) inj 5 mg per ml, 2 ml ampoule amendment to brand name
- Ethanol, dehydrated, inj 96% additional presentation
- Etoposide (Hospira) inj 20 mg per ml, 5 ml vial amendment to brand name
- Fluorouracil (Hospira) inj 25 mg per ml, 100 ml vial amendment to brand name
- Gemcitabine (Gemcitabine Actavis 200) inj 200 mg vial, and (Gemcitabine Actavis 1000) inj 1 g vial delist from 1 July 2014
- Gentamicin sulphate (Hospira) inj 10 mg per ml, 1 ml ampoule amendment to brand name
- Glyceryl trinitrate (Hospira) inj 5 mg per ml, 10 ml ampoule amendment to brand name
- Heparin sodium (Hospira) inj 1,000 iu per ml, 1 ml ampoule and inj 5,000 iu per ml, 1 ml ampoule – amendment to brand name
- Hyoscine hydrobromide (Hospira) inj 400 mcg per ml, 1 ml ampoule
   amendment to brand name
- Loperamide hydrochloride (Diamide Relief) cap 2 mg price reduction and addition of HSS
- Naloxone hydrochloride (Hospira) inj 400 mcg per ml, 1 ml ampoule
   amendment to brand name
- Olanzapine (Zyprexa Relprevv) inj 210 mg, 300 mg and 405 mg vial

   amendment to restriction

### Summary of PHARMAC decisions – effective 1 May 2014 (continued)

- Oxytocin (BNM) inj 10 iu per ml, 1 ml ampoule amendment to brand name
- Paliperidone (Invega Sustenna) inj 25 mg, 50 mg, 75 mg, 100 mg and 150 mg syringe new listing
- Papaverine hydrochloride (Hospira) inj 12 mg per ml, 10 ml ampoule
   amendment to brand name
- Paraffin liquid with wool fat (Poly-Visc) eye oint 3% with wool fat 3%
   new listing and addition of HSS
- Promethazine hydrochloride (Hospira) inj 25 mg per ml, 2 ml ampoule
   amendment to brand name
- Risperidone (Risperdal Consta) inj 25 mg, 37.5 mg and 50 mg vial
   amendment to restriction
- Tacrolimus (Tacrolimus Sandoz) cap 0.5 mg, 1 mg and 5 mg new listing and addition of HSS
- Tacrolimus (Prograf) cap 0.5 mg, 1 mg and 5 mg delisting from 1 November 2014
- Tiaprofenic acid (Surgam) tab 300 mg delist from 1 May 2014
- Vinblastine sulphate (Hospira) inj 1 mg per ml, 10 ml vial amendment to brand name

(e)	Price x man. Excl. G	ST)	Brand or Generic
,	\$	Per	Manufacturer

# Section H changes to Part II

Effective 1 May 2014

## **ALIMENTARY TRACT AND METABOLISM**

12	LOPERAMIDE HYDROCHLORIDE (‡ price and addition of HSS) Cap 2 mg - 1% DV Jul-14 to 2016	400	Diamide Relief
BL0	DD AND BLOOD FORMING ORGANS		
30	HEPARIN SODIUM (amendment to brand name) Inj 1,000 iu per ml, 1 ml ampoule	50 5	Hospira <del>Mayne</del> Hospira <del>Mayne</del>
32	CALCIUM GLUCONATE (amendment to brand name) Inj 10%, 10 ml ampoule21.40	10	Hospira <del>Mayne</del>
CAR	DIOVASCULAR SYSTEM		
41	CLONIDINE (4 price and addition of HSS)  Patch 2.5 mg, 100 mcg per day – <b>1% DV Jul-14 to 2017</b> 12.80  Patch 5 mg, 200 mcg per day – <b>1% DV Jul-14 to 2017</b> 18.04  Patch 7.5 mg, 300 mcg per day – <b>1% DV Jul-14 to 2017</b> 22.68	4 4 4	Catapres-TTS-1 Catapres-TTS-2 Catapres-TTS-3
44	GLYCERYL TRINITRATE (amendment to brand name) Inj 5 mg per ml, 10 ml ampoule40.00	5	Hospira <del>Mayne</del>
44	ADRENALINE (amendment to brand name) Inj 1 in 1,000, 1 ml ampoule	5 5	Hospira <del>Mayne</del> Hospira <del>Mayne</del>
46	PAPAVERINE HYDROCHLORIDE (amendment to brand name) Inj 12 mg per ml, 10 ml ampoule73.12	5	Hospira <del>Mayne</del>
GEN	TO-URINARY SYSTEM		
57	OXYTOCIN (amendment to brand name) Inj 10 iu per ml, 1 ml ampoule – 1% DV Feb-14 to 20155.98	5	<del>Oxytocin</del> BNM
INFE	CTIONS – AGENTS FOR SYSTEMIC USE		
65	GENTAMICIN SULPHATE (amendment to brand name) Inj 10 mg per ml, 1 ml ampoule8.56	5	Hospira <del>Mayne</del>
67	AMOXYCILLIN Cap 500 mg – <b>1% DV Jul-14 to 2016</b> 20.94 Note – Alphamox to be delisted from 1 July 2014.	500	Apo-Amoxi

Price (ex man. Excl. G	CT)	Brand or Generic
 \$	Per	Manufacturer

### Changes to Section H Part II - effective 1 May 2014 (continued)

### MUSCULOSKELETAL SYSTEM

96	TIAPROFENIC ACID		
	Tab 300 mg19.26	60	Surgam
	Note – Sugram tab 300 mg to be delisted from 1 May 2014.		

### **NERVOUS SYSTEM**

99	BUPIVACAINE HYDROCHLORIDE (addition of HSS) Inj 5 mg per ml, 4 ml ampoule – <b>1% DV Jul-14 to 2017</b> 50.00 Inj 2.5 mg per ml, 100 ml bag – <b>1% DV Jul-14 to 2017</b> 150.00	5 5	Marcain Isobaric Marcain
107	DIAZEPAM (amendment to brand name) Inj 5 mg per ml, 2 ml ampoule	5	Hospira <del>Mayne</del>
112	HYOSCINE HYDROBROMIDE (amendment to brand name) Inj 400 mcg per ml, 1 ml ampoule	5	Hospira <del>Mayne</del>
116	PALIPERIDONE  → Inj 25 mg syringe 194.25  → Inj 50 mg syringe 271.95  → Inj 75 mg syringe 357.42  → Inj 100 mg syringe 435.12  → Inj 150 mg syringe 435.12	1 1 1 1	Invega Sustenna Invega Sustenna Invega Sustenna Invega Sustenna Invega Sustenna

### Restricted

### Initiation

Re-assessment required after 12 months

Either:

- 1 The patient has had an initial Special Authority approval for risperidone depot injection or olanzapine depot injection; or
- 2 All of the following:
  - 2.1 The patient has schizophrenia or other psychotic disorder; and
- 2.2 The patient has tried but failed to comply with treatment using oral atypical antipsychotic agents; and
- 2.3 The patient has been admitted to hospital or treated in respite care, or intensive outpatient or home-based treatment for 30 days or more in the last 12 months.

### Continuation

Re-assessment required after 12 months

The initiation of paliperidone depot injection has been associated with fewer days of intensive intervention than was the case during a corresponding period of time prior to the initiation of an atypical antipsychotic depot injection.

	Price		Brand or
(ex	man. Excl. G	ST)	Generic
	\$	Per	Manufacturer

### Changes to Section H Part II - effective 1 May 2014 (continued)

116 OLANZAPINE (amendment to restriction)

→ Inj 210 mg vial	280.00	1	Zyprexa Relprevv
→ Inj 300 mg vial	460.00	1	Zyprexa Relprevv
→ Inj 405 mg vial	560.00	1	Zyprexa Relprevv

### Restricted

### Initiation

Re-assessment required after 6 12 months

### Fither.

- 1 The patient has had an initial Special Authority approval for risperidone depot injection or paliperidone depot injection; or
- 2 All of the following:
  - 2.1 The patient has schizophrenia; and
  - 2.2 The patient has tried but failed to comply with treatment using oral atypical antipsychotic agents; and
  - 2.3 The patient has been admitted to hospital or treated in respite care, or intensive outpatient or home-based treatment for 30 days or more in the last 12 months.

### Continuation

Re-assessment required after 12 months

Either:

- 1 The patient has had less than 12 months' treatment with olanzapine depot injection and there is no clinical reason to discontinue treatment; or
- 2 The initiation of olanzapine depot injection has been associated with fewer days of intensive intervention than was the case during a corresponding period of time prior to the initiation of an atypical antipsychotic olanzapine depot injection.
- 117 RISPERIDONE (1 price and amendment to restriction)

→	· Inj 25 mg vial	135.98	1	Risperdal Consta
<b>→</b>	· Inj 37.5 mg vial	178.71	1	Risperdal Consta
<b>→</b>	Inj 50 mg vial	217.56	1	Risperdal Consta

### Restricted

### Initiation

Re-assessment required after 6 12 months

### Either:

- 1 The patient has had an initial Special Authority approval for paliperidone depot injection or olanzapine depot injection; or
- 2 All of the following:
  - 2.1 The patient has schizophrenia or other psychotic disorder; and
  - 2.2 The patient has tried but failed to comply with treatment using oral atypical antipsychotic agents; and
  - 2.3 The patient has been admitted to hospital or treated in respite care, or intensive outpatient or home-based treatment for 30 days or more in the last 12 months.

### Continuation

Re-assessment required after 12 months

### Fither:

- 1 The patient has had less than 12 months' treatment with risperidone depot injection and there is no clinical reason to discontinue treatment; or
- 2 The initiation of risperidone depot injection has been associated with fewer days of intensive intervention than was the case during a corresponding period of time prior to the initiation of an atypical antipsychotic risperidone depot injection.

Price		Brand or
(ex man. Excl. GS	ST)	Generic
\$	Per	Manufacturer

# Changes to Section H Part II – effective 1 May 2014 (continued)

ONCO	DLOGY AGENTS AND IMMUNOSUPPRESSANTS		
124	FLUOROURACIL (amendment to brand name) Inj 25 mg per ml, 100 ml vial13.55	1	Hospira <del>Mayne</del>
124	GEMCITABINE Inj 200 mg vial12.50	1	Gemcitabine Actavis 200
	Inj 1 g vial62.50	1	Gemcitabine Actavis 1000
	Note – Gemcitabine Actavis 200 and 1000 to be delisted from 1 July 2014.		
126	ETOPOSIDE (amendment to brand name) Inj 20 mg per ml, 5 ml vial	1	Hospira <del>Mayne</del>
131	VINBLASTINE SULPHATE (amendment to brand name) Inj 1 mg per ml, 10 ml vial	5	Hospira <del>Mayne</del>
133	TACROLIMUS  → Cap 0.5 mg − 1% DV Nov-14 to 31/10/18	100 100 50 er 2014.	Tacrolimus Sandoz Tacrolimus Sandoz Tacrolimus Sandoz
RESP	IRATORY SYSTEM AND ALLERGIES		
160	PROMETHAZINE HYDROCHLORIDE (amendment to brand name) Inj 25 mg per ml, 2 ml ampoule	5	Hospira <del>Mayne</del>
SENS	ORY ORGANS		
169	ATROPINE SULPHATE (addition of HSS) Eye drops 1% – <b>1% DV Jul-14 to 2017</b> 17.36	15 ml	Atropt
170	PARAFFIN LIQUID WITH WOOL FAT Eye oint 3% with wool fat 3% – 1% DV Jul-14 to 2017	3.5 g	Poly-Visc
VARI	DUS		
171	NALOXONE HYDROCHLORIDE (amendment to brand name) Inj 400 mcg per ml, 1 ml ampoule	5	Hospira <del>Mayne</del>
171	ETHANOL, DEHYDRATED (additional presentation) Inj 96%		

Price		Brand or
(ex man. Excl. G	ST)	Generic
\$	Per	Manufacturer

# Changes to Section H Part II - effective 1 April 2014

### **BLOOD AND BLOOD FORMING ORGANS**

30 TRISODIUM CITRATE Inj 46.7%, 3 ml syringe

### **CARDIOVASCULAR SYSTEM**

40	DILTIAZEM HYDROCHLORIDE (HSS suspended and new listing) Cap long-acting 180 mg		
	- 5% DV Feb-13 to <b>31/03/14</b> <del>2015</del>	500 30	Apo-Diltiazem CD Cardizem CD
	Cap long-acting 240 mg - 5% DV Feb-13 to <b>31/03/14</b> <del>2015</del>	500 30	Apo-Diltiazem CD Cardizem CD
INFE	CTIONS – AGENTS FOR SYSTEMIC USE		
66	CEFEPIME (HSS suspended)  → Inj 1 g vial – 1% DV Oct-12 to <del>2015</del> <b>31/03/14</b>	1 1	DBL Cefepime DBL Cefepime
MUS	CULOSKELETAL SYSTEM		
94	SUXAMETHONIUM CHLORIDE (‡ price and addition of HSS) Inj 50 mg per ml, 2 ml ampoule – 1% DV Jun-14 to 201778.00	50	AstraZeneca
NER	OUS SYSTEM		
98	PERGOLIDE (delisting)  Tab 0.25 mg – <b>1% DV Sep-11 to 2014</b>	100 100	Permax Permax
111	BETAHISTINE DIHYDROCHLORIDE (‡ price and addition of HSS) Tab 16 mg – <b>1% DV Jun-14 to 2017</b> 4.95	84	Vergo 16
112	PROCHLORPERAZINE (4 price and addition of HSS) Tab 5 mg – <b>1% DV Jun-14 to 2017</b> 9.75	500	Antinaus

		Price (ex man. Excl. GST) \$ P	'er	Brand or Generic Manufacturer		
Char	nges to Section H Part II – effective 1 Ap	oril 2014 (continued)				
121	NICOTINE (amendment to HSS)  Gum 2 mg – 5% DV Oct-11 to 31/03/14 2014  1% DV Apr-14 to 2017	36.47	384	Habitrol (Classic) Habitrol (Fruit) Habitrol (Mint)		
	Gum 4 mg – 5% DV Oct-11 to <b>31/03/14</b> <del>2014</del> <b>1% DV Apr-14 to 2017</b>	42.04	384	Habitrol (Classic) Habitrol (Fruit) Habitrol (Mint)		
	Patch 7 mg per 24 hours – 5% DV Oct-11 to 31/03/14 2014 1% DV Apr-14 to 2017 Patch 14 mg per 24 hours – 5% DV Oct-11 to	18.13	28	Habitrol		
	31/03/14 2014 1% DV Apr-14 to 2017	18.81	28	Habitrol		
	Patch 21 mg per 24 hours – 5% DV Oct-11 to 31/03/14 2014 1% DV Apr-14 to 2017 Lozenge 1 mg – 5% DV Oct-11 to 31/03/14 20:		28	Habitrol		
	1% DV Apr-14 to 2017 Lozenge 2 mg – 5% DV Oct-11 to 31/03/14 <del>201</del>	19.94	216	Habitrol		
	1% DV Apr-14 to 2017		216	Habitrol		
ONC	ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS					
123	METHOTREXATE  Tab 2.5 mg – 1% DV Jun-14 to 2015  Tab 10 mg – 1% DV Jun-14 to 2015  Note – Methoblastin tab 2.5 mg and 10 mg to be of	26.25	30 50 4.	Trexate Trexate		
129	IMATINIB MESILATE  Cap 100 mg – <b>1% DV Jul-14 to 2017</b>	298.90	60	Imatinib-AFT		
	Note: Imatinib-AFT is not a registered for the treatn brand of imatinib mesilate (supplied by Novartis) ru with unresectable and/or metastatic malignant GIS	nent of Gastro Intestinal emains fully subsidised (	under Specia	I Authority for patients		
152	AZATHIOPRINE Tab 50 mg – <b>1% DV Jun-14 to 2016</b> Note – Imuprine tab 50 mg to be delisted from 1 Ji		100	Azamun		

Price		Brand or
(ex man. Excl. G	GST)	Generic
\$	Per	Manufacturer

### Changes to Section H Part II – effective 1 April 2014 (continued)

149 RITUXIMAB (amendment to restriction)

<b>→</b>	Inj 10 mg per ml,	10 ml vial	1,075.50	2	Mabthera
<b>→</b>	Inj 10 mg per ml,	50 ml vial	2,688.30	1	Mabthera

Initiation - ANCA associated vasculitis

### Rheumatologist or nephrologist

Limited to 4 weeks' treatment

All of the following:

- 1 Patient has been diagnosed with ANCA associated vasculitis\*; and
- 2 Either:
  - 2.1 Patient does not have MPO-ANCA positive vasculitis\*; or
  - 2.2 Mycophenolate mofetil has not been effective in those patients who have MPO-ANCA positive vasculitis\*; and
- 3 The total rituximab dose would not exceed the equivalent of 375 mg/m² of body-surface area per week for a total of 4 weeks; and
- 4 Any of the following:
  - 4.1 Induction therapy with daily oral or pulse intravenous cyclophosphamide has failed to achieve complete absence of disease after at least 3 months; or
  - 4.2 Patient has previously had a cumulative dose of cyclophosphamide >15 g or a further repeat 3 month induction course of cyclophosphamide would result in a cumulative dose >15 g; or
  - 4.3 Cyclophosphamide and methotrexate are contraindicated; or
  - 4.4 Patient is a female of child-bearing potential; or
  - 4.5 Patient has a previous history of haemorrhagic cystitis, urological malignancy or haematological malignancy.

Note: Indications marked with \* are Unapproved Indications.

Continuation - ANCA associated vasculitis

Rheumatologist or nephrologist

ODAL FEED 1 E I/CAL/ML (delicting)

Limited to 4 weeks' treatment

All of the following:

- 1 Patient has been diagnosed with ANCA associated vasculitis\*; and
- 2 Patient has previously responded to treatment with rituximab but is now experiencing an acute flare of vasculitis; and
- 3 The total rituximab dose would not exceed the equivalent

### **SPECIAL FOODS**

105

193	→ Liquid 5.5 g protein, 21.1 g carbohydrate and 4.81 g fat per 100 ml, can	237 ml	Ensure Plus (Strawberry)
	Note – Ensure Plus (Strawberry) to be delisted from 1 June 2014.		, , ,
195	ORAL FEED  → Powder 16 g protein, 59.8 g carbohydrate and 14 g fat per 100 g, can	850 g	Ensure (Chocolate)
	Note – Ensure (Chocolate) in the 900 g pack size to be delisted from 1 Jul	ne 2014.	

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New Zealand Permit No. 478



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