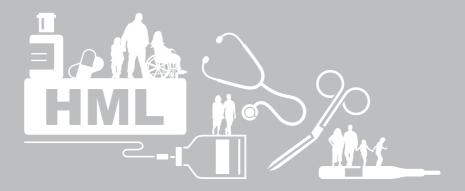
The Hospital Medicines List (HML)

Section H for Hospital Pharmaceuticals

Update effective 1 November 2015

Cumulative for August, September, October and November 2015





Contents

Summary of decisions effective 1 November 2015	. 3
Continue II also a months Doubli	_
Section H changes to Part II	. 6
Index	27

Summary of decisions EFFECTIVE 1 NOVEMBER 2015

- Aciclovir (Aciclovir-Claris) inj 250 mg vial new listing and addition of HSS
- Acivlocir (Zovirax IV) inj 250 mg vial to be delisted 1 January 2016
- Amino acid formula (without phenylalanine) (e.g. PKU Anamix Junior) powder
 36 g protein, 32 g carbohydrate and 12.5 g fat per 100 g, 36 g sachet
 amended presentation description
- Amino acid formula (without phenylalanine and tyrosine) (e.g. TYR Anamix Junior) powder 36 g protein, 32 g carbohydrate and 12.5 g fat per 100 g, 36 g sachet amended presentation description
- Aqueous cream (Pharmacy Health SLS-free) crm 100 g new listing and addition of HSS
- Aqueous cream (AFT) crm 100 g to be delisted 1 January 2016
- Atracurium besylate (Tracrium) inj 10 mg per ml, 2.5 ml and 5 ml ampoules
 price increase and addition of HSS
- Bisacodyl (Lax-Suppositories) suppos 10 mg new listing and addition of HSS
- Bisacodyl (Dulcolax) suppos 10 mg to be delisted 1 January 2016
- Blood ketone diagnostic test meter (Freestyle Optium Neo) meter new listing
- Blood ketone diagnostic test meter (Freestyle Optium) meter to be delisted
 1 May 2016
- Busulfan (Myleran) tab 2 mg price increase
- Cefoxitin (Cefoxitin Actavis) inj 1 g vial new listing and addition of HSS
- Cefoxitin (Hospira) inj 1 g vial to be delisted 1 January 2016
- Cetomacrogol (healthE) crm BP, 100 g price decrease and addition of HSS
- Citalopram hydrobromide (PSM Citalopram) tab 20 mg new listing and addition of HSS
- Citalopram hydrobromide (Arrow-Citalopram) tab 20 mg to be delisted 1 January 2016
- Cyclizine hydrochloride (Nauzene) tab 50 mg new listing and addition of HSS
- Cyclizine hydrochloride (Nausicalm) tab 50 mg to be delisted 1 January 2016
- Dexamethasone (Dexmethsone) tab 0.5 mg and 4 mg new listing and addition of HSS
- Dexamethasone (Douglas) tab 1 mg and 4 mg to be delisted 1 January 2016
- Factor eight inhibitor bypassing fraction (FEIBA NF) inj 500 U, 1,000 U and 2,500 U – amended chemical name and brand name
- Flucloxacillin (Flucloxin) inj 1 g vial addition of HSS

Summary of decisions - effective 1 November 2015 (continued)

- Flucloxacillin (DBL Flucloxacillin) inj 1 g vial to be delisted 1 January 2016
- Gabapentin (Neurontin) cap 100 mg, 300 mg and 400 mg new listing
- Lansoprazole (Lanzol Relief) cap 15 mg and 30 mg new listing and addition of HSS
- Lansoprazole (Solox) cap 15 mg and 30 mg to be delisted 1 January 2016
- Letrozole (Letrole) tab 2.5 mg new listing and addition of HSS
- Letrozole (Letraccord) tab 2.5 mg to be delisted 1 January 2016
- Lisinopril (Ethics Lisinopril) tab 5 mg, 10 mg and 20 mg new listing and addition of HSS
- Lisinopril (Arrow-Lisinopril) tab 5 mg, 10 mg and 20 mg to be delisted
 1 January 2016
- Mask for spacer device (e-chamber Mask) small new listing
- Mask for spacer device (EZ-fit Paediatric Mask) size 2 to be delisted
 1 February 2016
- Mirtazapine (Apo-Mirtazapine) tab 30 mg and 45 mg restriction removed
- Mixed salt solution for eye irrigation (Balanced Salt Solution) eye irrigation solution, 15 ml dropper bottle and 500 ml bottle – amended chemical name and presentation descriptions, new listing and addition of HSS
- Mixed salt solution for eye irrigation (e.g. Balanced Salt Solution) eye irrigation solution, 250 ml amended chemical name and presentation description
- Nonacog alfa [recombinant factor IX] (BeneFIX) inj 3,000 iu vial new listing
- Peak flow meter low range (Mini-Wright AFS Low Range) and normal range (Mini-Wright Standard) new listing
- Peak flow meter (Breath-Alert) low range and normal range to be delisted
 1 February 2016
- Sertraline (Arrow-Sertraline) tab 50 mg HSS suspended
- Spacer device 220 ml (single patient) (e-chamber Turbo) and 510 ml (single patient) (e-chamber La Grande) new listing
- Spacer device (Space Chamber Plus) spacer device 230 ml (single patient) to be delisted 1 February 2016
- Thiamine hydrochloride (e.g. Benerva) inj 100 mg per ml, 1 ml vial new listing
- Voriconazole (Vttack) tab 50 mg and 200 mg new listing and addition of HSS
- Voriconazole (Vfend) tab 50 mg and 200 mg to be delisted 1 January 2016
- Ziprasidone (Zusdone) cap 20 mg, 40 mg, 60 mg and 80 mg new listing and addition of HSS

Summary of decisions – effective 1 November 2015 (continued) • Ziprasidone (Zeldox) cap 20 mg, 40 mg, 60 mg and 80 mg – to be delisted 1 January 2016

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

Section H changes to Part II

Effective 1 November 2015

ALIM	ENTARY TRACT AND METABOLISM		
15	LANSOPRAZOLE Cap 15 mg – 1% DV Jan-16 to 2018	100 100	Lanzol Relief Lanzol Relief
20	BISACODYL Suppos 10 mg – 1% DV Jan-16 to 2018 3.78 Note – Dulcolax suppos 10 mg to be delisted from 1 January 2016.	10	Lax-Suppositories
25	THIAMINE HYDROCHLORIDE (new listing) Inj 100 mg per ml, 1 ml vial		e.g. Benerva
BL00	D AND BLOOD FORMING ORGANS		
29	FACTOR EIGHT INHIBITOR\$ BYPASSING FRACTION (amended chemical nar → Inj 500 U	me and brand 1 1 1	d name) FEIBA NF FEIBA FEIBA NF FEIBA FEIBA NF FEIBA
29	NONACOG ALFA [RECOMBINANT FACTOR IX] (new listing) → Inj 3,000 iu vial	1	BeneFIX
CARD	IOVASCULAR SYSTEM		
37	LISINOPRIL Tab 5 mg – 1% DV Jan-16 to 2018	90 90 90 January 20	Ethics Lisinopril Ethics Lisinopril Ethics Lisinopril 16.
DERN	IATOLOGICALS		
51	AQUEOUS CREAM Crm 100 g – 1% DV Jan-16 to 2018 1.00	100 g	Pharmacy Health SLS-free
	Note: DV limit applies to the pack sizes of 100 g or less. Note – AFT aqueous cream 100 g to be delisted from 1 January 2016.		OLO-1166
51	CETOMACROGOL (‡ price and addition of HSS) Crm BP, 100 g – 1% DV Jan-16 to 2018 1.47	1	healthE

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

Changes to Section H Part II – effective 1 November 2015 (continued)

HORM	ONE	DDED	ADAT	IUNG.
HUNIN	UNE	FNEF	ANAI	IUNO

weeks); or

60	DEXAMETHASONE Tab 0.5 mg – 1% DV Jan-16 to 2018	30 30 anuary 2010	Dexmethsone Dexmethsone 3.
INFE	CTIONS		
70	CEFOXITIN Inj 1 g vial – 1% DV Jan-16 to 2018 58.00 Note – Hospira cefoxitin inj 1 g vial to be delisted from 1 January 2016.	10	Cefoxitin Actavis
72	FLUCLOXACILLIN (addition of HSS) Inj 1 g vial – 1% DV Jan-16 to 2017 11.60 Note – DBL Flucloxacillin inj 1 g vial to be delisted from 1 January 2016.	10	Flucloxin
77	VORICONAZOLE → Tab 50 mg – 1% DV Jan-16 to 2018130.00 → Tab 200 mg – 1% DV Jan-16 to 2018500.00 Note – Vfend tab 50 mg and 200 mg to be delisted from 1 January 2016.	56 56	Vttack Vttack
88	ACICLOVIR Inj 250 mg vial – 1% DV Jan-16 to 2018 10.10 Note – Zovirax IV inj 250 mg vial to be delisted from 1 January 2016.	5	Aciclovir-Claris
MUS	CULOSKELETAL SYSTEM		
98	ATRACURIUM BESYLATE († price and addition of HSS) Inj 10 mg per ml, 2.5 ml ampoule – 1% DV Jan-16 to 2018 10.00 Inj 10 mg per ml, 5 ml ampoule – 1% DV Jan-16 to 2018 12.50	5 5	Tracrium Tracrium
NERV	OUS SYSTEM		
111	MIRTAZAPINE (restriction removed) Tab 30 mg – 1% DV Nov-15 to 2018	30 30	Apo-Mirtazapine Apo-Mirtazapine

2.1 The patient must have had a trial of two different antidepressants and was unable to tolerate the

treatments or failed to respond to an adequate dose over an adequate period of time (usually at least four

Price	
(ex man. Excl. GST)	
\$	Per

Brand or Generic Manufacturer

Neurontin

100

Changes to Section H Part II - effective 1 November 2015 (continued)

continued...

113

2.2 Both:

2.2.1 The patient is currently a hospital in-patient as a result of an acute depressive episode; and

2.2.2 The patient must have had a trial of one other antidepressant and either could not tolerate it or failed to respond to an adequate dose over an adequate period of time.

Continuation

Re-assessment required after two years

The patient has a high risk of relapse (prescriber determined)

112 CITALOPRAM HYDROBROMIDE

Tab 20 mg – 1% DV Jan-16 to 2018 Note – Arrow-Citalopram tab 20 mg to be delisted from		84	PSM Citalopran	n
GABAPENTIN (new listing)				
→ Cap 100 mg	7.16	100	Neurontin	
→ Cap 300 mg	11.00	100	Neurontin	

113 SERTRALINE (HSS suspended)

117 CYCLIZINE HYDROCHLORIDE

Tab 50 mg – 1% DV Jan-16 to 2018	0.59	20	Nauzene
Note - Nausicalm tab 50 mg to be delisted from 1 January	2016.		

120 ZIPRASIDONE

ZII TI/TOIDOINE			
→ Cap 20 mg – 1% DV Jan-16 to 2018	14.56	60	Zusdone
→ Cap 40 mg – 1% DV Jan-16 to 2018	24.75	60	Zusdone
→ Cap 60 mg – 1% DV Jan-16 to 2018	33.87	60	Zusdone
→ Cap 80 mg – 1% DV Jan-16 to 2018	39.74	60	Zusdone
Note - Zeldox cap 20 mg, 40 mg, 60 mg and 80 mg to be of	delisted from 1 Ja	anuary 201	6.

ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

128	BUSULFAN († price) Tab 2 mg89.25	100	Myleran
141	LETROZOLE Tab 2.5 mg – 1% DV Jan-16 to 2018 2.95 Note – Letraccord tab 2.5 mg to be delisted from 1 January 2016.	30	Letrole

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
Char	nges to Section H Part II – effective 1 November 2015 (cor	ntinued)	
SENS	SORY ORGANS		
179	MIXED SALT SOLUTION FOR EYE IRRIGATION CALCIUM CHLORIDE WARD SOLUTION FOR EYE IRRIGATION CALCIUM CHLORIDE WAS (amended chemical name and presentation description) Eye irrigation solution dreps calcium chloride 0.048% with magnesium chloride 0.03%, potassium chloride 0.075%, sodium acetate 0.39%, sodium chloride 0.64% and sodium citrate 0.17%, 15 ml dropper bottle		,
	- 1% DV Jan-16 to 2018 (new listing)	15 ml	Balanced Salt Solution e.g. Balanced Sali Solution
	Eye irrigation solution drops calcium chloride 0.048% with magnesium chloride 0.03%, potassium chloride 0.075%, sodium acetate 0.39%, sodium chloride		Columbia
	0.64% and sodium citrate 0.17%, 250 ml		e.g. Balanced Sal Solution
	Eye irrigation solution drops calcium chloride 0.048% with magnesium chloride 0.03%, potassium chloride 0.075%, sodium acetate 0.39%, sodium chloride 0.64% and sodium citrate 0.17%, 500 ml bottle		
	- 1% DV Jan-16 to 2018 (new listing)10.50	500 ml	Balanced Salt Solution e.g. Balanced Sal Solution
SPEC	CIAL FOODS		
197	AMINO ACID FORMULA (WITHOUT PHENYLALANINE) (presentation des → Powder 36 g 29 g protein, 32 g 38 g carbohydrate and	cription amend	•
	12.5 g fat 13.5 g fibre per 100 g, 36 g 29 g sachet		e.g. PKU Anamix Junior
98	AMINO ACID FORMULA (WITHOUT PHENYLALANINE AND TYROSINE) (→ Powder 36 g 29 g protein, 32 g 38 g carbohydrate and	presentation de	·
	12.5 g 13.5 g fat per 100 g, 36 g 29 g g sachet		e.g. TYR Anamix Junior
PART	TIII – OPTIONAL PHARMACEUTICALS		

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

214	MASK FOR SPACER DEVICE Small Note – EZ-fit Paediatric Mask mask for spacer device		-	
214	PEAK FLOW METER			
	Low range	9.54	1	Mini-Wright AFS Low Range
	Normal range	9.54	1	Mini-Wright Standard
	Note – Breath-Alert peak flow meter low range and no	ormal range to be deli	sted from	1 February 2016.
215	SPACER DEVICE			
	220 ml (single patient)	2.95	1	e-chamber Turbo
	510 ml (single patient)	5.12	1	e-chamber La Grande
	Note – Space Chamber Plus spacer device 230 ml (s	single patient) to be de	listed fron	n 1 February 2016.

Effective 1 October 2015

18

ALIMENTARY TRACT AND METABOLISM

10	Tab immediate-release 850 mg – 1% DV Dec-15 to 20187.82	500	Metformin Mylan
	Note – Apotex metformin hydrochloride tab immediate-release 850 mg to	be delist	ed from 1 December 2015.
18	PIOGLITAZONE		
	Tab 15 mg – 1% DV Dec-15 to 2018	90	Vexazone
	Tab 30 mg – 1% DV Dec-15 to 2018	90	Vexazone
	Tab 45 mg – 1% DV Dec-15 to 2018	90	Vexazone
18	PIOGLITAZONE (↓ price and delisting)		
	Tab 15 mg	28	Pizaccord
	Tab 30 mg1.57	28	Pizaccord
	Tab 45 mg2.21	28	Pizaccord
	Note – Pizaccord tab 15 mg, 30 mg and 45 mg to be delisted from 1 Dec	ember 20	15.
20	BISACODYL (delisting)		
	Suppos 5 mg3.00	6	Dulcolax
	Note – Dulcolax suppos 5 mg to be delisted from 1 December 2015.		
23	MULTIVITAMIN AND MINERAL SUPPLEMENT (new listing and amended res	striction)	
	→ Cap	180	Clinicians Multivit & Mineral Boost e.g.Clinicians Multivit &
			Mineral Boost

METEORMIN HYDROCHLORIDE (amended chemical name, new listing and addition of HSS)

Restricted

Limited to 3 months' treatment

Both:

- 1 Patient was admitted to hospital with burns; and
- 2 Any of the following:
 - 2.1 Burn size is greater than 15% of total body surface area (BSA) for all types of burns; or
 - 2.2 Burn size is greater than 10% of BSA for mid-dermal or deep dermal burns; or

continued...

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

Changes to Section H Part II - effective 1 October 2015 (continued)

continued...

2.3 Nutritional status prior to admission or dietary intake is poor.

Note: Multivitamin and mineral supplement capsule composition includes vitamin A 250 IU, thiamine 2.5 mg, riboflavin 2.5 mg, nicotinamide 12.5 mg, vitamin B5 10 mg, pyridoxine 5 mg, vitamin B12 6.2 meg, vitamin C 125 mg, cholecalciferol 2.5 meg, vitamin E 25 mg, betaine 12.5 mg, biotin 12.5 meg, boron 250 meg, calcium 25 mg, choline 6.2 mg, chromium 25 meg, citric acid 50mg, citrus bioflavonoid complex 50mg, co-enzyme Q10 1.2 mg, copper 125 meg, folic acid 37.5 meg, inositol 6.2 mg, iodine 25 meg, iron 250 meg, L-Glutamine 6.2 mg, magnesium 12.5 mg, molybdenum 12.5 meg, manganese 0.5 mg, potassium 5 mg, selenium 18.7 meg, zinc 1.9 mg.

24	MUL	_TIVIT	AMIN	RENAL

Either:

- 1 The patient has chronic kidney disease and is receiving either peritoneal dialysis or haemodialysis; or
- 2 The patient has chronic kidney disease grade 5, defined as patient with an estimated glomerular filtration rate of <15 ml/min/1.73 m² body surface area (BSA).</p>

BLOOD AND BLOOD FORMING ORGANS

DLUU	D AND BLOOD FORWING ORGANS		
28	ALUMINIUM CHLORIDE → Topical soln 20% w/v Restricted For use as a haemostasis agent.		e.g. Driclor
29	FACTOR EIGHT INHIBITORS BYPASSING FRACTION → Inj 2,500 U	1	FEIBA
CARD	IOVASCULAR SYSTEM		
39	FLECAINIDE ACETATE (delisting) Tab 100 mg68.78 Note – Tambocor tab 100 mg to be delisted from 1 December 2015.	60	Tambocor
45	DOBUTAMINE HYDROCHLORIDE (amended presentation description and n Inj 12.5 mg per ml, 20 ml ampoule vial - 1% DV Jan-16 to 2018	ew listing) 5	Dobutamine-Claris
DERN	IATOLOGICALS		
50	CALAMINE (4 price and addition of HSS) Crm, aqueous, BP – 1% DV Dec-15 to 2018	100 g 2,000 ml	Pharmacy Health PSM
50	ISOTRETINOIN (4 price and delisting) Cap 10 mg	120 120 015.	Oratane Oratane

Price	
(ex man. Excl. GST)	
\$	Per

Brand or Generic Manufacturer

Changes to Section H Part II – effective 1 October 2015 (continued)

HORMONE PREPARATIONS

59 **OXANDROLONE** OXANDROLINE (chemical name change)

→ Tab 2.5 mg

MUSCULOSKELETAL SYSTEM

100	DICLOFENAC SODIUM
100	DIOLOI LIVAO OODIOIVI

Tab EC 25 mg - 1% DV Dec-15 to 2018	1.30	50	Diclofenac Sandoz
Tab EC 50 mg - 1% DV Dec-15 to 2018	1.00	50	Diclofenac Sandoz
Tab long-acting 75 mg - 1% DV Dec-15 to 2018	15.20	500	Apo-Diclo SR
Tab long-acting 100 mg - 1% DV Dec-15 to 2018	26.20	500	Apo-Diclo SR
Note - Apo-Diclo tab EC 25 mg and 50 mg, and Dicla	x SR tab long-acting	75 mg a	nd 100 mg to be delisted
from 1 December 2015.			

NERVOUS SYSTEM

NERV	009 9191EM		
104	BUPIVACAINE HYDROCHLORIDE (Pharmacode change) Inj 5 mg per ml, 10 ml ampoule sterile pack – 1% DV Sep-15 to 2018	5 359 to be de	Marcain listed from 1 October
107	PARACETAMOL Suppos 125 mg – 1% DV Dec-15 to 2018	10 10 mber 2015.	Gacet Gacet
110	OXYCODONE HYDROCHLORIDE (\$\pi\ price and addition of HSS) Inj 50 mg per ml, 1 ml ampoule — 1% DV Dec-15 to 201851.00	5	OxyNorm
113	GABAPENTIN → Tab 600 mg Note – Gabapentin tab 600 mg to be delisted 1 November 2015.		
115	LAMOTRIGINE (delisting) Tab dispersible 25 mg	56 56 56 from 1 Dece	Mogine Mogine Mogine ember 2015.
115	PHENOBARBITONE († price and addition of HSS) Tab 15 mg – 1% DV Dec-15 to 2018	500 500	PSM PSM
117	DOMPERIDONE (‡ price and addition of HSS) Tab 10 mg – 1% DV Dec-15 to 2018	100	Prokinex
118	ARIPIPRAZOLE → Tab 5 mg123.54	30	Abilify

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
Char	nges to Section H Part II – effective 1 October 2015 (continu	ied)	
124	ZOPICLONE Tab 7.5 mg – 1% DV Dec-15 to 2018	30 500	Zopiclone Actavis Zopiclone Actavis
	Note – Apo-Zopiclone tab 7.5 mg to be delisted from 1 December 2015		Zupicione Actavis
125	DEXAMFETAMINE SULFATE († price and addition of HSS) → Tab 5 mg – 1% DV Dec-15 to 201817.00	100	PSM
ONC	OLOGY AGENTS AND IMMUNOSUPPRESSANTS		
141	ETANERCEPT (↓ price) → Inj 25 mg vial	4 4 4	Enbrel Enbrel Enbrel
RESI	PIRATORY SYSTEM AND ALLERGIES		
173	SALBUTAMOL Aerosol inhaler, 100 mcg per dose CFC free	200 dose	SalAir
174	FLUTICASONE Aerosol inhaler 50 mcg per dose	120 dose 120 dose 120 dose	Floair Floair Floair
174	SALMETEROL Aerosol inhaler 25 mcg per dose26.46	120 dose	Meterol
175	FLUTICASONE WITH SALMETEROL Aerosol inhaler 50 mcg with salmeterol 25 mcg	120 dose 120 dose	RexAir RexAir
SENS	SORY ORGANS		
178	PREDNISOLONE SODIUM PHOSPHATE (new listing) Eye drops 0.5%, single dose (preservative free)	20 dose	Minims Prednisolone
180	DORZOLAMIDE WITH TIMOLOL Eye drops 2% with timolol 0.5% – 1% DV Dec-15 to 20183.45 Note – Cosopt eye drops to be delisted from 1 December 2015.	5 ml	Arrow-Dortim

	Price		Brand or
(ex r	nan. Excl. 6	GST)	Generic
	\$	Per	Manufacturer

Changes to Section H Part II - effective 1 October 2015 (continued)

SPECIAL FOODS

207 ORAL FEED

Formula (Vanilla)

Note: Community subsidy of Sustagen Hospital Formula is subject to both Special Authority criteria and a manufacturer's surcharge. Higher subsidy by endorsement is available for patients meeting the following endorsement criteria; fat malabsorption, fat intolerance or chyle leak.

Note – Sustagen Hospital Formual (chocolate and vanilla) powder 900 g can to be delisted from 1 April 2016.

Effective 1 September 2015

ALIMENTARY TRACT AND METABOLISM

18 METEORMIN

Tab immediate-release 500 mg – **1% DV Nov-15 to 2018**........9.59 1,000 **Metchek** Note – Apotex metformin tab immediate-release 500 mg to be delisted from 1 November 2015.

BLOOD AND BLOOD FORMING ORGANS

29 EPTACOG ALFA [RECOMBINANT FACTOR VIIA] (amended restriction)

)	Inj 1 mg syringe	1,163.75	1	NovoSeven RT
-	Inj 2 mg syringe	2,327.50	1	NovoSeven RT
-	Inj 5 mg syringe	5,818.75	1	NovoSeven RT
	, , , ,	9,310.00	1	NovoSeven RT
_	, , , , , ,	•		

Restricted

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

29 FACTOR EIGHT INHIBITORS BYPASSING **FRACTION** AGENT (amended chemical name and restriction, and ↓ price)

→	· Inj 500 U	1,450.00	1	FEIBA
→	Inj 1,000 U	2,900.00	1	FEIBA

Restricted

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

29 MOROCTOCOG ALFA [RECOMBINANT FACTOR VIII] (amended presentation description and 1 price)

→ Inj 250 iu prefilled syringe vial	210.00	1	Xyntha
→ Inj 500 iu prefilled syringe vial	420.00	1	Xyntha
→ Inj 1,000 iu prefilled syringe vial	840.00	1	Xyntha
→ Inj 2,000 iu prefilled syringe vial	1,680.00	1	Xyntha
→ Inj 3,000 iu prefilled syringe vial	2,520.00	1	Xyntha

		Price (ex man. Excl. GST) \$ F	Per	Brand or Generic Manufacturer
Chai	nges to Section H Part II – effective 1	September 2015 (conf	inued)	
29	NONACOG ALFA [RECOMBINANT FACTOR IX]	(amended restriction)		
	→ Inj 250 iu vial		1	BeneFIX
	→ Inj 500 iu vial		1	BeneFIX
	→ Inj 1,000 iu vial		1	BeneFIX
	→ Inj 2,000 iu vial	2,480.00	1	BeneFIX
	Restricted When used in the treatment of haemophilia, ac Group in conjunction with the National Haemo		managed by	the Haemophilia Treaters
29	OCTOCOG ALFA [RECOMBINANT FACTOR VIII] (↓ price)		
	→ Inj 250 iu vial	237.50	1	Kogenate FS
	→ Inj 500 iu vial		1	Kogenate FS
	→ Inj 1,000 iu vial	950.00	1	Kogenate FS
	→ Inj 2,000 iu vial	1,900.00	1	Kogenate FS
	→ Inj 3,000 iu vial	2,850.00	1	Kogenate FS
00	OOTOOOO ALEA IDEOOMBINANT FAOTOD WIII	1 (A mains)		
29	OCTOCOG ALFA [RECOMBINANT FACTOR VIII		1	Advoto
	→ Inj 250 iu vial		1	Advate
	→ Inj 500 iu vial → Inj 1,000 iu vial		1	Advate
			1	Advate
	→ Inj 1,500 iu vial			Advate
	→ Inj 2,000 iu vial		1 1	Advate
	→ Inj 3,000 iu vial	3,450.00	I	Advate
31	HEPARIN SODIUM (amended presentation des Inj 1,000 iu per ml, 35 ml vial ampoule	cription)		
32	TICAGRELOR (amended restriction)			
	→ Tab 90 mg	90.00	56	Brilinta
	Restricted Restricted to treatment of acute coronary synd 60 days) been diagnosed with an ST-elevation fibrinolytic therapy has not been given in the la	or a non-ST-elevation acute	coronary sy	
CAR	DIOVASCULAR SYSTEM			
48	EPOPROSTENOL (new listing)			
	→ Inj 0.5 mg vial	36.61	1	Veletri
	→ Inj 1.5 mg vial		1	Veletri
	Restricted			
	For use as a bridge to transplant for patients w list for lung transplantation.	ith Pulmonary Arterial Hype	tension who	are on the active waiting
DER	MATOLOGICALS			
49	HYDROGEN PEROXIDE Soln 3% (10 vol) – 1% DV Nov-15 to 2018	1.40	100 ml	Pharmacy Health
50	DIMETHICONE			
	Crm 10% pump bottle – 1% DV Nov-15 to 2	2018 4.90	500 ml	healthE Dimethicone 10%

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
Cha	nges to Section H Part II – effective 1 September 2015 (co	ntinued)	
50	ISOTRETINOIN Cap 10 mg - 1% DV Nov-15 to 2018	100 100 r 2015.	Isotane 10 Isotane 20
51	CETOMACROGOL Crm BP, 500 g – 1% DV Nov-15 to 2018 2.74 Note – Pharmacy Health cetomacrogol crm BP, 500 g to be delisted f		healthE per 2015.
52	MOMETASONE FUROATE Crm 0.1% – 1% DV Nov-15 to 2018	15 g 50 g	Elocon Alcohol Free Elocon Alcohol Free
	Oint 0.1% – 1% DV Nov-15 to 2018	15 g 50 g	Elocon Elocon
GEN	ITO-URINARY SYSTEM	, delisted from	T NOVELLIDE ZOTO.
57	FINASTERIDE (pack size change) → Tab 5 mg – 1% DV Dec-14 to 20172.08 Note – The pack size has changed from 28 to 30 tab. The 28 tab pace 2015.		Finpro ed from 1 November
57	OXYTOCIN (\$\psi\$ price and addition of HSS) Inj 5 iu per ml, 1 ml ampoule — 1% DV Nov-15 to 2018		Oxytocin BNM Oxytocin BNM BNM
INFE	ECTIONS		
69	TOBRAMYCIN († price) → Inj 40 mg per ml, 2 ml vial38.00	5	DBL Tobramycin
72	PIPERACILLIN WITH TAZOBACTAM (new listing) → Inj 4 g with tazobactam 0.5 g vial	1	Hospira
72	PIPERACILLIN WITH TAZOBACTAM (delisting) → Inj 4 g with tazobactam 0.5 g vial - 1% DV Oct-13 to 31/8/15 2016		Tazocin EF er 2015.
82	NEVIRAPINE (↓ price and addition of HSS) → Tab 200 mg – 1% DV Nov-15 to 201865.00	60	Nevirapine Alphapharm
MUS	SCULOSKELETAL SYSTEM		
98	DANTROLENE (new listing) Inj 20 mg vial800.00	6	Dantrium IV e.g. Dantrium IV

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

Changes to Section H Part II – effective 1 September 2015 (continued)

NERVOUS SYSTEM

104	PROPOFOL (delisting) Inj 10 mg per ml, 100 ml vial30.00 Note – Diprivan inj 10 mg per ml, 100 ml vial to be delisted from 1 Nover	1 nber 2015.	Diprivan
107	PARACETAMOL (‡ price and addition of HSS) Suppos 500 mg – 1% DV Nov-15 to 201812.60	50	Paracare
110	PETHIDINE HYDROCHLORIDE († price and addition of HSS) Tab 50 mg – 1% DV Nov-15 to 2018	10 10	PSM PSM
111	MIRTAZAPINE → Tab 30 mg – 1% DV Nov-15 to 2018	30 30 15.	Apo-Mirtazapine Apo-Mirtazapine
120	ZIPRASIDONE Inj 20 mg Inj 100 mg Note – Ziprasidone inj 20 mg and 100 mg to be delisted from 1 March 20	016.	
122	ZUCLOPENTHIXOL DECANOATE (new listing) Inj 500 mg per ml, 1 ml ampoule		e.g. Clopixol Conc
ONCO	LOGY AGENTS AND IMMUNOSUPPRESSANTS		
128	THIOTEPA (new listing) Inj 100 mg vial		
129	EPIRUBICIN HYDROCHLORIDE Inj 2 mg per ml, 25 ml vial – 1% DV Nov-15 to 2018	1 1 1 00 ml vials	Epirubicin Ebewe Epirubicin Ebewe Epirubicin Ebewe to be delisted from 1
129	IDARUBICIN HYDROCHLORIDE († price and addition of HSS) Inj 5 mg vial – 1% DV Nov-15 to 2018	1 1	Zavedos Zavedos
133	CISPLATIN Inj 1 mg per ml, 50 ml vial – 1% DV Nov-15 to 2018	1 1 sted from 1	DBL Cisplatin DBL Cisplatin November 2015.
138	CALCIUM FOLINATE († price) Tab 15 mg104.26	10	DBL Leucovorin

Calcium

Pri	ice		Brand or
(ex man. I	Excl. GST)		Generic
	\$	Per	Manufacturer

Changes to Section H Part II – effective 1 August 2015

ALIMENTARY	TRACT AND	METABOLISM

ALIN	IENTARY TRACT AND METABULISM		
14	HYDROCORTISONE ACETATE (amended presentation, † price and addition Rectal foam 10%, CFC free (14 applications) - 1% DV Oct-15 to 2018 26.55	of HSS) 21.1 g	Colifoam
16	ACARBOSE Tab 50 mg – 1% DV Oct-15 to 2018	90 90 5.	Glucobay Glucobay
18	PANCREATIC ENZYME Cap EC 10,000 BP u lipase, 9,000 BP u amylase and 210 BP u protease – 1% DV 0ct-15 to 2018	100	Creon 10000
	1,000 BP u protease – 1% DV Oct-15 to 2018 94.38	100	Creon 25000
20	BISACODYL († price and addition of HSS) Tab 5 mg – 1% DV Oct-15 to 2018	200	Lax-Tabs
Char	nges to Section H Part II – effective 1 August 2015 (continued	d)	
BLO	DD AND BLOOD FORMING ORGANS		
28	FOLIC ACID Tab 0.8 mg – 1% DV Oct-15 to 2018	1,000 500	Apo-Folic Acid Apo-Folic Acid
35	POTASSIUM DIHYDROGEN PHOSPHATE Inj 1 mmol per ml, 10 ml ampoule - 1% DV Oct-15 to 2018151.80	10	Hospira
35	SODIUM DIHYDROGEN PHOSPHATE [SODIUM ACID PHOSPHATE] Inj 1 mmol per ml, 20 ml ampoule - 1% DV Oct-15 to 201847.50	5	Biomed
CARI	DIOVASCULAR SYSTEM		
37	QUINAPRIL WITH HYDROCHLOROTHIAZIDE († price and addition of HSS) Tab 10 mg with hydrochlorothiazide 12.5 mg	20	Accuretic 10
	- 1% DV Oct-15 to 2018	30 30	Accuretic 20
39	MEXILETINE HYDROCHLORIDE († price) Cap 150 mg162.00	100	Mexiletine
	Cap 250 mg202.00	100	Hydrochloride USP Mexiletine Hydrochloride USP

		Price (ex man. Excl. GST) \$ Pe	r	Brand or Generic Manufacturer
Chai	nges to Section H Part II – effective 1 Aug	gust 2015 (continued)		
40	NADOLOL († price and addition of HSS) Tab 40 mg – 1% DV Oct-15 to 2018 Tab 80 mg – 1% DV Oct-15 to 2018		100 100	Apo-Nadolol Apo-Nadolol
43	BEZAFIBRATE (addition of HSS) Tab 200 mg – 1% DV Oct-15 to 2018 (‡ price) Tab long-acting 400 mg – 1% DV Oct-15 to 20 1		90 30	Bezalip Bezalip Retard
46	ALPROSTADIL HYDROCHLORIDE († price and add Inj 500 mcg per ml, 1 ml ampoule – 1% DV Oct-15 to 2018	,	5	Prostin VR
47	BOSENTAN → Tab 62.5 mg – 1% DV Jan-16 to 2018 → Tab 125 mg – 1% DV Jan-16 to 2018 Note – Pms-Bosentan and Tracleer tab 62.5 mg	375.00	56 56 d from 1 J	Mylan-Bosentan Mylan-Bosentan anuary 2016.
DER	MATOLOGICALS			
49	LINDANE [GAMMA BENZENE HEXACHLORIDE] Crm 1% Note – Lindane [gamma benzene hexachloride c	ream 1% to be delisted fro	om 1 Janu	ary 2016.
GEN	ITO-URINARY SYSTEM			
56	NORETHISTERONE Tab 350 mcg – 1% DV Oct-15 to 2018	6.25	84	Noriday 28
HOR	MONE PREPARATIONS			
59	CYPROTERONE ACETATE Tab 50 mg – 1% DV Oct-15 to 2018 Tab 100 mg – 1% DV Oct-15 to 2018 Note – Siterone tab 50 mg and 100 mg to be de	30.40	50 50 5.	Procur Procur
60	METHYLPREDNISOLONE (AS SODIUM SUCCINATI Tab 4 mg – 1% DV Oct-15 to 2018		HSS) 100 20 1 1	Medrol Medrol Solu-Medrol Solu-Medrol
60	METHYLPREDNISOLONE (AS SODIUM SUCCINATI Inj 500 mg vial – 1% DV Oct-15 to 2018 Inj 1 g vial – 1% DV Oct-15 to 2018	9.00	HSS) 1 1	Solu-Medrol Solu-Medrol
60	METHYLPREDNISOLONE ACETATE († price and ac Inj 40 mg per ml, 1 ml vial – 1% DV Oct-15 to 2		5	Depo-Medrol

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

Changes to Section H Part II - effective 1 August 2015 (continued)

60 METHYLPREDNISOLONE ACETATE WITH **LIDOCAINE** [LIGNOCAINE] (amended chemical and presentation descriptions, † price and addition of HSS)

Inj 40 mg with lidocaine [lignocaine] 10 mg per ml,

INFECTIONS

70 CEFEPIME

→ Inj 1 g vial – 1% DV Oct-15 to 2018	3.95	1	Cefepime-AFT
→ Inj 2 g vial – 1% DV Oct-15 to 2018	6.92	1	Cefepime-AFT
Note - DBL Cefepime inj 1 g and 2 g vials to	be delisted from 1 October 20	15.	-

71 AZITHROMYCIN (amended presentation description. † price and addition of HSS)

50 **TMP**

MUSCULOSKELETAL SYSTEM

94 ZOLEDRONIC ACID (amended restriction)

Inherited bone fragility disorders

Patient has been diagnosed with an inherited bone fragility disorder (e.g. osteogenesis imperfecta).

Osteoporosis

Both:

- 1 Any of the following:
 - 1.1 History of one significant osteoporotic fracture demonstrated radiologically and documented bone mineral density (BMD) ≥ 2.5 standard deviations below the mean normal value in young adults (i.e. T-Score ≤ -2.5) (see Note); or
 - 1.2 History of one significant osteoporotic fracture demonstrated radiologically, and either the patient is elderly, or densitometry scanning cannot be performed because of major logistical, technical or pathophysiological reasons.

It is unlikely that this provision would apply to many patients under 75 years of age; or

- 1.3 History of two significant osteoporotic fractures demonstrated radiologically; or
- 1.4 Documented T-Score ≥ -3.0 (see Note); or
- 1.5 A 10-year risk of hip fracture ≥ 3%, calculated using a published risk assessment algorithm (e.g. FRAX or Garvan) which incorporates BMD measurements (see Note); or
- 1.6 Patient has had a Special Authority approval for alendronate (Underlying cause Osteoporosis) or raloxifene: and
- 2 The patient will not be prescribed more than 5 mg of zoledronic acid one infusion in a 12-month period.

Initiation - glucocorticosteroid therapy

Re-assessment required after 12 months

All of the following:

- 1 The patient is receiving systemic glucocorticosteroid therapy (≥ 5 mg per day prednisone equivalents) and has already received or is expected to receive therapy for at least three months; and
- 2 Any of the following:

continued...

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

Changes to Section H Part II – effective 1 August 2015 (continued)

continued...

- 2.1 The patient has documented BMD ≥ 1.5 standard deviations below the mean normal value in young adults (i.e. T-Score ≤ -1.5) (see Note); or
- 2.2 The patient has a history of one significant osteoporotic fracture demonstrated radiologically; or
- 2.3 The patient has had a Special Authority approval for alendronate (Underlying cause glucocorticosteroid therapy) or raloxifene: and
- 3 The patient will not be prescribed more than 5 mg of zoledronic acid one infusion in the 12-month approval period.

Continuation - glucocorticosteroid therapy

Re-assessment required after 12 months

Both:

- 1 The patient is continuing systemic glucocorticosteriod therapy (≥ 5 mg per day prednisone equivalents); and
- 2 The patient will not be prescribed more than 5 mg of zoledronic acid one infusion in the 12-month approval period.

Initiation - Paget's disease

Re-assessment required after 12 months

All of the following:

- 1 Paget's disease; and
- 2 Any of the following:
 - 2.1 Bone or articular pain; or
 - 2.2 Bone deformity: or
 - 2.3 Bone, articular or neurological complications; or
 - 2.4 Asymptomatic disease, but risk of complications; or
 - 2.5 Preparation for orthopaedic surgery; and
- 3 The patient will not be prescribed more than 5 mg of zoledronic acid one infusion in the 12-month approval period.

Continuation - Paget's disease

Re-assessment required after 12 months

Both:

- 1 Any of the following:
 - 1.1 The patient has relapsed (based on increases in serum alkaline phosphatase); or
 - 1.2 The patient's serum alkaline phosphatase has not normalised following previous treatment with zoledronic acid: or
 - 1.3 Symptomatic disease (prescriber determined); and
- 2 The patient will not be prescribed more than 5 mg of zoledronic acid one infusion in the 12-month approval period.

Notes:

- 1 BMD (including BMD used to derive T-Score) must be measured using dual-energy x-ray absorptiometry (DXA). Quantitative ultrasound and quantitative computed tomography (QCT) are not acceptable.
- 2 Evidence suggests that patients aged 75 years and over who have a history of significant osteoporotic fracture demonstrated radiologically are very likely to have a T-Score ≤ -2.5 and, therefore, do not require BMD measurement for treatment with bisphosphonates.
- 3 Osteoporotic fractures are the incident events for severe (established) osteoporosis and can be defined using the WHO definitions of osteoporosis and fragility fracture. The WHO defines severe (established) osteoporosis as a T-score below -2.5 with one or more associated fragility fractures. Fragility fractures are fractures that occur as a result of mechanical forces that would not ordinarily cause fracture (minimal trauma). The WHO has quantified this as forces equivalent to a fall from a standing height or less.
- 4 A vertebral fracture is defined as a 20% or greater reduction in height of the anterior or mid portion of a vertebral body relative to the posterior height of that body, or a 20% or greater reduction in any of these heights compared to the vertebral body above or below the affected vertebral body.

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

Changes to Section H Part II – effective 1 August 2015 (continued)

97 BENZBROMARONE (amended restriction)

Restricted

All of the following Both:

- 1 Patient has been diagnosed with gout; and
- 2 Any of the following:
 - 24.1 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 addition of probenecid at doses of up to 2 g per day or maximum tolerated dose; or
 - 24.2 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 use of probenecid at doses of up to 2 g per day or maximum tolerated dose; or
 - 21.3 Both:
 - 24.3.1 The patient has renal impairment such that probenecid is contraindicated or likely to be ineffective and serum urate remains greater than 0.36 mmol/l despite optimal treatment with allopurinol (see Notes): and
 - 24.3.2 The patient has a rate of creatinine clearance greater than or equal to 20 ml/min; or
 - **2**⁺.4 All of the following:
 - 24.4.1 The patient is taking azathioprine and requires urate-lowering therapy; and
 - 24.4.2 Allopurinol is contraindicated; and
 - 24.4.3 Appropriate doses of probenecid are ineffective or probenecid cannot be used due to reduced renal function; and
- 32 The patient is receiving monthly liver function tests.

Notes: Benzbromarone has been associated with potentially fatal hepatotoxicity. In chronic renal insufficiency, particularly when the glomerular filtration rate is 30 ml/minute or less, probenecid may not be effective.

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

www.rheumatology.org.nz/downloads/Benzbromarone-prescriber-information-NZRA-V2.pdf

98 FEBUXOSTAT (amended restriction)

→ Tab 80 mg	39.50	28	Adenurio
→ Tab 120 mg	39.50	28	Adenurio

Restricted

Roth:

- 1 Patient has been diagnosed with gout; and
- 2 Any of the following:
 - 2.1 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 addition of probenecid at doses of up to 2 g per day or maximum tolerated dose: or
 - 2.2 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 use of probenecid at doses of up to 2 q per day or maximum tolerated dose; or
 - 2.3 Both:
 - 3.1 The patient has renal impairment such that probenecid is contraindicated or likely to be ineffective and serum urate remains greater than 0.36 mmol/l despite optimal treatment with allopurinol (see Note); and

continued...

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

Changes to Section H Part II - effective 1 August 2015 (continued)

continued...

3.2 The patient has a rate of creatinine clearance greater than or equal to 30 ml/min.

Note: In chronic renal insufficiency, particularly when the glomerular filtration rate is 30 ml/minute or less, probenecid may not be effective. The efficacy and safety of febuxostat have not been fully evaluated in patients with severe renal impairment (creatinine clearance less than 30 ml/minute). No dosage adjustment of febuxostat is necessary in patients with mild or moderate renal impairment. 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.

NERVOUS SYSTEM

LIDOCVINE (LIGNOCVINE)

105

105	Crm 4%	30 g 5	LMX4 LMX4
107	PARACETAMOL Tab soluble 500 mg – 1% DV Oct-15 to 2017	20	Paragesic Soluble
108	MORPHINE HYDROCHLORIDE (addition of HSS) Oral liq 1 mg per ml – 1% DV Oct-15 to 2018	200 ml	RA-Morph
108	MORPHINE HYDROCHLORIDE († price and addition of HSS) Oral liq 2 mg per ml – 1% DV Oct-15 to 2018	200 ml 200 ml 200 ml	RA-Morph RA-Morph RA-Morph
110	OXYCODONE HYDROCHLORIDE (4 price and addition of HSS) Cap immediate-release 5 mg – 1% DV Oct-15 to 2018	20 20 20	OxyNorm OxyNorm OxyNorm
111	MOCLOBEMIDE († price and addition of HSS) Tab 150 mg – 1% DV Oct-15 to 2018	500 100	Apo-Moclobemide Apo-Moclobemide
113	PHENYTOIN SODIUM Inj 50 mg per ml, 2 ml ampoule – 1% DV Oct-15 to 2018 88.63 Inj 50 mg per ml, 5 ml ampoule – 1% DV Oct-15 to 2018 133.92	5 5	Hospira Hospira
118	ARIPIPRAZOLE (amended restriction) → Tab 10 mg	30 30 30 30	Abilify Abilify Abilify Abilify

- 1 Patient is suffering from schizophrenia or related psychoses; and
- 2 Either:
 - 2.1 An effective dose of risperidone or quetiapine has been trialled and has been discontinued, or is in the process of being discontinued, because of unacceptable side effects; or continued...

Price	
(ex man. Excl. GST)	
\$	Per

Brand or Generic Manufacturer

Changes to Section H Part II – effective 1 August 2015 (continued)

continued...

2.2 An effective dose of risperidone or quetiapine has been trialled and has been discontinued, or is in the process of being discontinued, because of inadequate clinical response.

Initiation – Autism spectrum disorder*

Psychiatrist or paediatrician

All of the following:

- 1 The patient has been diagnosed with an autism spectrum disorder* and has symptoms of severe irritability; and
- 2 An effective dose of risperidone has been trialled and has been discontinued because of unacceptable side effects or inadequate response; and
- 3 The patient is aged less than 18 years.

Note: Indications marked with * are Unapproved Indications

120 QUETIAPINE

Note – These are listing for new Pharmacodes, 2476266 and 2476274. The old Pharmacodes will be delisted from 1 February 2016.

127 NICOTINE (new listing)

→ Oral spray 1 mg per dose

e.g. Nicorette QuickMist Mouth Spray

Restricted

Any of the following:

- 1 For perioperative use in patients who have a 'nil by mouth' instruction; or
- 2 For use within mental health inpatient units; or
- 3 For acute use in agitated patients who are unable to leave the hospital facilities.

ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

128	BLEOMYCIN SULPHATE Inj 15,000 iu (10 mg) vial – 1% DV Oct-15 to 2018 150.48	1	DBL Bleomycin Sulfate
128	CYCLOPHOSPHAMIDE (addition of HSS) Inj 1 g vial – 1% DV Oct-15 to 2018	1	Endoxan Endoxan
128	DACTINOMYCIN [ACTINOMYCIN D] Inj 0.5 mg vial145.00	1	Cosmegen
130	FLUOROURACIL († price and addition of HSS) Inj 50 mg per ml, 20 ml vial – 1% DV Oct-15 to 201810.00	1	Fluorouracil Ebewe
130	FLUOROURACIL (\$\frac{1}{2}\$ price and addition of HSS) Inj 50 mg per ml, 50 ml vial - 1% DV Oct-15 to 2018	1	Fluorouracil Ebewe Fluorouracil Ebewe

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

Changes to Section H Part II – effective 1 August 2015 (continued)

130	FLUOROURACIL (delist) Inj 25 mg per ml, 100 ml vialInj 50 mg per ml, 10 ml vial Note – Hospira inj 25 mg per ml, 100 ml vial and Fluorouracil from 1 October 2015.	26.25	1 5) mg per ml,	Hospira Fluorouracil Ebewe 10 ml vial to be delisted
139	MEGESTROL ACETATE († price and addition of HSS) Tab 160 mg – 1% DV Oct-15 to 2018	54.30	30	Apo-Megestrol
141	TACROLIMUS (amended restriction) → Cap 0.5 mg – 1% DV Nov-14 to 31 Oct 2018 → Cap 1 mg – 1% DV Nov-14 to 31 Oct 2018 → Cap 5 mg – 1% DV Nov-14 to 31 Oct 2018 → Inj 5 mg per ml, 1 ml ampoule	.171.20	100 100 50	Tacrolimus Sandoz Tacrolimus Sandoz Tacrolimus Sandoz

Initiation – organ transplant recipients

For use in organ transplant recipients

Initiation - Steroid-resistant nephrotic syndrome*

Fither

Restricted

- 1 The patient is a child with steroid-resistant nephrotic syndrome* (SRNS) where ciclosporin has been trialled in combination with prednisone and discontinued because of unacceptable side effects or inadequate clinical response; or
- 2 All of the following:
 - 2.1 The patient is an adult with SRNS; and
 - 2.2 Ciclosporin has been trialled in combination with prednisone and discontinued because of unacceptable side effects or inadequate clinical response; and
 - 2.3 Cyclophosphamide or mycophenolate have been trialled and discontinued because of unacceptable side effects or inadequate clinical response, or these treatments are contraindicated.

Note: Indications marked with * are Unapproved Indications

RESPIRATORY SYSTEM AND ALLERGIES

- 171 BFF VFNOM
 - → Ini 120 mcg vial with diluent, 6 vial

Note – Bee venom inj 120 mcg vial with diluent, 6 vial to be delisted from 1 October 2015.

VACCINES

209 DIPHTHERIA, TETANUS AND PERTUSSIS VACCINE (amended restriction)

1 Boostrix 10 Boostrix

Restricted

Funded for any of the following:

- 1 A single vaccine for pregnant woman between gestational weeks 28 and 38 during epidemics; or
- 2 A course of up to four vaccines is funded for children from age 7 up the age of 18 years inclusive to complete full primary immunisation: or

continued...

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

Changes to Section H Part II – effective 1 August 2015 (continued)

continued...

3 An additional four doses (as appropriate) are funded for (re-)immunisation for patients post haematopoietic stem cell transplantation or chemotherapy; pre or post splenectomy; pre- or post solid organ transplant, renal dialvsis and other severely immunosuppressive regimens

Note: Tdap is not registered for patients aged less than 10 years. Please refer to the Immunisation Handbook for the appropriate schedule for catch up programmes.

212 INFLUENZA VACCINE

Restricted

Any of the following:

- 1 All people 65 years of age and over; or
- 2 People under 65 years of age who:
 - 2.1 Have any of the following cardiovascular diseases:
 - 2.1.1 Ischaemic heart disease; or
 - 2.1.2 Congestive heart failure; or
 - 2.1.3 Rheumatic heart disease; or
 - 2.1.4 Congenital heart disease; or 2.1.5 Cerebro-vascular disease; or
 - 2.2 Have any of the following chronic respiratory diseases:
 - 2.2.1 Asthma, if on a regular preventative therapy; or
 - 2.2.2 Other chronic respiratory disease with impaired lung function; or
 - 2.3 Have diabetes; or
 - 2.4 Have chronic renal disease: or
 - 2.5 Have any cancer, excluding basal and squamous skin cancers if not invasive; or
 - 2.6 Have any of the following other conditions:
 - 2.6.1 Autoimmune disease: or
 - 2.6.2 Immune suppression or immune deficiency; or
 - 2.6.3 HIV: or
 - 2.6.4 Transplant recipients: or
 - 2.6.5 Neuromuscular and CNS diseases/ disorders; or
 - 2.6.6 Haemoglobinopathies; or
 - 2.6.7 Are children on long term aspirin; or
 - 2.6.8 Have a cochlear implant; or
 - 2.6.9 Errors of metabolism at risk of major metabolic **decompensation** decomposition; or
 - 2.6.10 Pre and post splenectomy; or
 - 2.6.11 Down syndrome: or
 - 2.7 Are pregnant, or
 - 2.8 Are children aged four and under who have been hospitalised for respiratory illness or have a history of significant respiratory illness: or
- 3 Patients who are compulsorily detained long-term in a forensic unit within a DHB hospital in the 2015 season. Note: The following conditions are excluded from funding:
- asthma not requiring regular preventative therapy; and
- $\bullet \quad \hbox{hypertension and/or dyslipidaemia without evidence of end-organ disease.} \\$

Index

Pharmaceuticals and brands

A		Cefepime	. 20
Abilify 12, 23	3	Cefepime-AFT	. 20
Acarbose 18	8	Cefoxitin	7
Accuretic 10 18		Cefoxitin Actavis	
Accuretic 20 18		Cetomacrogol	
Aciclovir		Cisplatin	
Aciclovir-Claris	7	Citalopram hydrobromide	8
Aclasta 2	0	Clinicians Multivit & Mineral Boost	. 10
Actinomycin D 24	4	Clinicians Renal Vit	. 11
Adenuric 2	2	Colifoam	. 18
Advate 1:	5	Cosmegen	. 24
Alprostadil hydrochloride 19	9	Creon 10000	. 18
Aluminium chloride 1	1	Creon 25000	. 18
Amino acid formula (without phenylalanine)	9	Cyclizine hydrochloride	8
Amino acid formula (without phenylalanine		Cyclophosphamide	. 24
and tyrosine)		Cyproterone acetate	
Apo-Diclo SR		D	
Apo-Folic Acid	8	Dactinomycin	. 24
Apo-Megestrol		Dantrium IV	
Apo-Mirtazapine	7	Dantrolene	. 16
Apo-Moclobemide	3	DBL Bleomycin Sulfate	. 24
Apo-Nadolol1	9	DBL Cisplatin	. 17
Aqueous cream		DBL Leucovorin Calcium	
Aripiprazole		DBL Tobramycin	. 16
Arrow-Dortim1		Depo-Medrol	
Arrow-Sertraline	8	Depo-Medrol with Lidocaine	. 20
Atracurium besylate		Dexamethasone	
Azithromycin		Dexamfetamine sulfate	. 13
В		Dexmethsone	7
Balanced Salt Solution	9	Diclofenac Sandoz	. 12
Bee venom	5	Diclofenac sodium	. 12
BeneFIX 6, 1	5	Dimethicone	. 15
Benzbromaron AL 100 23	2	Diphtheria, tetanus and pertussis vaccine	. 25
Benzbromarone		Diprivan	
Bezafibrate 19	9	Dobutamine-Claris	. 11
Bezalip1	9	Dobutamine hydrochloride	. 11
Bezalip Retard 1	9	Domperidone	. 12
Bisacodyl	8	Dorzolamide with timolol	. 13
Bleomycin sulphate 24	4	Dulcolax	. 10
Blood ketone diagnostic test meter	9	E	
Boostrix 2	5	e-chamber La Grande	. 10
Bosentan 1	9	e-chamber Mask	. 10
Brilinta 1:		e-chamber Turbo	. 10
Bupivacaine hydrochloride 13	2	Elocon	. 16
Busulfan	8	Elocon Alcohol Free	. 16
C		Enbrel	. 13
Calamine1	1	Endoxan	. 24
Calcium chloride with magnesium chloride,		Epirubicin Ebewe	. 17
potassium chloride, sodium acetate, sodium		Epirubicin hydrochloride	
chloride and sodium citrate	9	Epoprostenol	. 15
Calcium folinate 1	7	Eptacog alfa	. 14

Index

Pharmaceuticals and brands

Ethics Lisinopril. 6 Lisinopril 6 CA CART STATE	Etanercept	13	Lindane	19
Factor eight inhibitor bypassing fraction. 6	Ethics Lisinopril	6		
Factor eight inhibitors bypassing agent 14 Marcain 12 Factor eight inhibitors bypassing fraction 6 11 14 Mask for spacer device 11 12 13 Medicol 15 Medicol 15 Medicol 15 Medicol 15 Medicol 15 Medicol 16 Meticol	F		LMX4	23
Factor eight inhibitors bypassing fraction	Factor eight inhibitor bypassing fraction	6	M	
Factor eight inhibitors bypassing fraction	Factor eight inhibitors bypassing agent	14	Marcain	12
Febusosiat 22 Medrol 11 15 15 15 15 15 15 1		14	Mask for spacer device	10
FEIBA 6, 11, 14 Megestrol acetate. 22	· · · · · · · · · · · · · · · · · · ·		•	19
FEIBA NF				
Finasteride				
Fingro				
Flecainide acetate				
Floair	•		·	
Fluarix			,	
Flucloxacillin				
Flucioxim				
Fluorouracil 24, 25				
Fluticasone 13			71 \	
Fluticasone 13	•			
Fluticasone with salmeterol				
Folic acid				
Freestyle Optium Neo 9 Mirtazapine 7, 15 G Mixed salt solution for eye irrigation 9 Gabapentin 8, 12 Moclobemide 25 Gacet 12 Mogine 15 Gamma benzene hexachloride 19 Mornetasone furoate 16 Glucobay 18 Moroctocog alfa 14 H Morphine hydrochloride 25 healthE Dimethicone 10% 15 Multivitamin and mineral supplement 16 Heparin sodium 15 Multivitamin renal 17 Hydrocortisone acetate 18 Mylan-Bosentan 15 Hydrogen peroxide 15 Multivitamin renal 17 Hydrogen peroxide 15 Myleran 8 I N I N Idarubicin hydrochloride 17 Nadolol 15 Influenza vaccine 26 Nauzene 26 Influenza vaccine 26 Neurontin 26 Isotane 10 16 Nevirapine Alphapharm			9	
G Mixed salt solution for eye irrigation S. G. Gabapentin S. 12 Moclobernide S. G. Gacet S. Moclobernide S. G. Gacet S. Moclobernide S. G. Gacet S. G. Gacet Moclobernide S. G. Gacet S. G. Gacet Moclobernide S. G. Gacet S. G. Gacet Moclobernide Moclobernice Moclobernide Moclobernice Moclobernice Moclobernice Moclobernice Moclobernic				
Gabapentin 8, 12 Moclobemide 23 Gacet 12 Mogine 15 Gamma benzene hexachloride 19 Mometasone furoate 16 Glucobay 18 Moroctocog alfa 14 H Morphine hydrochloride 23 healthE Dimethicone 10% 15 Multivitamin and mineral supplement 10 Heparin sodium 15 Multivitamin renal 11 Hydrocortisone acetate 18 Mylan-Bosentan 16 Hydrogen peroxide 15 Multivitamin renal 17 I N N N Idarubicin hydrochloride 17 Nadolol 19 Influenza vaccine 26 Nauzene 8 Influenza vaccine 26 Nauzene 8 Influenza vaccine 26 Neurontin 8 Isotane 10 16 Nevirapine Alphapharm 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotane FS 15 Norethisterone		9		
Gacet 12 Mogine 12 Gamma benzene hexachloride 19 Mometasone furoate 16 Glucobay 18 Moroctocog alfa 12 H Morphine hydrochloride 23 healthE Dimethicone 10% 15 Multivitamin and mineral supplement 16 Heparin sodium 15 Multivitamin renal 11 Hydrocortisone acetate 18 Mylan-Bosentan 15 Hydrogen peroxide 15 Myleran 8 I N I I Idarubicin hydrochloride 17 Nadolol 19 Influenza vaccine 26 Nauzene 16 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 22 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L		10		
Gamma benzene hexachloride 19 Mometasone furoate 16 Glucobay 18 Moroctocog alfa 1- H Morphine hydrochloride 25 healthE Dimethicone 10% 15 Multivitamin and mineral supplement 10 Heparin sodium 15 Multivitamin renal 1- Hydrocortisone acetate 18 Mylan-Bosentan 15 Hydrogen peroxide 15 Myleran 8 I N N I Idarubicin hydrochloride 17 Nadolol 15 Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11,16 Nicotine 22 K Nonacog alfa 6,15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 <td>·</td> <td></td> <td></td> <td></td>	·			
Section 18				
H Morphine hydrochloride. 23 healthE Dimethicone 10% 15 Multivitamin and mineral supplement 10 Heparin sodium 15 Multivitamin renal 11 Hydrocortisone acetate 18 Mylan-Bosentan 19 Hydrogen peroxide 15 Myleran 8 I N N I Idarubicin hydrochloride 17 Nadolol 15 Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 22 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 19 L Noriday 28 19 Lamotrigine 12 NovoSeven RT 14 Lanzol Relief 6 Octocog alfa 15 Lax- Suppositories 6				
healthE Dimethicone 10% 15 Multivitamin and mineral supplement 10 Heparin sodium 15 Multivitamin renal 11 Hydrocortisone acetate 18 Mylan-Bosentan 15 Hydrogen peroxide 15 Myleran 6 I N N Idarubicin hydrochloride 17 Nadolol 15 Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 11 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lanso prazole 6 O Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed		18		
Heparin sodium 15 Multivitamin renal 1 Hydrocortisone acetate 18 Mylan-Bosentan 15 Hydrogen peroxide 15 Myleran 8 I N I Idarubicin hydrochloride 17 Nadolol 16 Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 16 Lamotrigine 12 NovoSeven RT 12 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrozole 8 Oxandroline 12	==	4-	Morphine nydrochioride	23
Hydrocortisone acetate 18 Mylan-Bosentan 19 Hydrogen peroxide 15 Myleran 8 I N N Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 12 Lanzol Relief 6 O O Lanz-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrozole 8 Oxandroline 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Ligocaine 20, 23 Oxylorm 12, 23				
Hydrogen peroxide 15 Myleran 8 I N N Influenza vaccine 26 Nauzene 8 Influenza vaccine 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 12 Lanzol Relief 6 O O Lanz-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Ligonaine 20, 23 Oxylorm 12, 23	•			
Nadolol				
Idarubicin hydrochloride 17 Nadolol 19 Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 22 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Ligoncaine 20, 23 OxyNorm 12, 23 <	Hydrogen peroxide	15		8
Influenza vaccine 26 Nauzene 8 Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 2e K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Ligonocaine 20, 23 OxyNorm 12, 23	Idaruhicin hydrochloride	17	**	10
Influvac 26 Neurontin 8 Isotane 10 16 Nevirapine 16 Isotane 20 16 Nevirapine Alphapharm 16 Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				
Isotane 10				
Isotane 20. 16 Nevirapine Alphapharm 16 Isotretinoin. 11, 16 Nicotine. 24 K Nonacog alfa. 6, 15 Kogenate FS. 15 Norethisterone. 15 L Noriday 28. 15 Lamotrigine. 12 NovoSeven RT. 14 Lansoprazole. 6 0 Lanzol Relief. 6 Octocog alfa. 15 Lax-Suppositories. 6 Oral feed. 14 Lax-Tabs. 18 Oratane. 11 Letrole. 8 Oxandroline. 12 Letrozole. 8 Oxandrolone. 15 Lidocaine. 23 Oxycodone hydrochloride. 12, 23 Lignocaine. 20, 23 OxyNorm. 12, 23				
Isotretinoin 11, 16 Nicotine 24 K Nonacog alfa 6, 15 Kogenate FS 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23			•	
K Nonacog alfa 6, 15 Kogenate FS. 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				
Kogenate FS. 15 Norethisterone 15 L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 15 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23	v	10		
L Noriday 28 15 Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 11 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 15 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23	Noganata ES	15	,	
Lamotrigine 12 NovoSeven RT 14 Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23	I	13		
Lansoprazole 6 0 Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23	L ametrigine	10		
Lanzol Relief 6 Octocog alfa 15 Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				14
Lax-Suppositories 6 Oral feed 14 Lax-Tabs 18 Oratane 17 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23	•			46
Lax-Tabs 18 Oratane 1 Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 Oxylorm 12, 23				
Letrole 8 Oxandroline 12 Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				
Letrozole 8 Oxandrolone 12 Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				
Lidocaine 23 Oxycodone hydrochloride 12, 23 Lignocaine 20, 23 OxyNorm 12, 23				
Lignocaine				
	Lignocaine	23	UXYINOIM 12,	23

Index

Pharmaceuticals and brands

Oxytocin	16	Sertraline
Oxytocin BNM	16	Sodium a
P		Sodium o
Pancreatic enzyme	18	Solu-Med
Paracare	17	Spacer d
Paracetamol	23	Sustagen
Paragesic Soluble	23	Sustagen
Peak flow meter	10	T
Pethidine hydrochloride	17	Tacrolim
Pharmacy Health SLS-free	6	Tacrolim
Phenobarbitone	12	Tamboco
Phenytoin sodium	23	Tazocin E
Pioglitazone	10	Thiamine
Piperacillin with tazobactam	16	Thiotepa.
Pizaccord	10	Ticagrelo
Potassium dihydrogen phosphate	18	TMP
Prednisolone sodium phosphate	13	Tobramy
Procur	19	Tracrium
Prokinex	12	Trimetho
Propofol	17	V
Prostin VR	19	Veletri
PSM Citalopram	8	Vexazone
Q		Voricona
Quetapel	24	Vttack
Quetiapine	24	Χ
Quinapril with hydrochlorothiazide	18	Xyntha
R		Z
RA-Morph	23	Zavedos.
Recombinant factor IX	15	Ziprasido
Recombinant factor VIIA	14	Zithroma
Recombinant factor VIII	15	Zoledroni
RexAir	13	Zopiclone
\$		Zopiclone
SalAir	13	Zuclopen
Salbutamol	13	Zusdone.
Salmeterol	13	

Sertraline	8
Sodium acid phosphate	18
Sodium dihydrogen phosphate	18
Solu-Medrol	19
Spacer device	10
Sustagen Hospital Formula (Chocolate)	14
Sustagen Hospital Formula (Vanilla)	14
T	•
Tacrolimus	25
Tacrolimus Sandoz	25
Tambocor	11
Tazocin EF	16
Thiamine hydrochloride	6
Thiotepa	17
Ticagrelor	15
TMP	20
Tobramycin	16
Tracrium	
Trimethoprim	20
V	
Veletri	15
Vexazone	10
Voriconazole	
Vttack	
X	-
Xvntha	14
Z	-
Zavedos	17
Ziprasidone	
Zithromax	20
Zoledronic acid	20
Zopiclone	13
Zopiclone Actavis	13
Zuclopenthixol decanoate	17
Zusdone	
LUJUUIIU	

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