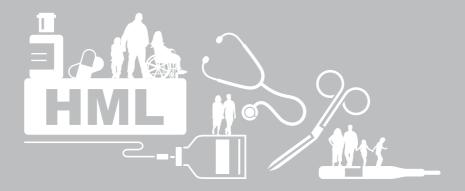
# Section H for Hospital Medicines List (HML) for Hospital Pharmaceuticals

# **Update Effective 1 February 2014**

Cumulative for November, December 2013, January and February 2014





# **Contents**

Summary of decisions effective 1 February 2014	3
Section H changes to Part II	4
Section H changes to Part III	18
Index	19

# Summary of decisions EFFECTIVE 1 FEBRUARY 2014

- Atomoxetine (Strattera) cap 10 mg, 18 mg, 25 mg, 40 mg, 60 mg, 80 mg and 100 mg – amendment to restriction
- Dexamethasone phosphate (Dexamethasone-hameln) inj 4 mg per ml, 1 ml ampoule and 2 ml ampoule – new listing
- Dexamethasone phosphate (Hospira) inj 4 mg per ml, 1 ml ampoule and 2 ml ampoule – delisting from 1 April 2014
- Dimethicone (healthE Dimethicone 5%) crm 5%, 100 g and crm 5% pump bottle, 500 ml new listing
- Dimethyl sulfoxide, soln 99% new listing
- Ferrous sulphate (Ferrograd) tab long-acting 325 mg (105 elemental)
   new listing
- Ferrous sulphate (Ferodan) oral liq 30 mg (6 mg elemental) per ml
   price decrease and addition of HSS
- Fluoxetine hydrochloride (Arrow-Fluoxetine) tab dispersible 20 mg, scored and cap 20 mg – new listing
- Fluoxetine hydrochloride (Fluox) tab dispersible 20 mg, scored and cap 20 mg
   delisting from 1 April 2014
- Isosorbide mononitrate (Ismo 40 Retard) tab long-acting 40 mg new listing
- Isosorbide mononitrate (Corangin) tab long-acting 40 mg HSS suspended 31 January 2014 and delisting from 1 August 2014
- Mesalazine (Pentasa) suppos 1 g delisting 28 packsize from 1 April 2014, the 30 packsize will remain listed
- Olanzapine (Olanzine) tab 2.5 mg delisting from 1 April 2014, the Zypine brand remains listed.
- Phenoxymethlypenicillin (Penicillin V) (AFT) grans for oral liq 125 mg per 5 ml and 250 mg per 5 ml – amendment to presentation description, price decrease and addition of HSS
- Spironolactone (Spiractin) tab 25 mg addition of HSS
- Spironolactone (Spirotone) tab 25 mg HSS suspended 31 January 2014 and delisting from 1 April 2014
- Sugammadex (Bridion) inj 100 mg per ml, 2 ml vial and inj 100 mg per ml, 5 ml vial amendment to restriction
- A range of Optional Pharmaceuticals, including some wound care products and disposable laparoscopic equipment are listed in an addendum to Part III which is available at www.pharmac.govt.nz.

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

**Section H changes to Part II** 

Effective 1 February 2014

# **ALIMENTARY TRACT AND METABOLISM**

12	MESALAZINE (delisting) Suppos 1 g	28 80 packsize w	Pentasa vill remain listed.
20	FERROUS SULPHATE  Tab long-acting 325 mg (105 mg elemental) (new listing)2.06  Oral liq 30 mg (6 mg elemental) per ml  - 1% DV Apr-14 to 2016 (addition of HSS and ↓ price)10.28	30 500 ml	Ferrograd <b>Ferodan</b>
CARI	DIOVASCULAR SYSTEM		
39	SPIRONOLACTONE         Tab 25 mg – 1% DV Feb-14 to 2016 (addition of HSS).       3.65         Tab 25 mg – 1% DV Sep-13 to 31/01/14 2016.       3.65         Note – Spirotone tab 25 mg to be delisted 1 April 2014.	100 100	<b>Spiractin</b> Spirotone
41	ISOSORBIDE MONONITRATE Tab long-acting 40 mg – 1% DV Jun-11 to 31/01/14 20147.50	30	Corangin Ismo 40 Retard
	Note – Corangin tab long-acting 40 mg to be delisted from 1 August 2014.		
DERI	MATOLOGICALS		
46	DIMETHICONE Crm 5% tube – <b>1% DV Apr-14 to 2016</b>	100 g	healthE Dimethicone 5%
	Crm 5% pump bottle – <b>1% DV Apr-14 to 2016</b> 4.73	500 ml	healthE Dimethicone 5%
HORI	MONE PREPARATIONS - SYSTEMIC EXCLUDING CONTRACEP	TIVE HORI	MONES
55	DEXAMETHASONE PHOSPHATE Inj 4 mg per ml, 1 ml ampoule – 1% DV Apr-14 to 201625.80	10	Dexamethasone-
	Inj 4 mg per ml, 2 ml ampoule – 1% DV Apr-14 to 2016 17.98	5	Dexamethasone-
	Note – Hospira inj 4 mg per ml, 1 ml ampoule and 2 ml vial to be delisted 1	April 2014.	hameln
INFE	CTIONS		
63	PHENOXYMETHYLPENICILLIN [PENICILLIN V]  (amendment to presentation description, ↓ price and addition of HSS)  Grans for oral liq 125 mg per ml 5 ml 25 mg per ml	100!	AFT
	<ul> <li>1% DV Apr-14 to 2016</li></ul>	100 ml	AFT

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

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

### MUSCULOSKELETAL SYSTEM

89	SUGAMMADEX (amendment to restriction)		
	→ Inj 100 mg per ml, 2 ml vial1,200.00	10	Bridior
	→ Inj 100 mg per ml, 5 ml vial	10	Bridior

### Restricted

Any of the following:

- 1 Patient requires reversal of profound neuromuscular blockade following a rapid sequence induction that has been undertaken using rocuronium (i.e. suxamethonium is contraindicated or undesirable); or
- 2 Severe neuromuscular degenerative disease where the use of neuromuscular blockade is required; or
- 3 Patient has an unexpectedly difficult airway that cannot be intubated and requires a rapid reversal of anaesthesia and neuromuscular blockade; or
- 4 The duration of the patient's surgery is unexpectedly short; or
- 5 Neostigmine or a neostigmine/anticholinergic combination is contraindicated (for example the patient has ischaemic heart disease, morbid obesity or COPD); or
- 6 Patient has a partial residual block after conventional reversal.

### **NERVOUS SYSTEM**

109

102	FILIOXETIME	HYDROCHLORIDE

Tab dispersible 20 mg, scored – <b>1% DV Apr-14 to 2016</b> 2.50 30	Arrow-Fluoxetine
Cap 20 mg – <b>1% DV Apr-14 to 2016</b> 1.74 90	Arrow-Fluoxetine
Note – Fluox tab dispersible 20 mg, scored and cap 20 mg to be delisted 1 April 2014.	

# OLANZAPINE (delisting)

Tab 2.5 mg	2.00	28	Olanzine
Note – Olanzine tab 2.5 mg to be delisted 1 April 2014. The Zy	pine brand re	mains listed.	

### 113 ATOMOXETINE (amendment to restriction)

→ Cap 10 mg	107.03	28	Strattera
→ Cap 18 mg	107.03	28	Strattera
→ Cap 25 mg	107.03	28	Strattera
→ Cap 40 mg	107.03	28	Strattera
→ Cap 60 mg	107.03	28	Strattera
→ Cap 80 mg	139.11	28	Strattera
→ Cap 100 mg	139.11	28	Strattera

### Restricted

All of the following:

- 1 Patient has ADHD (Attention Deficit and Hyperactivity Disorder) diagnosed according to DSM-IV or ICD 10 criteria: and
- 2 Once-daily dosing; and
- 3 Any of the following:
  - 3.1 Treatment with a subsidised formulation of a stimulant has resulted in the development or worsening of serious adverse reactions or where the combination of subsidised stimulant treatment with another agent would pose an unacceptable medical risk; or
  - 3.2 Treatment with a subsidised formulation of a stimulant has resulted in worsening of co-morbid substance abuse or there is a significant risk of diversion with subsidised stimulant therapy; or
  - 3.3 An effective dose of a subsidised formulation of a stimulant has been trialled and has been discontinued because of inadequate clinical response; or
  - 3.4 Treatment with a subsidised formulation of a stimulant is considered inappropriate because the

continued...

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

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

continued...

### patient has a history of psychoses or has a first-degree relative with schizophrenia; and

4 The patient will not be receiving treatment with atomoxetine in combination with a subsidised formulation of a stimulant, except for the purposes of transitioning from subsidised stimulant therapy to atomoxetine.

Note: A "subsidised formulation of a stimulant" refers to currently listed methylphenidate hydrochloride tablet formulations (immediate release, sustained-release and extended-release) or dexamphetamine sulphate tablets.

### **VARIOUS**

168 DIMETHYL SULFOXIDE Soln 99%

### Effective 1 January 2014

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

25	FI	TRI	AI/IR	UDVC

<b>→</b>	Tab 25 mg1,771.00	28	Revolade
	Tab 50 mg	28	Revolade

### Restricted

Haematologist

### Initiation (idiopathic thrombocytopenic purpura – post-splenectomy)

Re-assessment required after 6 weeks.

All of the following:

- 1. Patient has had a splenectomy; and
- Two immunosuppressive therapies have been trialled and failed after therapy of 3 months each (or 1 month for rituximab); and
- 3. Fither:
  - 3.1. Patient has a platelet count of ≤20,000 platelets per microlitre and has evidence of active bleeding; or
  - 3.2. Patient has a platelet count of ≤10,000 platelets per microlitre.

### Initiation - (idiopathic thrombocytopenic purpura - preparation for splenectomy)

Re-assessment required after 6 weeks.

The patient requires eltrombopag treatment as preparation for splenectomy.

### Continuation – (idiopathic thrombocytopenic purpura – post-splenectomy)

Re-assessment required after 12 months

The patient has obtained a response (see note) from treatment during the initial approval or subsequent renewal periods and further treatment is required.

Note: Response to treatment is defined as a platelet count of >30,000 platelets per microlitre.

### 28 ASPIRIN

Tab 100 mg – <b>1% DV Mar-14 to 2016</b>	990	Ethics Aspirin EC
1.60	90	Ethics Aspirin EC

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

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

# **CARDIOVASCULAR SYSTEM**

33	CILAZAPRIL WITH HYDROCHLOROTHIAZIDE Tab 5 mg with hydrochlorothiazide 12.5 mg - 1% DV Mar-14 to 201610.72	100	Apo-Cilazapril/ Hydrochlorothiazide
	Note – Inhibace Plus tab 5 mg with hydrochlorothiazide 12.5 mg to be delic	sted from 1 I	March 2014.
37	DILTIAZEM HYDROCHLORIDE (HSS suspended and new brand listed) Cap long-acting 120 mg		
	- 5% DV Feb-13 to <b>31/12/13</b> <del>2015</del>	500	Apo-Diltiazem CD
	1.91	30	Cardizem CD
43	BOSENTAN (↓ price)  → Tab 62.5 mg	60 60	pms-Bosentan pms-Bosentan

### **DERMATOLOGICALS**

49 POTASSIUM PERMANGANATE Crystals

ETHINVI OFCEDADIOL WITH LEVONODOCCEDEL

### **GENITO-URINARY SYSTEM**

ונ	Tab 20 mcg with levonorgestrel 100 mcg and 7 inert tablets 2.65 Tab 30 mcg with levonorgestrel 150 mcg and 7 inert tablets 2.30	84 84	Ava 20 ED Ava 30 ED
54	SODIUM CITRO-TARTRATE († price) Grans eff 4 g sachets	28	Ural

### INFECTIONS

61	CEFTRIAXONE Inj 500 mg vial – <b>1% DV Mar-14 to 2016</b> Inj 1 g vial – <b>1% DV Mar-14 to 2016</b> Inj 2 g vial – <b>1% DV Mar-14 to 2016</b> Note – Veracol inj 500 mg vial, inj 2 g vial and Aspen Ceftriaxon	5.22 2.75	1 5 1 to be deliste	Ceftriaxone-AFT Ceftriaxone-AFT Ceftriaxone-AFT d from 1 March 2014.
62	AMOXYCILLIN Cap 250 mg – <b>1% DV Mar-14 to 2016</b> Note – Alphamox cap 250 mg to be delisted from 1 March 2014		500	Apo-Amoxi
63	PHENOXYMETHYLPENICILLIN [PENICILLIN V] († price) Cap 250 mgCap 500 mg		50 50	Cilicaine VK Cilicaine VK
66	NYSTATIN († price) Tab 500,000 u Cap 500,000 u		50 50	Nilstat Nilstat

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

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

80 ZANAMIVIR

### Restricted

Fither:

- 1. Only for hospitalised patient with known or suspected influenza; or
- 2. For prophylaxis of influenza in hospitalised patients as part of a DHB hospital approved infections control plan.

### **MUSCULOSKELETAL SYSTEM**

90 IBUPROFEN (‡ price and addition of HSS)
Oral lig 20 mg per ml – 1% DV Mar-14 to 2016.......1.89 200 ml Fenpaed

### **NERVOUS SYSTEM**

112 LORAZEPAM († price)

Tab 1 mg	19.82	250	Ativan
Tab 2.5 mg	12 /0	100	Ativan

### **ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS**

121 ERLOTINIB (1 price and amendment of restriction)

<b>→</b>	Tab 100 mg	1,133.00	30	Tarceva
<b>→</b>	Tah 150 ma	1.700.00	30	Tarceva

Restricted

Initiation

Re-assessment required after 3 months

### Both:

- 1 Patient has advanced, unresectable, Non Small Cell Lung Cancer (NSCLC); and
- 2 Patient has documented disease progression following treatment with first line platinum basedehemotherapy:

### Either

- 1 All of the following:
  - 1.1 Patient has locally advanced or metastatic, unresectable, non-squamous Non Small Cell Lung Cancer (NSCLC); and
  - 1.2 There is documentation confirming that the disease expresses activating mutations of EGFR tyrosine kinase; and
  - 1.3 Either
    - 1.3.1 Patient is treatment naïve: or
    - 1.3.2 Both:
      - 1.3.2.1 Patient has documented disease progression following treatment with first line platinum based chemotherapy; and
      - 1.3.2.2 Patient has not received prior treatment with gefitinib; and
  - 1.4 Erlotinib is to be given for a maximum of 3 months, or
- 2 The patient received funded erlotinib prior to 31 December 2013 and radiological assessment (preferably including CT scan) indicates NSCLC has not progressed.

### Continuation

Re-assessment required after 6 months

Radiological assessment (preferably including CT scan) indicates NSCLC has not progressed.

	Price (ex man. Excl. GST)		Brand or Generic
	\$	Per	Manufacturer
Char	nges to Section H Part II – effective 1 January 2014 (contin	iued)	
148	AZATHIOPRINE († price) Inj 50 mg vial	1	Imuran
RESI	PIRATORY SYSTEM AND ALLERGIES		
154	FLUTICASONE WITH SALMETEROL — Restricted (removal of restriction) Aerosol inhaler 50 mcg with salmeterol 25 mcg	120 dose 60 dose 120 dose 60 dose	Seretide Seretide Accuhaler Seretide Seretide Accuhaler
SENS	SORY ORGANS		
159	TIMOLOL (addition of HSS)  Eye drops 0.25%, gel forming – <b>1% DV Mar-14 to 2016</b>	2.5 ml 2.5 ml	Timoptol XE Timoptol XE
VAC	CINES		
189	INFLUENZA VACCINE  → Inj 45 mcg in 0.5 ml syringe	10	Influvac Fluarix

### **Effective 1 December 2013**

### **GENERAL RULES**

### 2 Introducing PHARMAC

PHARMAC, the Pharmaceutical Management Agency, is a Crown entity established pursuant to the New Zealand Public Health and Disability Act 2000 (The Act). The primary objective of PHARMAC is to secure for eligible people in need of pharmaceuticals, the best health outcomes that are reasonably achievable from pharmaceutical treatment and from within the amount of funding provided.

The PHARMAC Board consists of up to six members appointed by the Minister of Health. All decisions relating to PHARMAC's operation are made by or under the authority of the Board. More information on the Board can be found at www.pharmac.govt.nz.

The functions of PHARMAC are to perform set out in section 48 of the Act. PHARMAC is required to perform these functions within the amount of funding provided to it in the Pharmaceutical Budget or to DHBs from their own budgets for the use of pharmaceuticals in their hospitals, as applicable, and in accordance with its annual-plan statement of intent and any directions given by the Minister (Section 103 of the Crown Entities Act).

The Government has agreed that PHARMAC will assume responsibility for the assessment, prioritisation and procurement of medical devices on behalf of DHBs. Medical devices come within the definition of Pharmaceuticals in the Act.

PHARMAC is assuming responsibility for procurement of some medical devices categories immediately, as a first step to full PHARMAC management of these categories within the Pharmaceutical Schedule.

continued...

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

# Changes to Section H Part II – effective 1 December 2013 (continued) continued...

- "Give" means to administer, provide or dispense (or, in the case of a Medical Device, use) a Pharmaceutical, or to arrange for the administration, provision or dispensing (or, in the case of a Medical Device, use) of a Pharmaceutical, and "Given" has a corresponding meaning.
- "National Contract" means a contractual arrangement between PHARMAC and a Pharmaceutical supplier which sets out the basis on which any Pharmaceutical may be purchased for use in a DHB Hospital, including an agreement as to a national price.
- 6 2 Hospital Pharmaceuticals
  - 2.2 Section H Part III lists Optional Pharmaceuticals that PHARMAC and the relevant pharmaceutical supplier Pharmaceutical supplier have entered into contractual arrangements for the purchase of, including an agreement on a national price and other obligations such as HSS. DHB Hospitals may choose whether or not to fund the Optional Pharmaceuticals listed in Part III of Section H, but if they do, they must comply with any National Contract requirements. obligations.
  - 2.3 Section H Part II does not encompass the provision of pharmaceutical treatments for DHB Hospital staff as part of an occupational health and safety programme. DHBs Hospitals may choose whether or not to fund pharmaceutical treatments for such use, but if they do, they must comply with any National Contract requirements. obligations.
- 6 2.1 Section H Part II contains the list of Hospital Pharmaceuticals that must be funded by DHB Hospitals. Section H Part II does not currently encompass the following categories of pharmaceuticals except for any items specifically listed in this Section H Part II:
  - a) Medical Devices:
  - b) whole or fractionated blood products:
  - c) diagnostic products which have an ex vivo use, such as pregnancy tests and reagents;
  - d disinfectants and sterilising products, except those that are to be used in or on a patient:
  - e) foods and probiotics:
  - f) radioactive materials;
  - g) medical gases; and
  - h) parenteral nutrition.
  - Subject to rule 2.2, the funding of pharmaceuticals identified in a) h) above is a decision for individual DHB Hospitals.
- 8 18 Hospital Pharmaceutical Contracts
  - 18.1 A DHB Hospital may enter into a contract for the purchase of any Pharmaceutical, including any Medical Device, that it is entitled to fund in accordance with this Schedule H and that is not a National Contract Pharmaceutical, provided that such a contract:
    - does not oblige the relevant DHB Hospital to purchase a volume of that Pharmaceutical, if that Pharmaceutical is a DV Pharmaceutical, that is greater than the relevant DV Limit;
    - enables PHARMAC to access and use future price and volume data in respect of that Pharmaceutical; and
    - enables the relevant DHB Hospital to terminate the contract or relevant parts of the contract in order to give full effect to the national National contract Contract on no more than 3 months' written notice to the pharmaceutical supplier.
  - 18.3 Following written notification from PHARMAC that a Pharmaceutical is a National Contract Pharmaceutical, either through Section H updates or otherwise, DHB Hospitals must, unless PHARMAC expressly notifies otherwise:
    - a) take any steps available to them to terminate pre-existing contracts or relevant parts of such a contract, and
    - not to enter any new contracts or extend the period of any current contracts, for the supply of that National Contract Pharmaceutical or the relevant chemical entity or Medical Device.

continued...

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

# Changes to Section H Part II – effective 1 December 2013 (continued)

- continued...
  - 19 National Contract Pharmaceuticals
    - 19.1 DHB Hospitals must take all necessary steps to enable any contracts between PHARMAC and a pharmaceutical Pharmaceutical supplier in relation to National Contract Pharmaceuticals to be given full effect
    - 19.2 The contractual arrangement between PHARMAC and the relevant pharmaceutical supplier of a National Contract Pharmaceutical requires it to be made available by for purchase at the relevant Price by any or all of the following:
      - a) DHB Hospitals at Designated Delivery Points; and/or
      - b) Contract Manufacturers (expressly for the purpose of compounding).

In the case of Medical Devices, a National Contract may require the Medical Device to be purchased by, and/or supplied to, a third party logistics provider.

22.1 DHB Hospitals must provide to PHARMAC, on a monthly basis in accordance with PHARMAC's requirements, any volume data and, unless it would result in a breach of a pre-existing contract, price data held by those DHB Hospitals in respect of any Hospital Pharmaceuticals listed in Part II of Section H of the Schedule Pharmaceutical (including any Medical Device) listed in Section H.

### **ALIMENTARY TRACT AND METABOLISM**

12	MESALAZINE		
	Suppos 1 g54.60	30	Pentasa

OILY PHENOL [PHENOL OILY]
Inj 5%, 5 ml vial

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

25 EPTACOG ALFA [RECOMBINANT FACTOR VIIA] (addition of restrictions and amendment to presentation description)

<b>→</b>	Inj 1 mg syringe vial		1	NovoSeven RT
		2,327.50		NovoSeven RT
		5,818.75		NovoSeven RT
		9,310.00		NovoSeven RT

### Restricted

When used in the treatment of haemophilia, treatment is managed by the Haemophilia Treaters Group in conjunction with the National Haemophilia Management Group.

25 MOROCTOCOG ALFA [RECOMBINANT FACTOR VIII] (addition of restrictions)

<b>→</b>	Inj 250 iu vial	225.00	1	Xyntha
<b>→</b>	lnj 500 iu vial	450.00	1	Xyntha
	Inj 1,000 iu vial		1	Xyntha
<b>→</b>	Inj 2,000 iu vial	1,800.00	1	Xyntha
	Inj 3,000 iu vial		1	Xyntha

### Restricted

When used in the treatment of haemophilia, treatment is managed by the Haemophilia Treaters Group in conjunction with the National Haemophilia Management Group.

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

# Changes to Section H Part II – effective 1 December 2013 (continued)

25	NONACOG ALFA	[RECOMBINANT FACTOR IX	1	(addition of restriction)	١

<b>→</b>	Inj 250 iu vial	310.00	1	BeneFIX
	lnj 500 iu vial		1	BeneFIX
	Inj 1,000 iu vial		1	BeneFIX
	Inj 2,000 iu vial		1	BeneFIX

### Restricted

When used in the treatment of haemophilia, treatment is managed by the Haemophilia Treaters Group in conjunction with the National Haemophilia Management Group.

25 FACTOR EIGHT INHIBITORS BYPASSING AGENT (move from Part III and addition of restriction)

<b>→</b>	Inj 500 U	1,640.00	1	FEIBA
<b>→</b>	Ini 1.000 U	3.280.00	1	FEIBA

### Restricted

When used in the treatment of haemophilia, treatment is managed by the Haemophilia Treaters Group in conjunction with the National Haemophilia Management Group.

26 OCTOCOG ALFA [RECOMBINANT FACTOR VIII] (addition of restriction)

UU	I O O O O ALI A [I ILO O IVIDIIVAIVI I AO I O II VII	iij (audition of restriction)		
<b>→</b>	Inj 250 iu vial	237.50	1	Advate
		250.00		Kogenate FS
<b>→</b>	Inj 500 iu vial	475.00	1	Advate
	,	500.00		Kogenate FS
<b>→</b>	Inj 1,000 iu vial	950.00	1	Advate
		1,000.00		Kogenate FS
<b>→</b>	Inj 1,500 iu vial	1,425.00	1	Advate
<b>→</b>	Inj 2,000 iu vial	1,900.00	1	Advate
		2,000.00		Kogenate FS
<b>→</b>	Inj 3,000 iu vial	2,850.00	1	Advate
	•	3.000.00		Kogenate FS

### Restricted

When used in the treatment of haemophilia, treatment is managed by the Haemophilia Treaters Group in conjunction with the National Haemophilia Management Group.

### CARDIOVASCULAR SYSTEM

39 SPIF	RONOL	.ACTONE
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Tab 25 mg	3.65	100	Spiractin
Tab 100 mg	11.80	100	Spiractin

### **DERMATOLOGICALS**

47 CETOMACROGOL WITH GLYCEROL (amendment to presentation description and brand name)

Crm 90% with glycerol 10%, <del>100 g</del> 2.1	100	g Pharmacy Health
2.0	00	Pharmacy Health
3.2	20	healthE
Crm 90% with glycerol 10%, <del>1,000 ml</del>	50 1,000	ml Pharmacy Health Sorbolene with
		Glycerin
Crm 90% with glycerol 10% , <del>500 ml</del>	50 500 r	nl Pharmacy Health
		Sorbolene with
		Glycerin

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

# Changes to Section H Part II - effective 1 December 2013 (continued)

### **GENITO-URINARY SYSTEM**

52 OXYTOCIN

52 LEVONORGESTREL (amendment to restrictions)

→ Intra-uterine system, 20 mcg per day

Restricted

### Obstetrician or gynaecologist

### Initiation – heavy menstrual bleeding

All of the following:

- 1 The patient has a clinical diagnosis of heavy menstrual bleeding; and
- 2 The patient has failed to respond to or is unable to tolerate other appropriate pharmaceutical therapies as per the Heavy Menstrual Bleeding Guidelines; and
- 3 Either Any of the following:
  - 3.1 Serum ferritin level < 16 mcg/l (within the last 12 months); or
  - 3.2 Haemoglobin level < 120 g/l; or
  - 3.3 The patient has had a uterine ultrasound and either a hysteroscopy or endometrial biopsy.

### Continuation - heavy menstrual bleeding

Fither:

- 1 Patient demonstrated clinical improvement of heavy menstrual bleeding; or
- 2 Previous insertion was removed or expelled within 3 months of insertion.

### Initiation - endometriosis

The patient has a clinical diagnosis of endometriosis confirmed by laparoscopy.

Continuation - endometriosis

Either:

- 1 Patient demonstrated satisfactory management of endometriosis: or
- 2 Previous insertion was removed or expelled within 3 months of insertion.

Note – Endrometriosis is an unregistered indication.

### **INFECTIONS**

64 DAPTOMYCIN

→ Inj 500 mg vial

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

### Changes to Section H Part II – effective 1 December 2013 (continued)

### MUSCULOSKELETAL SYSTEM

88	BENZBROMARONE (addition of note)			
	→ Tab 100 mg	45.00	100	Benzbromaron
				AL 100

### Restricted

Both:

- 1 Any of the following:
  - 1.1 The patient has a serum urate level greater than 0.36 mmol/l despite treatment with allopurinol at doses of at least 600 mg/day and appropriate doses of probenecid: or
  - 1.2 The patient has experienced intolerable side effects from allopurinol such that treatment discontinuation is required and serum urate remains greater than 0.36 mmol/l despite appropriate doses of probenecid; or 1.3 Both:
    - 1.3.1 The patient has renal impairment and serum urate remains greater than 0.36 mmol/l despite optimal treatment with allopurinol (see Note); and
    - 1.3.2 The patient has a rate of creatinine clearance greater than or equal to 20 ml/min; or
  - 1.4 All of the following:
    - 1.4.1 The patient is taking azathioprine and requires urate-lowering therapy; and
    - 1.4.2 Allopurinol is contraindicated; and
    - 1.4.3 Appropriate doses of probenecid are ineffective or probenecid cannot be used due to reduced renal function; and
- 2 The patient is receiving monthly liver function tests.

Notes: Benzbromarone has been associated with potentially fatal hepatotoxicity. Optimal treatment with allopurinol in patients with renal impairment is defined as treatment to the creatinine clearance-adjusted dose of allopurinol then, if serum urate remains greater than 0.36 mmol/l, a gradual increase of the dose of allopurinol to 600 mg or the maximum tolerated dose.

The New Zealand Rheumatology Association has developed information for prescribers which can be accessed from its website at http://www.rheumatology.org.nz/benzbromarone prescriber information.cfm

### **NERVOUS SYSTEM**

90	ROPINIROLE HYDROCHLORIDE  Tab 0.25 mg - 1% DV Mar-14 to 2016	5.32 7.72 14.48	100 100 100 100 100	Apo-Ropinirole Apo-Ropinirole Apo-Ropinirole Apo-Ropinirole
93	PRAMIPEXOLE HYDROCHLORIDE Tab 0.25 mg Tab 1 mg		100 100	Ramipex Ramipex
99	MORPHINE SULPHATE (‡ price and addition of HSS) Cap long-acting 10 mg - 1% DV Feb-14 to 2016 Cap long-acting 30 mg - 1% DV Feb-14 to 2016 Cap long-acting 60 mg - 1% DV Feb-14 to 2016 Cap long-acting 100 mg - 1% DV Feb-14 to 2016	2.50 5.40	10 10 10 10	m-Eslon m-Eslon m-Eslon m-Eslon

	(e)	Price ( man. Excl. GST) \$	Per	Brand or Generic Manufacturer
Chan	nges to Section H Part II – effective 1 Decer	mber 2013 (co	ntinued)	
103	GABAPENTIN  → Cap 100 mg  → Cap 300 mg  → Cap 400 mg	11.00	100 100 100	Arrow-Gabapentin Arrow-Gabapentin Arrow-Gabapentin
109	OLANZAPINE Tab 2.5 mg Tab 5 mg Tab orodispersible 5 mg Tab 10 mg Tab orodispersible 10 mg	2.00 3.85 6.36 6.35	28 28 28 28 28	Zypine Zypine Zypine ODT Zypine Zypine ODT
ONC	OLOGY AGENTS AND IMMUNOSUPPRESSANT	rs		
122	IMATINIB <b>MESILATE</b> (amendment to chemical name)  → Tab 100 mg	2,400.00	60	Glivec
RESP	PIRATORY SYSTEM AND ALLERGIES			
153	MONTELUKAST (amendment to restriction)  → Tab 4 mg	wheezing (possible dose of up to 400 least one month; de episodes in the exacerbations at least one month at	ly viral) in chil meg per day and previous 12 reast one of wi ment) in the p	beclomethasone or months of acute wheeze nich required hospitalisation ast 12 months. nhaled corticosteroids and
VARI	IOUS			
163	CHLORHEXIDINE WITH CETRIMIDE (amendment to pro Crm 0.1% 1% with cetrimide 0.5%	esentation descrip	otion)	
163	DESFERRIOXAMINE MESILATE (change to brand name Inj 500 mg vial		10	Hospira <del>Mayne</del>
165	IOHEXOL (new packsize) Inj 350 mg per ml, 200 ml bottle Note – Omnipaque inj 350 mg per ml, 200 ml bottle pa			Omnipaque oruary 2014.

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

### Changes to Section H Part II - effective 1 December 2013 (continued)

### **SPECIAL FOODS**

173 CARBOHYDRATE SUPPLEMENT (delisting)

→ Powder 95 g carbohydrate per 100 g, 368 g can

Note – Moducal is to be delisted from 1 February 2014.

e.a. Moducal

### **Effective 1 November 2013**

### ALIMENTARY TRACT AND METABOLISM

17 LACTULOSE
Oral liq 10 g per 15 ml – 1% DV May-14 to 2016 .......3.84 500 ml Laevolac
Note – Laevolac oral liq 10 g per 15 ml, 1,000 ml pack size will be delisted from 1 May 2014.

ZINC CHLORIDE (amendment to presentation description) Inj 5.3 mg per ml (5.1 mg per ml elemental), 2 ml ampoule

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

28 EPTIFIBATIDE (amendment to restriction)

<b>→</b>	Inj 750 mcg per ml, 100 ml vial	324.00	1	Integrilin
$\rightarrow$	Inj 2 mg per ml, 10 ml vial	111.00	1	Integrilin

### Restricted

Either:

- 1. For use in patients with acute coronary syndromes undergoing percutaneous coronary intervention; or
- 2. For use in patients with definite or strongly suspected intra-coronary thrombus on coronary angiography.

### CARDIOVASCULAR SYSTEM

33	ENALAPRIL MALEATE	

Tab 5 mg1.07	90	m-Enalapril
Tab 10 mg	90	m-Enalapril
Tab 20 mg	90	m-Enalapril

Note – m-Enalapril tab 5 mg, 10 mg and 20 mg will be delisted from 1 January 2014. The Ethics Enalapril brand remains listed.

42 HYDRALAZINE HYDROCHLORIDE (remove S29)

42 MINOXIDIL (correction to listing)

Restricted

For patients with severe refractory hypertension which has failed to respond to extensive multiple therapies.

	Price (ex man. Excl. GST) \$ P	er	Brand or Generic Manufacturer
Char	nges to Section H Part II – effective 1 November 2013 (cont	inued)	
DER	MATOLOGICALS		
49	HYDROCORTISONE WITH MICONAZOLE (correction to listing) Crm 1% with miconazole nitrate 2%	15 g	Micreme H
INFE	CTIONS		
80	OSELTAMIVIR  Powder for oral suspension 6 mg per ml		
	Restricted Either:		
	<ol> <li>Only for hospitalised patient with known or suspected influenza; or</li> <li>For prophylaxis of influenza in hospitalised patients as part of a DHB hosplan.</li> </ol>	spital approv	ved infections control
	Note – Oseltamivir powder for oral suspension 12 mg per ml will be deliste	d from 1 No	vember 2013.
MUS	CULOSKELETAL SYSTEM		
88	BENZBROMARONE (amendment to brand name)  Tab 100 mg45.00	100	Benzbromaron <b>AL</b> 100
NER	VOUS SYSTEM		
93	LEVODOPA WITH BENSERAZIDE (amendment to brand name) Tab dispersible 50 mg with benserazide 12.5 mg10.00	100	Madopar <del>Dispersible</del> <b>Rapid</b>
100	IMIPRAMINE HYDROCHLORIDE (remove S29) Tab 10 mg6.58	60	Tofranil <del>S29</del>
102	PAROXETINE HYDROCHLORIDE Tab 20 mg4.32 Note – Loxamine tab 20 mg (30 packsize) will be delisted from 1 January 2	90 1014	Loxamine
	Total Loran mile and Lee may (see particular), mile de acticular nom nominal y l		
107	ONDANSETRON Tab 4 mg – <b>1% DV Jan-14 to 2016</b>	50 50	Onrex Onrex
	Note – Dr Reddy's Ondansetron tab 4 mg and 8 mg will be delisted from 1		
ONC	OLOGY AGENTS AND IMMUNOSUPPRESSANTS		
119	METHOTREXATE Inj 7.5 mg prefilled syringe - 1% DV Jan-14 to 2016	1 1 1 1 1	Methotrexate Sando Methotrexate Sando Methotrexate Sando Methotrexate Sando Methotrexate Sando Methotrexate Sando

Price		
(ex man. Excl. G	Generic	
\$	Per	Manufacturer

# Changes to Section H Part II – effective 1 November 2013 (continued)

148	AZATHIOPRINE			
	Tab 50 mg	18.45	100	Imuran
	Note – Imuran tab 50 mg will be delisted from 1 November 2013	3. The Imuprir	ne brand i	remains listed.

### **RESPIRATORY SYSTEM AND ALLERGIES**

152	SALBUTAMOL		
	Oral liq 400 mcg per ml - 1% DV Jan-14 to 20162.06	150 ml	Ventolin
	Note – Salapin oral liq 400 mcg per ml to be delisted 1 January 2014.		

### **SPECIAL FOODS**

185

ORAL FEED (change of packsize)			
→ Powder 16 g protein, 59.8 g carbohydrate			
and 14 g fat per 100 g, can	13.00	850 g	Ensure (Vanilla)
Note - Ensure (Vanilla) 900 g packsize to be delisted	1 February 2014.		

# **Section H changes to Part III**

# Effective 1 February 2014

### 192 OPTIONAL PHARMACEUTICALS

In addition to the products expressly listed here in Part III: Optional Pharmaceuticals, a number of additional Optional Pharmaceuticals, including some wound care products and disposable laparoscopic equipment, are listed in an addendum to Part III which is available at www.pharmac.govt.nz. The Optional Pharmaceuticals listed in the addendum are deemed to be listed in Part III, and the Rules of the Pharmaceutical Schedule applying to products listed in Part III apply to them.

# Index

# Pharmaceuticals and brands

Advate 12 Fenpaed 6 Anoxycillin 77 Ferrodan 4 Apo-Amoxi 77 Ferrodan 4 Apo-Amoxi 77 Ferrograd 4 Apo-Dilliazem CD 77 Ferrous sulphate 4 Apo-Dilliazem CD 77 Ferrous sulphate 6 Apo-Dilliazem CD 77 Fluarix 9 Apo-Ropinirole 14 Fluoxetine hydrochloride 5 Aprosoline 16 Fluticasone with salmeterol 6 Arrow-Fluoxetine 75 Gabapentin 15 Gabapentin 15 Gabapentin 15 Galvec 11 Ativan 8 Glycopyrronium bromide 11 Attwan 8 Glycopyronium bromide 11 Attwan 9 Elb 11 B	A		FEIBA	
Apo-Cilazapril/Hydrochlorothiazide 7 Ferrograd. 4 Apo-Cilazapril/Hydrochlorothiazide 7 Ferrogra sulphate 4 Apo-Diltiazem CD. 7 Fluarix. 5 Apo-Ropiliriole 14 Fluoxetine hydrochloride. 5 Arrow-Fluoxetine 5 Arrow-Fluoxetine 5 Arrow-Gabapentin 15 Gabapentin 15 Aspirin 66 Gilvec 11 Attivan 8 Glycopyrronium bromide. 16 Attivan 8 Glycopyrronium bromide. 16 Attivan 8 Glycopyrronium bromide. 16 Attivan 8 Hydrochloride 16 Attivan 8 Hydrochloride 16 Attivan 8 Hydrochloride 16 Attivan 8 Hydrochloride 16 Attivan 8 Hydrocortisone with miconazole 17 Ava 20 ED. 7 Hydralazine hydrochloride 16 BeneFIX 12 Ibuprofen 8 BeneFIX 12 Ibuprofen 8 BeneFIX 14 Imipramine hydrochloride 17 Benezbromaron AL 100 114, 17 Benezbromarone 14, 17 Benezbromarone 14, 17 Benezbromarone 14, 17 Benzbromarone 15 C Influenza vaccine 5 C Influenza vaccine 5 Carbohydrate supplement 16 Cardizem CD. 7 Iohexol 11 Cardixone AFT 7 K Cefoxitin 13 Ismo 40 Retard 4 Cardixanoe AFT 7 K Cetomacrogol with glycerol 12 Collicaine VK 7 L acevolac 16 Cilcaine VK 7 L acevolac 16 Cilcaine VK 7 L acevolac 16 Cilcaine VK 7 T L acevolac 16 Corangin 4 Levonorgestrel 17 Desamethasone phosphate 17 Desamethasone hosphate 17 Desamethasone hosphate 18 E	Advate	12	Fenpaed	8
Apo-Calizarpril/Hydrochlorothiazide 7 Ferrous sulphate 4 Apo-Dilitazem CD 7 Fluarix 5 Ferrous sulphate 4 Apo-Ropinirole 14 Fluoxetine hydrochloride 5 Fusicasone with salmeterol 6 Fluticasone with salmeterol 7 Fluarix 15 Gabapentin 15 Gabapentin 15 Gabapentin 15 Gabapentin 15 Gabapentin 16 Fluticasone with salmeterol 16 Fluticasone with salmeterol 17 Gapapentin 17 Gapapentin 18 Flutiva 18 Glivec 11 Gapapentin 18 Flutiva 18 Glivec 11 Gapapentin 18 Flutiva 18 Glivec 11 Gapapentin 19 Flutiva 19			Ferodan	4
Apo-Bitiazem CD. 7 Fluarix. 9 Supposition CD.			Ferrograd	4
Apo-Repolinicole.         14         Fluoxetine hydrochloride.         5           Apresoline.         16         Fluticasone with salmeterol.         5           Arrow-Eluoxetine.         5         G           Arrow-Gabapentin.         15         Gabapentin.         15           Aspirin.         6         Glivec.         15           Atternow-British and the state of the s	Apo-Cilazapril/Hydrochlorothiazide	7	Ferrous sulphate	4
Apresoline	Apo-Diltiazem CD	7	Fluarix	ć
Arrow-Gabapentin	Apo-Ropinirole	14		
Arrow-Gabapentin 15 Gabapentin 15 Aspirin 6 Gilvec 15 Aspirin 6 Gilvec 15 Aspirin 16 Gilvec 15 Aspirin 16 Gilvec 15 Aspirin 16 Ativan 8 Gilycopyrronium bromide 16 Ava 20 ED. 7 health's Dimethicone 5% 4 Ava 30 ED. 7 Hydralazine hydrochloride 16 Azathioprine 9, 18 Hydrocortisone with miconazole 17 Azathioprine 17 BeneDromaron AL 100 14, 17 Impramine hydrochloride 17 Benezbromarone 14, 17 Impramine hydrochloride 17 Benezbromarone 14, 17 Impramine hydrochloride 17 Benezbromarone 14, 17 Impramine hydrochloride 18 Influenza vaccine 5 Influenza vaccine 6 Influenza vacci	Apresoline	16	Fluticasone with salmeterol	ć
Aspirin 6 Glivec 15 Ativan 8 Glycopyrronium bromide 16 Attomoxetine 5 H Ava 20 ED. 7 health E Dimethicone 5%. 4 Ava 30 ED. 7 Hydralazine hydrochloride 16 Azathioprine 9, 18 Hydrocortisone with miconazole 17 BeneFIX 12 Ibuprofen 8 Benzbromaron AL 100 14, 17 Imatinib mesilate 15 Benzbromarone 14, 17 Imatinib mesilate 15 Bridion 5 Influenza vaccine 5 C Influenza vaccine 15 C Influenza vaccine 16 C Influenza vaccine 1	Arrow-Fluoxetine	5	G	
Ativanoxetine	Arrow-Gabapentin	15	Gabapentin	15
Atomoxetine	Aspirin	6		
Atomoxetine	Ativan	8	Glycopyrronium bromide	16
Ava 30 ED. 7 Hydralazine hydrochloride 16 Azathioprine 9, 18 Hydrocortisone with miconazole 17 B	Atomoxetine	5		
Azathioprine	Ava 20 ED	7	healthE Dimethicone 5%	4
BeneFIX	Ava 30 ED	7		
BeneFIX	Azathioprine 9,	18	Hydrocortisone with miconazole	17
Benzbromaron AL 100	•		ľ	
Benzbromaron AL 100	BeneFIX	12	lbuprofen	8
Bosentan	Benzbromaron AL 100 14,	17		
Bosentan	Benzbromarone 14.	17	Imipramine hydrochloride	17
Bridion	Bosentan	7		
Carbohydrate supplement         16         Integrilin         16           Cardizem CD         7         lohexol         15           Cefoxitin         13         Ismo 40 Retard         4           Ceftriaxone         7         Isosorbide mononitrate         4           Ceftriaxone-AFT         7         K           Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L         12           Chlorhexidine with hydrochlorothiazide         7         Lactulose         16           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levonorgestrel         17           Levonorgestrel         13         Loniten         16           Despermicamente mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Dilitazem hydrochloride         7         Madopar Rapid         17           Dimethyl sulfoxide         6         Mesalazine         4, 11 <td></td> <td></td> <td></td> <td></td>				
Carbohydrate supplement         16         Integrilin         16           Cardizem CD         7         lohexol         15           Cefoxitin         13         Ismo 40 Retard         4           Ceftriaxone         7         Isosorbide mononitrate         4           Ceftriaxone-AFT         7         K           Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L         12           Chlorhexidine with hydrochlorothiazide         7         Lactulose         16           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levonorgestrel         17           Levonorgestrel         13         Loniten         16           Despermicamente mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Dilitazem hydrochloride         7         Madopar Rapid         17           Dimethyl sulfoxide         6         Mesalazine         4, 11 <td>C</td> <td></td> <td>Influvac</td> <td>ç</td>	C		Influvac	ç
Cardizem CD         7         lohexol         15           Cefoxitin         13         Ismo 40 Retard         4           Ceftriaxone         7         Isosorbide mononitrate         4           Ceftriaxone-AFT         7         K           Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L         L           Ciliazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Desamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diitiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         M         15           Eltrombopag         6         Methotrexate         17           Eltrombopag         6         Methotrexate         17           Ensure (Vanill	Carbohydrate supplement	16		
Cefoxitin         13         Ismo 40 Retard         4           Ceftriaxone         7         Isosorbide mononitrate         4           Ceftriaxone-AFT         7         K           Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L         12           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desamethasone-hameln         4         Loxamine         17           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diitiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4,11           E         m-Eslon         14           Eltrombopag         6         Methotrexa			3	
Ceftriaxone				
Cettriaxone-AFT         7         K           Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4,11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H				
Cetomacrogol with glycerol         12         Kogenate FS         12           Chlorhexidine with cetrimide         15         L           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Desamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M         M           Dittiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4,11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recom				
Chlorhexidine with cetrimide         15         L           Cilazapril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptifibatide         16         Moducal         16           Erlotinib         8         Monteluka				12
Cilizazpril with hydrochlorothiazide         7         Lactulose         16           Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslorn         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6	0 0,		=	
Cilicaine VK         7         Laevolac         16           Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4,11           E         m-Eslorn         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorg			_	16
Corangin         4         Levodopa with benserazide         17           D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Eth				
D         Levonorgestrel         13           Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Dittiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4,11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14 <td></td> <td></td> <td></td> <td></td>				
Daptomycin         13         Loniten         16           Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M         M           Dilitiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14	•	'	•	
Desferrioxamine mesilate         15         Lorazepam         8           Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14	_	13	•	
Dexamethasone-hameln         4         Loxamine         17           Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N	• •			
Dexamethasone phosphate         4         M           Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N			•	
Diltiazem hydrochloride         7         Madopar Rapid         17           Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N				. ,
Dimethicone         4         m-Enalapril         16           Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N				17
Dimethyl sulfoxide         6         Mesalazine         4, 11           E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N			m-Fnalanril	16
E         m-Eslon         14           Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N				
Eltrombopag         6         Methotrexate         17           Enalapril maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N	•	U	,	
Enalaprii maleate         16         Methotrexate Sandoz         17           Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N	_	6		
Ensure (Vanilla)         18         Micreme H         17           Eptacog alfa [recombinant factor viia]         11         Minoxidil         16           Eptifibatide         16         Moducal         16           Erlotinib         8         Montelukast         15           Ethics Aspirin EC         6         Moroctocog alfa [recombinant factor viii]         11           Ethinyloestradiol with levonorgestrel         7         Morphine sulphate         14           F         N	. •			
Eptacog alfa [recombinant factor viia] 11 Minoxidil 16 Eptifibatide 16 Moducal 16 Erlotinib 8 Montelukast 15 Ethics Aspirin EC 6 Moroctocog alfa [recombinant factor viii] 11 Ethinyloestradiol with levonorgestrel 7 Morphine sulphate 14 F N	·			
Eptifibatide       16       Moducal       16         Erlotinib       8       Montelukast       15         Ethics Aspirin EC       6       Moroctocog alfa [recombinant factor viii]       11         Ethinyloestradiol with levonorgestrel       7       Morphine sulphate       14         F       N				
Erlotinib				
Ethics Aspirin EC	•			
Ethinyloestradiol with levonorgestrel		-		
F N				
Factor eight inhibitors bypassing agent		1	·	14
	Factor eight inhibitors bypassing agent	12	Nilstat	7

# Index

### Pharmaceuticals and brands

Nonacog alfa [recombinant factor ix] NovoSeven RT Nystatin	12 11 7
Octocog alfa [recombinant factor viii]	12
9 1	11
Olyppenol [phenol oily]	
Olanzapine	
Olanzine	15
Omnipaque	17
Ondansetron	17
Onex	18
Oral feed	
Oseltamivir	17
Oxytocin	13
Oxytocin BNM	13
P	
Paroxetine hydrochloride	
Pentasa	
Pharmacy Health Sorbolene with Glycerin	12
Phenoxymethylpenicillin [penicillin v] 4	
pms-Bosentan	7
Potassium permanganate	
Pramipexole hydrochloride	14
R	
Ramipex	14
Relenza Rotadisk	8
Revolade	6
Ropinirole hydrochloride	14

S	
Salbutamol	18
Seretide	9
Seretide Accuhaler	9
Singulair	
Sodium citro-tartrate	7
Spiractin 4,	
Spironolactone4,	12
Spirotone	4
Strattera	5
Sugammadex	5
Т	
Tarceva	8
Timolol	9
Timoptol XE	9
Tofranil	17
U	
Ural	7
V	
Ventolin	18
X	
Xyntha	11
Z	
Zanamivir	8
Zinc chloride	16
Zypine	15
Zypine ODT	15

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**Hospital Medicines List queries:** 

Freephone Information line 0800 66 00 50 (option 2)

Fax: 64 4 974 7819

Email: HML@pharmac.govt.nz

www.pharmac.health.nz/medicines/hospital-pharmaceuticals

Pharmaceutical Management Agency Level 9, 40 Mercer Street, PO Box 10-254, Wellington 6143, New Zealand Phone: 64 4 460 4990 - Fax: 64 4 460 4995 - www.pharmac.govt.nz Freephone Information line (9am-5pm weekdays) 0800 66 00 50

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