

Pharmaceutical Management Agency

Update

New Zealand Pharmaceutical Schedule

Effective 1 July 2013

Cumulative for May, June and July 2013



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Summary of PHARMAC decisions

EFFECTIVE 1 JULY 2013

New listings (page 22)

- Ascorbic acid (Cvite) tab 100 mg
- Vitamin B Complex (Bplex) tab, strong, BPC
- Vitamins (Mvite) tab (BPC cap strength)
- Ticagrelor (Brilinta) tab 90 mg – Special Authority – Retail pharmacy
- Pegfilgrastim (Neulastim) inj 6 mg per 0.6 ml syringe – Special Authority – Retail pharmacy
- Amiloride hydrochloride (Apo-Amiloride) tab 5 mg
- Cetomacrogol with glycerol (Pharmacy Health Sorbolene with Glycerine) crm 90% with glycerol 10%, 500 g OP
- Lidocaine [lignocaine] hydrochloride (Lidocaine-Claris) inj 1%, 5 ml and 20 ml ampoule
- Phenobarbitone (Martindale) inj 200 mg per ml, 1 ml ampoule – Special Authority – Retail pharmacy – S29
- Adalimumab (Humira) inj 20 mg per 0.4 ml prefilled syringe – Special Authority – Retail pharmacy
- Sodium hyaluronate (Hilo-Fresh) eye drops 1 mg per ml, 10 ml OP – Special Authority – Retail pharmacy – addition of note
- Retinol palmitate (VitA-POS) eye oint 138 mcg per g, 5 g OP – Special Authority – Retail pharmacy
- Pharmacy services (BSF Arrow-Quinapril) Brand switch fee – may only be claimed once per patient
- Fat supplement (Liquigen) oil, 250 ml, 4 OP – Special Authority – Hospital pharmacy [HP3]
- Renal oral feed 2 kcal/ml (Renilon 7.5) liquid (apricot), 125 ml, 4 OP and liquid (caramel), 125 ml, 4 OP – Special Authority – Hospital pharmacy [HP3]

Changes to restrictions (pages 25-45)

- Blood ketone diagnostic test meter (Freestyle Optium) – addition of 1 dev available on a PSO
- Ketone blood beta-ketone electrodes (Freestyle Optium Ketone) – addition of up to 10 test available on a PSO
- Blood glucose diagnostic test meter (CareSens) – addition of 1 dev available on a PSO, removal of Brand switch fee payable and addition of patient co-payment.
- Blood glucose diagnostic test strip (CareSens) – addition of up to 50 test available on a PSO
- Ursodeoxycholic acid (Ursosan) – Special Authority amendment

Summary of PHARMAC decisions – effective 1 July 2013 (continued)

- Mucilaginous laxatives (Konsyl-D) – amendment of chemical name to ispaghula (psyllium) husk
 - Protamine sulphate (Artex) inj 10 mg per ml, 5 ml – addition of S29
 - Quinapril (Arrow-Quinapril 5, Arrow-Quinapril 10, Arrow-Quinapril 20) tab 5 mg , 10 mg and 20 mg – addition of Brand switch fee payable
 - Cyproterone acetate with ethinyloestradiol (Ginet 84) – addition of up to 84 tab available on a PSO
 - Solifenacin succinate (Vesicare) tab 5 mg and 10 mg – Special Authority amendment
 - Tolterodine (Arrow-Tolterodine) tab 1 mg and 2 mg – Special Authority amendment
 - Prednisone (Apo-Prednisone) tab 1 mg – removal of stat symbol
 - Propylthiouracil (PTU) – addition of prescribing note
 - Cabergoline (Dostinex) tab 0.5 mg – Special Authority amendment
 - Danazol (Azol) cap 100 mg and 200 mg – removal of Retail pharmacy-Specialist
 - Cefazolin sodium (AFT) inj 500 mg and 1 g – amendment of endorsement restriction
 - Ceftriaxone sodium (Vercacol) inj 500 mg and 1 g (Aspen Ceftriazone) – amendment of endorsement restriction
 - Minocycline hydrochloride tab 50 mg – addition of Special Authority for higher subsidy
 - Gentamicin sulphate inj – amendment of endorsement restriction
 - Moxifloxacin (Avelox) tab 400 mg – Special Authority amendment
 - Vancomycin hydrochloride (Mylan) inj 500 mg – amendment of endorsement restriction
 - Fluconazole (Diflucan) powder for oral suspension 10 mg per ml – Special Authority amendment
 - Lamivudine (Zetlam) tab 100 mg and (Zeffix) oral liq 5 mg per ml – Special Authority amendment
 - Entecavir (Baraclude) tab 0.5 mg – Special Authority amendment
 - Tenofovir disoproxil fumarate (Viread) tab 300 mg – Special Authority amendment
 - Valaciclovir (Valtrex) tab 500 mg – Special Authority amendment
 - Antiretrovirals – Special Authority amendment
 - Guidelines for the use of interferon in the treatment of hepatitis C – amendment
 - Pegylated interferon alpha-2A – Special Authority amendment
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Summary of PHARMAC decisions – effective 1 July 2013 (continued)

- Lidocaine [lignocaine] hydrochloride inj 1%, 5 ml ampoule – amendment to PSO quantity
- Sumatriptan (Arrow-Sumatriptan) inj 12 mg per ml, 0.5 ml – addition of cartridge to presentation
- Hyoscine (Scopolamine) patch 1.5 mg – Special Authority amendment
- Adalimumab (Humira and HumiraPen) – Special Authority amendment
- Bee venom allergy treatment – Special Authority amendment
- Wasp venom allergy treatment – Special Authority amendment
- Special Foods – Special Authority amendments to
 - o Carbohydrate
 - o Carbohydrate and Fat
 - o Fat
 - o Protein
 - o Respiratory products
 - o Fat modified products
 - o High protein products
 - o Paediatric products
 - o Paediatric products for children with chronic renal failure
 - o Renal products
 - o Specialised and elemental products
 - o Adult products high calorie
 - o Extensively hydrolysed formula
- Renal oral feed 1 kcal/ml (Suplena) liquid – chemical amended to 2 kcal/ml

Decreased subsidy (pages 46-49)

- Insulin lispro with insulin lispro protamine (Humalog Mix 25) and (Humalog Mix 50)
 - Ispaghula (psyllium) husk (Konsyl-D) powder for oral soln
 - Sodium citrate with sodium lauryl sulphoacetate (Micolette) enema 90 mg with sodium lauryl sulphoacetate 9 mg per ml, 5 ml
 - Terazosin (Arrow) tab 1 mg, 2 mg and 5 mg
 - Cilazapril (Zapril) tab 0.5 mg, 2.5 mg and 5 mg
 - Clonidine hydrochloride (Dixarit) tab 25 mcg
 - Spironolactone (Spirotone) tab 25 mg and 100 mg
 - Ezetimibe (Ezetrol) tab 10 mg
 - Ezetimibe with simvastatin (Vytorin) tab 10 mg with simvastatin 10 mg, 10 mg with simvastatin 20 mg, 10 mg with simvastatin 40 mg and 10 mg with simvastatin 80 mg
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Summary of PHARMAC decisions – effective 1 July 2013 (continued)

- Levonorgestrel (Next Choice) tab 750 mcg
- Medroxyprogesterone acetate (Depo-Provera) inj 150 mg per ml, 1 ml syringe
- Clindamycin (Dalacin C) inj phosphate 150 mg per ml, 4 ml
- Aciclovir (Lovir) tab dispersible 200 mg, 400 mg and 800 mg
- Tetrabenazine (Motetis) tab 25 mg
- Lidocaine [lignocaine] hydrochloride (Xylocaine) inj 2%, 5 ml and 20 ml
- Dihydrocodeine tartrate (DHC Continus) tab long-acting 60 mg
- Morphine sulphate (Arrow-Morphine LA) tab long-acting 10 mg, 30 mg, 60 mg and 100 mg
- Sertraline (Arrow-Sertraline) tab 50 mg and 100 mg
- Venlafaxine (Arrow-Venlafaxine XR) tab 37.5 mg, 75 mg, 150 mg and 225 mg
- Venlafaxine (Efexor XR) cap 37.5 mg, 75 mg and 150 mg
- Sumatriptan (Arrow-Sumatriptan) tab 50 mg and 100 mg; and inj 12 mg per ml, 0.5 ml cartridge
- Naltrexone hydrochloride (Naltraccord) tab 50 mg
- Methotrexate (Hospira) inj 25 mg per ml, 2 ml and 20 ml
- Docetaxel (Taxotere) inj 20 mg per ml, 1 ml and 4 ml
- Docetaxel (Baxter) inj 1 mg for ECP
- Temozolomide (Temaccord) cap 5 mg, 20 mg, 100 mg and 250 mg
- Vincristine sulphate (Hospira) inj 1 mg per ml, 1ml and 2 ml
- Vincristine sulphate (Baxter) inj 1 mg for ECP
- Bacillus calmette-guerin (BCG) vaccine (OncoTICE) inj 2-8 x 100 million CFU
- Ipratropium bromide (Univent) nebuliser soln, 250 mcg per ml, 1 ml and 2 ml

Increased subsidy (pages 46-49)

- Fusidic acid (Foban) oint 2%, 15 g OP
- Morphine tartrate (Hospira) inj 80 mg per ml, 1.5 ml and 5 ml
- Nitrazepam (Nitrados) tab 5 mg

The new Hospital Medicines List (HML)

This month DHB hospitals start a transition to using a nationally-consistent list of funded hospital pharmaceuticals. The new Section H will be known as the HML (Hospital Medicines List) and will list all the medicines that all DHB hospitals need to provide for their patients. It includes both contracted products and non-contracted products.

The introduction of the HML will mean more alignment between the funding of medicines in hospitals and in the community.

Restrictions to reflect community restrictions

The use of some medicines in DHB hospitals will be restricted by prescriber type or indication. Each restriction will generally reflect current community restrictions such as Special Authorities.

Dispensing DHB prescriptions in the community

For community based patients' prescriptions originating from DHB hospitals, community pharmacies should dispense in accordance with the community Pharmaceutical Schedule listings.

A hospital pharmacy must dispense according to HML rules, including when dispensing to community-based patients.

Implementation/transition period

There are likely to be new listings and changes to restrictions in the community Pharmaceutical Schedule as work continues developing the HML and on aligning it with the community Schedule.



Further HML information can be found on our website, including:

- An electronic copy of the HML
- Notifications by therapeutic group of what medicines are included in the HML
- An alphabetical list of products that were considered for the HML

If you have questions or feedback relating to the HML, email HML@pharmac.govt.nz or call 0800 66 00 50 (option 2)

Please note that the changes to Section H that usually appear in this Update, are being reflected in a separate HML Update for the next few months.

Anti-Infectives restriction amendments

From 1 July 2013 there will be amendments to Special Authority criteria and endorsement criteria for a number of pharmaceuticals listed in the Infections Group. The changes are the result of a review of the pharmaceuticals in this group by the Anti-infective Subcommittee.

Sensory Organs changes

There will be a number of changes to the Sensory therapeutic group of products. These include:

- Hylo-Fresh (sodium hyaluronate) eye drops 1 mg per ml will be listed fully funded and Sole Supply from 1 July 2013. Hylo-Fresh will be fully funded for patients with severe secretory dry eye who have a confirmed allergic reaction to eye drop preservative. Note that Hylo-Fresh eye drops have a **six month expiry after opening**. Only the prescribed dosage to the nearest OP may be claimed. Prescriptions will not be subsidised as one bottle per month as detailed in the Pharmacy Procedures Manual.
 - VitA-POS (retinol palmitate) eye oint 138 mcg per g, 5 g OP will be listed fully funded from 1 July 2013.
 - Systane Unit Dose (macrogol 400 0.4% with propylene glycol 0.3%) 0.4 ml eye drops will be listed fully funded and Sole Supply from 1 August 2013.
 - Poly-Gel (carbomer) 0.3% ophthalmic gel, 0.5 g will be listed fully funded and Sole Supply from 1 August 2013.
 - Special Authority criteria will apply to all Preservative Free Ocular Lubricants from 1 July 2013.
 - The listing for Poly-Tears (hypromellose 0.3% with dextran 0.1%) eye drops will be amended from 1 August 2013 to include dextran.
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Funding of blood glucose and blood ketone testing products via PSO

From 1 July 2013 the following quantities of blood glucose and blood ketone testing products will be funded on a PSO (in addition to the current subsidy provisions):

- Blood glucose diagnostic test meter – 1 meter
- Blood glucose diagnostic test strip – up to 50 strips (1 OP) of **CareSens or CareSens N brands only**
- Blood ketone diagnostic test meter – 1 meter
- Ketone blood beta-ketone electrodes test strip – up to 10 strips (1 OP)

Ticagrelor subsidised for acute coronary syndrome

Ticagrelor (Brilinta) 90 mg tablets will be listed fully funded from 1 July 2013 subject to Special Authority criteria for acute coronary syndrome.

Pegfilgrastim – new listing

From 1 July 2013, pegfilgrastim (Neulastim) injection, 6 mg per 0.6 ml syringe, will be listed fully funded subject to Special Authority criteria for the prevention of neutropenia in patients undergoing high risk chemotherapy for cancer

Risedronate – new listing

From 1 September 2013, risedronate sodium (Risedronate Sandoz) 35 mg tablets will be listed fully funded.

Adalimumab new listing and extension to Special Authority criteria

From 1 July 2013, a 20 mg per 0.4 ml prefilled syringe of adalimumab (Humira) will be subsidised. The Special Authority criteria will also be amended from 1 July 2013 to include patients with juvenile idiopathic arthritis and fistulising Crohn's disease.

Phenobarbitone injection subsidised

Phenobarbitone (Martindale) injection 200 mg per ml, 1 ml ampoule will be listed fully funded from 1 July 2013 subject to Special Authority criteria for the treatment of terminal agitation unresponsive to other agents. The Martindale brand is an unapproved medicine and must be supplied in accordance with section 29 of the Medicines Act 1981.

m-Eslon 30 mg capsules out-of-stock

Multichem has advised that m-Eslon (morphine sulphate) 30 mg long-acting capsules are out-of-stock until mid-July. Pharmacists can dispense the equivalent dose of 10 mg long-acting capsules or contact the prescriber.

Ursodeoxycholic acid – amendment to Special Authority criteria

The Special Authority criteria for ursodeoxycholic acid will be widened from 1 July 2013 to include patients with:

- Alagille syndrome
- Progressive familial intrahepatic cholestasis (PFIC)
- Chronic severe drug induced cholestatic liver injury; and
- Total parenteral nutrition induced cholestasis (TPN-IC) in paediatric patients

The criteria will also be amended to specify the criteria for patients with cholestasis associated with pregnancy and for patients with cirrhosis, and to reduce the bilirubin level relating to decompensated cirrhosis from 170 $\mu\text{mol/L}$ to 100 $\mu\text{mol/L}$.

Prednisone 1 mg tablets – removal of Stat dispensing

Due to limited supply of Apo-Prednisone 1 mg tablets, the requirement to dispense prednisone 1 mg tablets 'stat' will be removed from 1 July 2013 until further notice.

Protamine sulphate – addition of section 29

From 1 July 2013, the section 29 symbol will be added to the listing of the Artex brand of protamine sulphate. New supplies of stock are unapproved and must be supplied in accordance with section 29 of the Medicines Act 1981.

New Pharmacode for Zyban

From 1 August 2013, there is a price and subsidy reduction for bupropion hydrochloride (Zyban) modified release tablets. GSK have advised that there will be a new Pharmacode for the stock supplied at the new price and claims for the new Pharmacode cannot be made until it is listed from 1 August 2013.

Permax (pergolide) 0.25 mg stock shortage

Permax (Pergolide) 0.25 mg, supplied by Aspen Pharmacare, is temporarily out-of-stock due to manufacturing issues. Resupply is expected in September 2013 at the earliest. Some stock of Permax 1 mg is available, but this presentation is also expected to go out-of-stock. Patients should consult with their prescribers regarding alternative treatment options.

News in brief

- A Brand Switch Fee will apply to dispensings of **Arrow-Quinapril** from 1 July 2013 to 1 October 2013.
- **Lincocin (lincomycin) injection** 300 mg per ml, 2 ml injection will be delisted 1 December 2013.
- The description of Konsyl-D is changing from **mucilaginous laxatives** to ispaghula (psyllium) husk from 1 July 2013.
- Please note due to the number of changes this month only the amended portion of the **Special Authority** criteria is shown. For the full text refer to the Pharmaceutical Schedule online at www.pharmac.govt.nz.
- From 1 July 2013 pharmacist are no longer able to dispense and claim for **blood glucose diagnostic test meters** without a prescription.



Tender News

Sole Subsidised Supply changes – effective 1 August 2013

Chemical Name	Presentation; Pack size	Sole Subsidised Supply brand (and supplier)
Amisulpride	Oral liq 100 mg per ml; 60 ml	Solian (Sanofi)
Amisulpride	Tab 100 mg; 30 tab	Solian (Sanofi)
Amisulpride	Tab 200 mg; 60 tab	Solian (Sanofi)
Amisulpride	Tab 400 mg; 60 tab	Solian (Sanofi)
Codeine phosphate	Tab 15 mg; 100 tab	PSM (PSM)
Codeine phosphate	Tab 30 mg; 100 tab	PSM (PSM)
Codeine phosphate	Tab 60 mg; 100 tab	PSM (PSM)

Looking Forward

This section is designed to alert both pharmacists and prescribers to possible future changes to the Pharmaceutical Schedule. It may also assist pharmacists, distributors and wholesalers to manage stock levels.

Possible decisions for future implementation 1 August 2013

- Removal of Special Authority on Arrow-Venlafaxine brand of venlafaxine
- Increase in price and subsidy for Marevan brand of warfarin
- Oxydone BNM brand of oxycodone hydrochloride – new listing

Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Abacavir sulphate	Oral liq 20 mg per ml Tab 300 mg	Ziagen Ziagen	2014
Acarbose	Tab 50 mg and 100 mg	Accarb	2015
Acetazolamide	Tab 250 mg	Diamox	2014
Acetylcysteine	Inj 200 mg per ml, 10 ml	Martindale Acetylcysteine	2015
Allopurinol	Tab 100 mg & 300 mg	Apo-Allopurinol	2014
Amantadine hydrochloride	Cap 100 mg	Symmetrel	2014
Aminophylline	Inj 25 mg per ml, 10 ml	DBL Aminophylline	2014
Amitriptyline	Tab 10 mg Tab 25 mg & 50 mg	Arrow-Amitriptyline Amitrip	2014
Amlodipine	Tab 2.5 mg Tab 5 mg & 10 mg	Apo-Amlodipine Apo-Amlodipine	2014
Amoxicillin	Inj 250 mg, 500 mg & 1 g	Ibiamox	2014
Amoxicillin clavulanate	Grans for oral liq amoxicillin 125 mg with potassium clavulanate 31.25 mg per 5 ml	Augmentin	2015
	Grans for oral liq amoxicillin 250 mg with potassium clavulanate 62.5 mg per 5 ml	Augmentin	
	Tab 500 mg with potassium clavulanate 125 mg	Curam Duo	2014
Aqueous cream	Crm	AFT	2014
Atenolol	Tab 50 mg & 100 mg	Mylan Atenolol	2015
Atorvastatin	Tab 10 mg, 20 mg, 40 mg & 80 mg	Zarator	2015
Atropine sulphate	Inj 600 mcg, 1 ml	AstraZeneca	2015
Azithromycin	Tab 500 mg	Apo-Azithromycin	2015
Baclofen	Tab 10 mg	Pacifen	2016
Bendrofluazide	Tab 2.5 mg & 5 mg	Arrow- Bendrofluazide	2014
Benzathine benzylpenicillin	Inj 1.2 mega u per 2.3 ml	Bicillin LA	2015
Benzylpenicillin sodium (Penicillin G)	Inj 600 mg	Sandoz	2014
Betaxolol hydrochloride	Eye drops 0.5% Eye drops 0.25%	Betoptic Betoptic S	2014
Bezafibrate	Tab 200 mg Tab long-acting 400 mg	Bezalip Bezalip Retard	2015
Bicalutamide	Tab 50 mg	Bicalaccord	2014
Blood glucose diagnostic test meter	Meter with 50 lancets, a lancing device and 10 diagnostic test strips	CareSens N CareSens N POP CareSens II	2015

*Expiry date of the Sole Subsidised Supply period is 30 June of the year indicated unless otherwise stated. Please note that Sole Subsidised Supply may have been awarded for a wider scope than just those presentation(s) listed in the above table.

Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Blood glucose diagnostic test strip	Blood glucose test strips	CareSens CareSens N	2015
Brimonidine tartrate	Eye drops 0.2%	Arrow-Brimonidine	2014
Cabergoline	Tab 0.5 mg	Dostinex	2015
Calamine	Lotn, BP	PSM	2015
Calcitonin	Inj 100 iu per ml, 1 ml	Miacalcic	2014
Calcium carbonate	Tab 1.25 g (500 mg elemental) Tab eff 1.75 g (1 g elemental)	Arrow-Calcium Calsource	2014
Calcium folinate	Tab 15 mg	DBL Leucovorin Calcium	2014
Candesartan	Tab 4 mg, 8 mg, 16 mg & 32 mg	Candestar	2015
Cefazolin sodium	Inj 500 mg & 1 g	AFT	2014
Cefuroxime sodium	Inj 750 mg	Multichem	2014
Cetirizine hydrochloride	Oral liq 1 mg per ml Tab 10 mg	Cetirizine - AFT Zetop	2014
Chloramphenicol	Eye oint 1% Eye drops 0.5%	Chlorsig Chlorafast	2015
Chlorhexidine gluconate	Mouthwash 0.2% Handrub 1% with ethanol 70% Soln 4%	healthE healthE Orion	2015 2014
Ciclopirox olamine	Nail-soln 8%	Apo-Ciclopirox	2015
Ciprofloxacin	Tab 250 mg, 500 mg & 750 mg	Cipflox	2014
Citalopram hydrobromide	Tab 20 mg	Arrow-Citalopram	2014
Clarithromycin	Tab 500 mg Tab 250 mg	Apo-Clarithromycin Apo-Clarithromycin	2014
Clomipramine hydrochloride	Tab 10 mg & 25 mg	Apo-Clomipramine	2015
Clonidine hydrochloride	Tab 150 mcg Inj 150 mcg per ml, 1 ml	Catapres	2015
Clotrimazole	Crn 1%	Clomazol	2014
Crotamiton	Crn 10%	Itch-Soothe	2015
Cyclizine hydrochloride	Tab 50 mg	Nausicalm	2015
Cyclosporin	Oral liq 100 mg per ml	Neoral	2015
Cyproterone acetate	Tab 50 mg & 100 mg	Siterone	2015
Cyproterone acetate with ethinyloestradiol	Tab 2 mg with ethinyloestradiol 35 mcg and 7 inert tabs	Ginet 84	2014
Desmopressin	Nasal spray 10 mcg per dose	Desmopressin-PH&T	2014
Dexamethasone	Tab 1 mg & 4 mg Eye oint 0.1%	Douglas Maxidex	2015 2014

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Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Dexamethasone with neomycin and polymyxin b sulphate	Eye oint 0.1% with neomycin sulphate 0.35% and polymyxin B sulphate 6,000 u per g	Maxitrol	2014
	Eye drops 0.1% with neomycin sulphate 0.35% and polymyxin B sulphate 6,000 u per ml	Maxitrol	
Dexamphetamine sulphate	Tab 5 mg	PSM	2015
Dextrose	Inj 50%, 10 ml	Biomed	2014
Diclofenac sodium	Tab EC 25 mg & 50 mg	Apo-Diclo	2015
	Tab long-acting 75 mg & 100 mg	Diclax SR	
	Inj 25 mg per ml, 3 ml	Voltaren	2014
	Eye drops 1 mg per ml Suppos 12.5 mg, 25 mg, 50 mg & 100 mg	Voltaren Ophtha Voltaren	
Diltiazem hydrochloride	Cap long-acting 120 mg, 180 mg & 240 mg	Apo-Diltiazem CD	2015
	Tab 30 mg & 60 mg	Dilzem	
Dipyridamole	Tab long-acting 150 mg	Pytazen SR	2014
Docusate sodium	Cap 50 mg	Laxofast 50	2014
	Cap 120 mg	Laxofast 120	
Domperidone	Tab 10 mg	Prokinex	2015
Doxazosin mesylate	Tab 2 mg & 4 mg	Apo-Doxazosin	2014
Doxycycline hydrochloride	Tab 100 mg	Doxine	2014
Emulsifying ointment	Oint BP	AFT	2014
Enalapril	Tab 5 mg, 10 mg & 20 mg	m-Enalapril	2015
Enoxaparin sodium	Inj 20 mg, 40 mg, 60 mg, 80 mg, 100 mg, 120 mg & 150 mg	Clexane	2015
Entacapone	Tab 200 mg	Entapone	2015
Ergometrine maleate	Inj 500 mcg per ml, 1 ml	DBL Ergometrine	2014
Etidronate disodium	Tab 200 mg	Arrow-Etidronate	2015
Ethinylestradiol	Tab 10 mcg	NZ Medical and Scientific	2015
Ethinylestradiol with levonorgestrel	Tab 20 mcg with levonorgestrel 100 mcg & 7 inert tab	Ava 20 ED	2014
	Tab 30 mcg with levonorgestrel 150 mcg & 7 inert tab	Ava 30 ED	
Exemestane	Tab 25 mg	Aromasin	2014
Felodopine	Tab long-acting 5 mg & 10 mg	Plendil ER	2015
	Tab long-acting 2.5 mg	Plendil ER	
Fentanyl citrate	Inj 50 mcg per ml, 2 ml & 10 ml	Boucher and Muir	2015
Filgrastim	Inj 300 mcg per 0.5 ml	Zarzio	31/12/15
	Inj 480 mcg per 0.5 ml	Zarzio	

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Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Finasteride	Tab 5 mg	Rex Medical	2014
Flucloxacillin sodium	Grans for oral liq 125 mg per 5 ml Grans for oral liq 250 mg per 5 ml Cap 250 mg & 500 mg Inj 250 mg, 500 mg & 1 g	AFT Staphlex Flucloxin	2015 2014
Fluconazole	Cap 50 mg, 150 mg & 200 mg	Ozole	2014
Fluorometholone	Eye drops 0.1%	Flucon	2015
Fluorouracil sodium	Crn 5%	Efudix	2015
Fluticasone propionate	Metered aqueous nasal spray, 50 mcg per dose	Flixonase Hayfever & Allergy	2015
Furosemide	Tab 500 mg Tab 40 mg	Urex Forte Diurin 40	2015
Gentamicin sulphate	Inj 40 mg per ml, 2 ml	Pfizer	2015
Gliclazide	Tab 80 mg	Apo-Gliclazide	2014
Glipizide	Tab 5 mg	Minidiab	2015
Glycerol	Suppos 3.6 g	PSM	2015
Glyceryl trinitrate	Aerosol spray 400 mcg per dose TDDS 5 mg & 10 mg Tab 600 mcg	Glytrin Nitroderm TTS Lycinate	2014
Hydrocortisone	Tab 5 mg & 20 mg Crn 1% Powder	Douglas Pharmacy Health ABM	2015 2014
Hydrocortisone acetate	Rectal foam 10%, CFC-Free (14 applications)	Colifoam	2015
Hydrocortisone butyrate	Lipocream 0.1% Milky emul 0.1% Oint 0.1% Scalp lotn 0.1%	Locoid Lipocream Locoid Crelo Locoid Locoid	2015
Hydrocortisone with wool fat and mineral oil	Lotn 1% with wool fat hydrous 3% and mineral oil	DP Lotn HC	2014
Hydroxocobalamin	Inj 1 mg per ml, 1 ml	ABM Hydroxocobalamin	2015
Hydroxychloroquine sulphate	Tab 200 mg	Plaquenil	2015
Hyoscine N-butylbromide	Inj 20 mg, 1 ml Tab 10 mg	Buscopan Gastrosoothe	2014
Ibuprofen	Tab 200 mg Tab long-acting 800 mg	Arrowcare Brufen SR	2014
Imiquimod	Crn 5%	Aldara	2014
Iron polymaltose	Inj 50 mg per ml, 2 ml	Ferrum H	2014
Isoniazid	Tab 100 mg	PSM	2015

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Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Isosorbide mononitrate	Tab 20 mg Tab long-acting 40 mg	Ismo 20 Corangin	2014
Isotretinoin	Cap 10 mg & 20 mg	Oratane	2015
Ketoconazole	Shampoo 2%	Sebizole	2014
Lamivudine	Tab 100 mg	Zetlam	2014
Lansoprazole	Cap 15 mg & 30 mg	Solox	2015
Latanoprost	Eye drops 50 mcg per ml	Hysite	2015
Letrozole	Tab 2.5 mg	Letraccord	2015
Levonorgestrel	Subdermal implant (2 x 75 mg rods)	Jadelle	31/12/13
Lignocaine hydrochloride	Viscous soln 2%	Xylocaine Viscous	2014
Lisinopril	Tab 5 mg, 10 mg & 20 mg	Arrow-Lisinopril	2015
Lithium carbonate	Tab 250 mg & 400 mg Cap 250 mg	Lithicarb FC Douglas	2015 2014
Lodoxamide trometamol	Eye drops 0.1%	Lomide	2014
Losartan	Tab 12.5 mg, 25 mg, 50 mg & 100 mg	Lostaar	2014
Losartan with hydrochlorothiazide	Tab 50 mg with hydrochlorothiazide 12.5 mg	Arrow-Losartan & Hydrochlorothiazide	2014
Mask for spacer device	Size 2	EZ-fit Paediatric Mask	2015
Mebendazole	Tab 100 mg	De-Worm	2014
Mebeverine hydrochloride	Tab 135 mg	Colofac	2014
Megestrol acetate	Tab 160 mg	Apo-Megestrol	2015
Methylprednisolone	Tab 4 mg & 100 mg	Medrol	2015
Methylprednisolone acetate	Inj 40 mg per ml	Depo-Medrol	2015
Methylprednisolone acetate with lignocaine	Inj 40 mg per ml with lignocaine 1 ml	Depo-Medrol with Lidocaine	2015
Mesalazine	Enema 1 g per 100 ml Suppos 500 mg	Pentasa Asacol	2015 2014
Metformin hydrochloride	Tab immediate-release 500 mg & 850 mg	Apotex	2015
Methadone hydrochloride	Oral liq 2 mg per ml Oral liq 5 mg per ml Oral liq 10 mg per ml	Biodone Biodone Forte Biodone Extra Forte	2015
Methylprednisolone sodium succinate	Inj 40 mg per ml, 1 ml; 62.5 mg per ml, 2 ml; 500 mg & 1 g	Solu-Medrol	2015

*Expiry date of the Sole Subsidised Supply period is 30 June of the year indicated unless otherwise stated. Please note that Sole Subsidised Supply may have been awarded for a wider scope than just those presentation(s) listed in the above table.

Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Metoclopramide hydrochloride	Inj 5 mg per ml, 2 ml Tab 10 mg	Pfizer Metamide	2014
Metoprolol succinate	Tab long-acting 23.75 mg, 47.5 mg, 95 mg & 190 mg	Metoprolol-AFT CR	2015
Metoprolol tartrate	Inj 1 mg per ml, 5 ml Tab 50 mg & 100 mg Tab long-acting 200 mg	Lopresor Lopresor Slow-Lopresor	2015
Miconazole	Oral gel 20 mg per g	Decozol	2015
Miconazole nitrate	Crn 2%	Multichem	2014
Mirtazapine	Tab 30 mg & 45 mg	Avanza	2015
Moclobemide	Tab 150 mg & 300 mg	Apo-Moclobemide	2015
Mometasone furoate	Crn 0.1% Oint 0.1%	m-Mometasone	2015
Morphine hydrochloride	Oral liq 1 mg per ml, 2 mg per ml, 5 mg per ml & 10 mg per ml	RA-Morph	2015
Morphine sulphate	Inj 5 mg per ml, 1 ml Inj 10 mg per ml, 1 ml Inj 15 mg per ml, 1 ml Inj 30 mg per ml, 1 ml	DBL Morphine Sulphate DBL Morphine Sulphate DBL Morphine Sulphate DBL Morphine Sulphate	2014
Naphazoline hydrochloride	Eye drops 0.1%	Naphcon Forte	2014
Nadolol	Tab 40 mg & 80 mg	Apo-Nadolol	2015
Naproxen	Tab 250 mg Tab 500 mg	Noflam 250 Noflam 500	2015
Neostigmine	Inj 2.5 mg per ml, 1 ml	AstraZeneca	2014
Nevirapine	Tab 200 mg	Nevirapine Alphapharm	2015
Nicotine	Gum 2 mg & 4 mg (classic, fruit, mint) Lozenge 1 mg & 2 mg Patch 7 mg, 14 mg & 21 mg	Habitrol Habitrol Habitrol	2014
Nicotinic acid	Tab 50 mg & 500 mg	Apo-Nicotinic Acid	2014
Norethisterone	Tab 350 mcg Tab 5 mg	Noriday 28 Primolut N	2015 2014
Norfloxacin	Tab 400 mg	Arrow-Norfloxacin	2014
Nortriptyline hydrochloride	Tab 10 mg & 25 mg	Norpress	2016
Nystatin	Oral liq 100,000 u per ml	Nilstat	2014

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Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Octreotide (somatostatin analogue)	Inj 50 mcg per ml, 1 ml Inj 100 mcg per ml, 1 ml Inj 500 mcg per ml, 1 ml	Octreotide Max Rx	2014
Oil in water emulsion	Crn	healthE Fatty Cream	2015
Omeprazole	Cap 10 mg, 20 mg & 40 mg Powder Inj 40 mg	Omezol Relief Midwest Dr Reddy's Omeprazole	2014
Oxazepam	Tab 10 mg & 15 mg	Ox-Pam	2014
Oxybutynin	Oral liq 5 mg per ml Tab 5 mg	Apo-Oxybutynin	2016
Oxycodone hydrochloride	Inj 50 mg per ml, 1 ml Inj 10 mg per ml, 1 ml & 2 ml	OxyNorm Oxycodone Orion	2015
Oxytocin	Inj 5 iu with ergometrine maleate 500 mcg per ml, 1 ml	Syntometrine	2015
Pamidronate disodium	Inj 3 mg per ml, 10 ml; 6 mg per ml, 10 ml & 9 mg per ml, 10 ml	Pamidronate BNM	2014
Pantoprazole	Inj 40 mg	Pantocid IV	2014
Paracetamol	Suppos 500 mg Tab 500 mg Oral liq 120 mg per 5 ml Oral liq 250 mg per 5 ml	Paracare Parafast Ethics Paracetamol Paracare Double Strength	2015 2014
Paracetamol with codeine	Tab paracetamol 500 mg with codeine phosphate 8 mg	Paracetamol + Codeine (Relieve)	2014
Peak flow meter	Low range & normal range	Breath-Alert	2015
Pergolide	Tab 0.25 mg & 1 mg	Permax	2014
Permethrin	Crn 5% Lotn 5%	Lyderm A-Scabies	2014
Pethidine hydrochloride	Tab 50 mg & 100 mg Inj 50 mg per ml, 1 ml Inj 50 mg per ml, 2 ml	PSM DBL Pethidine Hydrochloride DBL Pethidine Hydrochloride	2015 2014
Phenobarbitone	Tab 15 mg & 30 mg	PSM	2015
Pioglitazone	Tab 15 mg, 30 mg & 45 mg	Pizaccord	2015
Pizotifen	Tab 500 mcg	Sandomigran	2015
Poloxamer	Oral drops 10%	Coloxyl	2014
Potassium chloride	Tab long-acting 600 mg	Span-K	2015
Pravastatin	Tab 20 mg & 40 mg	Cholvastin	2014
Procaine penicillin	Inj 1.5 mega u	Cilicaine	2014
Promethazine hydrochloride	Oral liq 5 mg per 5 ml Tab 10 mg & 25 mg	Allersoothe Allersoothe	2015

*Expiry date of the Sole Subsidised Supply period is 30 June of the year indicated unless otherwise stated. Please note that Sole Subsidised Supply may have been awarded for a wider scope than just those presentation(s) listed in the above table.

Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Pyridostigmine bromide	Tab 60 mg	Mestinon	2014
Pyridoxine hydrochloride	Tab 25 mg Tab 50 mg	PyridoxADE Apo-Pyridoxine	2014
Quinapril	Tab 5 mg, 10 mg & 20 mg	Arrow-Quinapril	2015
Quinapril with hydrochlorothiazide	Tab 10 mg with hydrochlorothiazide 12.5 mg Tab 20 mg with hydrochlorothiazide 12.5 mg	Accuretic 10 Accuretic 20	2015
Ranitidine hydrochloride	Oral liq 150 mg per 10 ml Tab 150 mg & 300 mg	Peptisoothe Arrow-Ranitidine	2014
Ritonavir	Tab 100 mg	Norvir	2015
Rizatriptan	Tab orodispersible 10 mg	Rizamelt	2014
Roxithromycin	Tab 150 mg & 300 mg	Arrow-Roxithromycin	2015
Salbutamol	Nebuliser soln, 1 mg per ml & 2 mg per ml, 2.5 ml	Asthalin	2015
Salbutamol with ipratropium bromide	Nebuliser soln, 2.5 mg with ipratropium bromide 0.5 mg per vial, 2.5 ml	Duolin	2015
Sildenafil	Tab 25 mg, 50 mg & 100 mg	Silagra	2014
Simvastatin	Tab 10 mg Tab 20 mg Tab 40 mg Tab 80 mg	Arrow-Simva 10mg Arrow-Simva 20mg Arrow-Simva 40mg Arrow-Simva 80mg	2014
Sodium hyaluronate	Eye drops 1 mg per ml, 10 ml OP	Hylo-Fresh	2016
Spacer device	800 ml 230 ml (single patient)	Volumatic Space Chamber Plus	2015
Tamoxifen citrate	Tab 20 mg	Genox	2014
Tar with triethanolamine lauryl sulphate and fluorescein	Soln 2.3% with triethanolamine lauryl sulphate and fluorescein sodium, 500 ml & 1,000 ml	Pinetarsol	2014
Temazepam	Tab 10 mg	Normison	2014
Terbinafine	Tab 250 mg	Dr Reddy's Terbinafine	2014
Testosterone cypionate	Inj long-acting 100 mg per ml, 10 ml	Depo-Testosterone	2014
Testosterone undecanoate	Cap 40 mg	Andriol Testocaps	2015
Tetracosactrin	Inj 250 mcg Inj 1 mg per ml, 1 ml	Synacthen Synacthen Depot	2014
Timolol maleate	Eye drops 0.25% & 0.5%	Arrow-Timolol	2014
Tobramycin	Eye drops 0.3% Eye oint 0.3% Inj 40 mg per ml, 2 ml	Tobrex Tobrex DBL Tobramycin	2014

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Sole Subsidised Supply Products – cumulative to July 2013

Generic Name	Presentation	Brand Name	Expiry Date*
Tolcapone	Tab 100 mg	Tasmar	2014
Tramadol hydrochloride	Cap 50 mg	Arrow-Tramadol	2014
Triamcinolone acetonide	Inj 10 mg per ml, 1 ml Inj 40 mg per ml, 1 ml Crm 0.02% Oint 0.02% 0.1% in Dental Paste USP	Kenacort-A Kenacort-A40 Aristocort Aristocort Oracort	2014
Tropicamide	Eye drops 0.5% & 1%	Mydracyl	2014
Ursodeoxycholic acid	Cap 250 mg	Ursosan	2014
Vancomycin hydrochloride	Inj 500 mg	Mylan	2014
Verapamil hydrochloride	Tab 40 mg & 80 mg	Isoptin	2014
Zidovudine [AZT] with lamivudine	Tab 300 mg with lamivudine 150 mg	Alphapharm	2014
Zinc and castor oil	Oint BP	Multichem	2014
Zinc sulphate	Caps 137.4 mg (50 mg elemental)	Zincaps	2014

July changes are in bold type

*Expiry date of the Sole Subsidised Supply period is 30 June of the year indicated unless otherwise stated. Please note that Sole Subsidised Supply may have been awarded for a wider scope than just those presentation(s) listed in the above table.

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

New Listings

Effective 1 July 2013

41	ASCORBIC ACID a) No more than 100 mg per dose b Only on a prescription * Tab 100 mg	13.80	500	✓ Cvite
41	VITAMIN B COMPLEX * Tab, strong, BPC	4.70	500	✓ Bplex
42	VITAMINS * Tab (BPC cap strength)	8.00	1,000	✓ Mvite
45	TICAGRELOR – Special Authority see SA1382 – Retail pharmacy * Tab 90 mg	90.00	56	✓ Brilinta
<p>▶ SA1382 Special Authority for Subsidy Initial application (acute coronary syndrome) from any relevant practitioner. Approvals valid for 12 months for applications meeting the following criteria: Both: 1 Patient has recently been diagnosed with an ST-elevation or a non-ST-elevation acute coronary syndrome; and 2 Fibrinolytic therapy has not been given in the last 24 hours and is not planned. Renewal (subsequent acute coronary syndrome) from any relevant practitioner. Approvals valid for 12 months for applications meeting the following criteria: Both: 1 Patient has recently been diagnosed with an ST-elevation or a non-ST-elevation acute coronary syndrome; and 2 Fibrinolytic therapy has not been given in the last 24 hours and is not planned.</p>				
48	PEGFILGRASTIM – Special Authority see SA1384 – Retail pharmacy Inj 6 mg per 0.6 ml syringe.....	1,080.00	1	✓ Neulastim
<p>▶ SA1384 Special Authority for Subsidy Initial application only from a relevant specialist, vocationally registered general practitioner or medical practitioner on the recommendation of a relevant specialist. Approvals valid without further renewal unless notified where used for prevention of neutropenia in patients undergoing high risk chemotherapy for cancer (febrile neutropenia risk \geq 20%*).* *Febrile neutropenia risk \geq 20% after taking into account other risk factors as defined by the European Organisation for Research and Treatment of Cancer (EORTC) guidelines.</p>				
58	AMILORIDE HYDROCHLORIDE * Tab 5 mg	17.50	100	✓ Apo-Amiloride
69	CETOMACROGOL WITH GLYCEROL Crm 90% with glycerol 10%.....	4.50	500 g OP	✓ Pharmacy Health Sorbolene with Glycerin
119	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE Inj 1%, 5 ml ampoule – Up to 25 inj available on a PSO	8.75	25	✓ Lidocaine-Claris
	Inj 1%, 20 ml ampoule – Up to 5 inj available on a PSO	2.40	1	✓ Lidocaine-Claris

Patients pay a manufacturer's surcharge when the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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New Listings - effective 1 July 2013 (continued)

140	PHENOBARBITONE SODIUM – Special Authority see SA1386 – Retail pharmacy Inj 200 mg per ml, 1 ml ampoule	46.20	10	✓ Martindale S29
	<p>▶ SA1386 Special Authority for Subsidy Initial application from any relevant practitioner. Approvals valid without further renewal unless notified for applications meeting the following criteria: Both: 1 For the treatment of terminal agitation that is unresponsive to other agents; and 2 The applicant is part of a multidisciplinary team working in palliative care.</p>			
166	ADALIMUMAB – Special Authority see SA1371 – Retail pharmacy Inj 20 mg per 0.4 ml prefilled syringe	1,799.92	2	✓ Humira
185	SODIUM HYALURONATE – Special Authority see SA1388 – Retail pharmacy Eye drops 1 mg per ml	22.00	10 ml OP	✓ Hilo-Fresh
	<p>Note: Hilo-Fresh has a 6 month expiry after opening. The Pharmacy Handbook restriction allowing one bottle per month is not relevant and therefore only the prescribed dosage to the nearest OP may be claimed.</p>			
185	PRESERVATIVE FREE OCULAR LUBRICANTS ▶ SA1388 – Special Authority for Subsidy Initial application from any relevant practitioner. Approvals valid for 12 months for patients meeting the following criteria: Both: 1 Confirmed diagnosis by slit lamp of severe secretory dry eye; and 2 Either: 2.1 Patient is using eye drops more than four times daily on a regular basis; or 2.2 Patient has had a confirmed allergic reaction to preservative in eye drop. Renewal from any relevant practitioner. Approvals valid for 24 months where the patient continues to require lubricating eye drops and has benefited from treatment.			
185	RETINOL PALMITATE Eye oint 138 mcg per g	3.80	5 g OP	✓ Vita-POS
186	PHARMACY SERVICES – May only be claimed once per patient. * Brand switch fee	4.33	1 fee	✓ BSF Arrow-Quinapril
	<p>The Pharmacode for BSF Arrow-Quinapril is 2441497.</p>			
198	FAT SUPPLEMENT – Special Authority see SA1374 – Hospital pharmacy [HP3] Oil, 250 ml	114.92	4 OP	✓ Liquigen
202	RENAL ORAL FEED 2 KCAL/ML – Special Authority see SA1101 – Hospital pharmacy [HP3] Liquid (apricot), 125 ml	11.52	4 OP	✓ Renilon 7.5
	Liquid (caramel), 125 ml	11.52	4 OP	✓ Renilon 7.5

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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New Listings - effective 1 June 2013

39	MACROGOL 3350 – Special Authority see SA0891 – Retail pharmacy Powder 13.125 g, sachets - Maximum of 60 sach per prescription	18.14	30	✓ Movicol
62	BOSENTAN – Special Authority see SA0967 – Retail pharmacy Tab 62.5 mg	2,000.00	60	✓ pms-Bosentan
	Tab 125 mg	2,000.00	60	✓ pms-Bosentan
211	AMINOACID FORMULA WITHOUT PHENYLALANINE – Special Authority see SA1108 – Hospital pharmacy [HP3] Powder (unflavoured), 29 g	330.12	30	✓ PKU Anamix Junior
213	HIGH FAT LOW CARBOHYDRATE FORMULA – Special Authority see SA1197 – Retail pharmacy Powder (unflavoured)	35.50	300 g OP	✓ KetoCal 4:1

Effective 6 May 2013

72	MALATHION WITH PERMETHRIN AND PIPERONYL BUTOXIDE Spray 0.25% with permethrin 0.5% and piperonyl butoxide 2%.....	11.95	90 g OP	✓ Para Plus
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Effective 1 May 2013

52	PERINDOPRIL * Tab 2 mg	3.75	30	✓ Apo-Perindopril
	* Tab 4 mg	4.80	30	✓ Apo-Perindopril
57	CLONIDINE HYDROCHLORIDE * Tab 25 mcg.....	15.09	112	✓ Clonidine BNM
58	METOLAZONE – Special Authority see SA1323 – Retail pharmacy Tab 5 mg	CBS	50	✓ Zaroxolyn S29
61	HYDRALAZINE HYDROCHLORIDE – Special Authority see SA1321 – Retail pharmacy * Tab 25 mg	CBS	56	✓ Onelink S29
119	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE Inj 2%, 5 ml – Up to 5 inj available on a PSO.....	6.90	25	✓ Lidocaine-Claris
	Inj 2%, 20 ml – Up to 5 inj available on a PSO.....	2.40	1	✓ Lidocaine-Claris
186	PHARMACY SERVICES – May only be claimed once per patient Brand switch fee	4.33	1 fee	✓ BSF Apo-Diltiazem CD
	The Pharmacode for BSF Apo-Diltiazem CD is 2437775			

Check your Schedule for full details
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Generic Mnfr
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Changes to Restrictions

Effective 1 July 2013

29 BLOOD KETONE DIAGNOSTIC TEST METER

a) Meter funded for the purposes of blood ketone diagnostics only. Patient has had one or more episodes of ketoacidosis and is at risk of future episodes. Only one meter per patient will be subsidised every 5 years.

b) **Up to 1 dev available on a PSO.**

Meter 40.00 1 ✓ **Freestyle Optium**

29 KETONE BLOOD BETA-KETONE ELECTRODES

a) Maximum of 20 strip per prescription.

b) **Up to 10 test available on a PSO.**

Test strip – Not on a BSO 15.50 10 strip OP ✓ **Freestyle Optium
Ketone**

29 BLOOD GLUCOSE DIAGNOSTIC TEST METER – Subsidy by endorsement

a) Maximum of 1 pack per prescription.

b) **Up to 1 dev available on a PSO.**

c) A diagnostic blood glucose test meter is subsidised for a patient who:

- i is receiving insulin or sulphonylurea therapy; or
- ii is pregnant and has diabetes; or
- iii is on home TPN at risk of hypoglycaemia or hyperglycaemia; or
- iv has a genetic or an acquired disorder of glucose homeostasis excluding type 1 or type 2 diabetes and metabolic syndrome.

d) ~~CareSens N brand: Brand switch fee payable (Pharmacode 2423138) – see page 186 for details~~

e) ~~CareSens N POP brand: Brand switch fee payable (Pharmacode 2423154) – see page 186 for details~~

f) ~~CareSens II brand: Brand switch fee payable (Pharmacode 2423146) – see page 186 for details~~

g) ~~No patient co-payment payable~~

Only one CareSens meter per patient. No further prescriptions will be subsidised for patients who already have a CareSens meter. For the avoidance of doubt patients who have previously received a funded meter, other than CareSens, are eligible for a CareSens meter. The prescription must be endorsed accordingly. Pharmacists may annotate the prescription as endorsed where there exists a record of prior dispensing of insulin or sulphonylureas.

Meter with 50 lancets, a lancing device and 10 diagnostic test
strips – Note differing brand requirements 20.00 1 OP ✓ **CareSens II**
✓ **CareSens N**
✓ **CareSens N POP**

Note: Only 1 meter available per PSO

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 July 2013 (continued)

30 BLOOD GLUCOSE DIAGNOSTIC TEST STRIP

a) **Up to 50 test available on a PSO.**

b) The number of test strips available on a prescription is restricted to 50 unless:

- 1) Prescribed with insulin or a sulphonylurea but are on a different prescription and endorsed accordingly; or
- 2) Prescribed on the same prescription as insulin or a sulphonylurea in which case the prescription is deemed to be endorsed; or
- 3) Prescribed for a pregnant woman with diabetes and endorsed accordingly; or
- 4) Prescribed for a patient on home TPN at risk of hypoglycaemia or hyperglycaemia and endorsed accordingly; or
- 5) Prescribed for a patient with a genetic or an acquired disorder of glucose homeostasis excluding type 1 or type 2 diabetes and metabolic syndrome and endorsed accordingly.

Blood glucose test strips

– Note differing brand requirements below	10.56	50 test OP	✓ CareSens
	28.75		✓ CareSens N
			✓ Accu-Chek Performa
			✓ Freestyle Optium

a) Accu-Chek Performa brand: Special Authority see SA1294 – Retail pharmacy

b) Freestyle Optium brand: Special Authority see SA1291 – Retail pharmacy

Note: Accu-Chek Performa and Freestyle Optium are not available on a PSO

38 URSODEOXYCHOLIC ACID – Special Authority see **SA1383** ~~††88~~ – Retail pharmacy

Cap 250 mg – For ursodeoxycholic acid oral liquid

formulation refer, page 188	71.50	100	✓ Ursosan
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▶ **SA1383** ~~††88~~ Special Authority for Subsidy

Initial application – (Alagille syndrome or progressive familial intrahepatic cholestasis) - from any relevant practitioner. Approvals valid without further renewal unless notified for applications meeting the following criteria:

Either:

1. Patient has been diagnosed with Alagille syndrome; or
2. Patient has progressive familial intrahepatic cholestasis.

Initial application – (Chronic severe drug induced cholestatic liver injury) - from any relevant practitioner.

Approvals valid for 3 months for applications meeting the following criteria:

All of the following:

1. Patient has chronic severe drug induced cholestatic liver injury; and
2. Cholestatic liver injury not due to Total Parenteral Nutrition (TPN) use in adults; and
3. Treatment with ursodeoxycholic acid may prevent hospital admission or reduce duration of stay.

Initial application – (Cirrhosis) - from any relevant practitioner. Approvals valid for 6 months for applications meeting the following criteria:

Both:

1. Primary biliary cirrhosis confirmed by antimitochondrial antibody titre (AMA) > 1:80, and raised cholestatic liver enzymes with or without raised serum IgM or, if AMA is negative, by liver biopsy; and
2. Patient not requiring a liver transplant (bilirubin > 170 100 umol/l; decompensated cirrhosis).

Initial application – (Pregnancy/Cirrhosis) - from any relevant practitioner. Approvals valid for 6 months where the

Either:

patient diagnosed with cholestasis of pregnancy; or

†. **Both:**

- 1.1. Primary biliary cirrhosis confirmed by antimitochondrial antibody titre (AMA) > 1:80, and raised cholestatic liver enzymes with or without raised serum IgM or, if AMA is negative, by liver biopsy; and
- 1.2. Patient not requiring a liver transplant (bilirubin > 170 100 umol/l; decompensated cirrhosis)

Note: Liver biopsy is not usually required for diagnosis but is helpful to stage the disease.

continued...

Patients pay a manufacturer's surcharge when
the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

continued...

Initial application – (Haematological transplant) - from any relevant practitioner. Approvals valid for 6 months for applications meeting the following criteria:

Both:

1. Patient at risk of veno-occlusive disease or has hepatic impairment and is undergoing conditioning treatment prior to allogenic stem cell or bone marrow transplantation; and
2. Treatment for up to 13 weeks.

Initial application – (Total parenteral nutrition induced cholestasis) from any relevant practitioner. Approvals valid for 6 months for applications meeting the following criteria:

Both:

1. Paediatric patient has developed abnormal liver function as indicated on testing which is likely to be induced by Total Parenteral Nutrition (TPN); and
2. Liver function has not improved with modifying the TPN composition.

Renewal (Chronic severe drug induced cholestatic liver injury) from any relevant practitioner. Approvals valid for 6 months where the patient continues to benefit from treatment.

Renewal – (Pregnancy/cirrhosis) - from any relevant practitioner. Approvals valid for 2 years where the treatment remains appropriate and the patient is benefiting from treatment.

Renewal - (Total parenteral nutrition induced cholestasis) from any relevant practitioner. Approvals valid for 6 months where a paediatric patient continues to require TPN and who is benefiting from treatment, defined as a sustained improvement in bilirubin levels.

Note: Ursodeoxycholic acid is not an appropriate therapy for patients requiring a liver transplant (bilirubin > 170 100 µmol/l; decompensated cirrhosis). These patients should be referred to an appropriate transplant centre.

Treatment failure – doubling of serum bilirubin levels, absence of a significant decrease in ALP or ALT and AST, development of varices, ascites or encephalopathy, marked worsening of pruritus or fatigue, histological progression by two stages, or to cirrhosis, need for transplantation.

38	MUCILAGINOUS LAXATIVES ISPAGHULA (PSYLLIUM) HUSK – Only on a prescription * Dry Powder for oral soln.....	5.51	500 g OP	✓ Konsyl-D
47	PROTAMINE SULPHATE * Inj 10 mg per ml, 5 ml	22.40	10	Artex S29
		(101.61)		
52	QUINAPRIL – Brand switch fee payable (Pharmacode 2441497) * Tab 5 mg	3.44	90	✓ Arrow-Quinapril 5
	* Tab 10 mg	4.64	90	✓ Arrow-Quinapril 10
	* Tab 20 mg	6.34	90	✓ Arrow-Quinapril 20
78	CYPROTERONE ACETATE WITH ETHINYLOESTRADIOL * Tab 2 mg with ethinyloestradiol 35 mcg and 7 inert tabs – Up to 84 tab available on a PSO	3.89	84	✓ Ginet 84
80	SOLIFENACIN SUCCINATE – Special Authority see SA0998 – Retail pharmacy Tab 5 mg	56.50	30	✓ Vesicare
	Tab 10 mg	56.50	30	✓ Vesicare

▶ SA0998 Special Authority for Subsidy

Initial application from any relevant practitioner. Approvals valid without further renewal, unless notified, where the patient has overactive bladder and a documented intolerance of, or is non-responsive to oxybutynin.

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

80	TOLTERODINE – Special Authority see SA1272 – Retail pharmacy		
	Tab 1 mg	14.56	56 ✓ Arrow-Tolterodine
	Tab 2 mg	14.56	56 ✓ Arrow-Tolterodine

▶ SA1272 Special Authority for Subsidy

Initial application from any relevant practitioner. Approvals valid without further renewal, unless notified, where the patient has overactive bladder and a documented intolerance of, or is non-responsive to oxybutynin.

82	PREDNISON		
	Tab 1 mg	10.68	500 ✓ Apo-Prednisone

Note: the removal of the stat symbol will be temporary due to a potential out of stock

85	PROPYLTHIOURACIL – Special Authority see SA1199 – Retail pharmacy		
	Tab 50 mg	35.00	100 ✓ PTU ^{S29}

Note: Propylthiouracil is not recommended for patients under the age of 18 years unless the patient is pregnant and other treatments are contraindicated.

86	CABERGOLINE		
	Tab 0.5 mg – Maximum of 2 tab per prescription; can be waived by Special Authority see SA1370 †03†	6.25 25.00	2 8 ✓ Dostinex ✓ Dostinex

▶ SA1370 †03† Special Authority for Waiver of Rule

Initial application only from an obstetrician, endocrinologist or any relevant practitioner. Approvals valid without further renewal unless notified for applications meeting the following criteria where the patient has:

- 1) pathological hyperprolactinemia; or
- 2) acromegaly*.

Renewal (for patients who have previously been funded under Special Authority form SA1031) only from an obstetrician, endocrinologist or gynaecologist any relevant practitioner. Approvals valid without further renewal unless notified where the patient has previously held a Special Authority which has expired and the treatment remains appropriate and the patient is benefiting from treatment.

Indication marked with * is an Unapproved Indication.

86	DANAZOL – Retail pharmacy–Specialist		
	Cap 100 mg	68.33	100 ✓ Azol
	Cap 200 mg	97.83	100 ✓ Azol

88	CEFAZOLIN SODIUM – Subsidy by endorsement		
	Only if prescribed for dialysis or cystic fibrosis patient cellulitis in accordance with a DHB approved protocol and the prescription is endorsed accordingly.		
	Inj 500 mg	3.99	5 ✓ AFT
	Inj 1 g	3.99	5 ✓ AFT

88	CEFTRIAXONE SODIUM – Subsidy by endorsement		
	a) Up to 5 inj available on a PSO		
	b) Subsidised only if prescribed for a dialysis or cystic fibrosis patient, or the treatment of confirmed ciprofloxacin-resistant gonorrhoea, or the treatment of pelvic inflammatory disease, or the treatment of suspected meningitis in patients who have a known allergy to penicillin, and the prescription or PSO is endorsed accordingly.		
	Inj 500 mg	2.70	1 ✓ Veracol
	Inj 1 g	10.49	5 ✓ Aspen Ceftriaxone

Patients pay a manufacturer's surcharge when the Manufacturer's Price is greater than the Subsidy

^{S29} Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

91	MINOCYCLINE HYDROCHLORIDE * Tab 50 mg – Additional subsidy by Special Authority see SA1355 – Retail pharmacy	5.79 (12.05)	60	Mino-tabs
	SA1355 Special Authority for Manufacturers Price Initial application from any relevant practitioner. Approvals valid without further renewal unless notified where the patient has rosacea.			
92	GENTAMICIN SULPHATE Inj 10 mg per ml, 1 ml – Subsidy by endorsement	8.56	5	✓ Mayne
	Only if prescribed for a dialysis or cystic fibrosis patient or for prophylaxis of endocarditis or complicated urinary tract infection , and the prescription is endorsed accordingly.			
	Inj 10 mg per ml, 2 ml – Subsidy by endorsement	175.10	25	✓ APP Pharmaceuticals S29
	Only if prescribed for a dialysis or cystic fibrosis patient or for prophylaxis of endocarditis or complicated urinary tract infection , and the prescription is endorsed accordingly.			
	Inj 40 mg per ml, 2 ml – Subsidy by endorsement	6.50	10	✓ Pfizer
	Only if prescribed for a dialysis or cystic fibrosis patient or for prophylaxis of endocarditis or complicated urinary tract infection , and the prescription is endorsed accordingly.			
92	MOXIFLOXACIN – Special Authority see SA1358 1065 – Retail pharmacy No patient co-payment payable Tab 400 mg	52.00	5	✓ Avelox
	SA1358 1065 Special Authority for Subsidy Initial application - (Mycoplasma genitalium) from any relevant practitioner. Approvals valid for 1 month for applications meeting the following criteria: All of the following: 1. Has nucleic acid amplification test (NAAT) confirmed Mycoplasma genitalium*; and 2. Has tried and failed to clear infection using azithromycin; and 3. Treatment is only for 7 days. Initial application - (Penetrating eye injury) only from an ophthalmologist. Approvals valid for 1 month where the patient requires prophylaxis following a penetrating eye injury and treatment is for 5 days only. Note: Indications marked with * are Unapproved Indications (refer to Section A: General Rules, Part I (Interpretations and Definitions) and Part IV (Miscellaneous Provisions) rule 4.6).			
93	VANCOMYCIN HYDROCHLORIDE – Subsidy by endorsement Only if prescribed for a dialysis or cystic fibrosis patient or in the treatment of pseudomembranous colitis or for prophylaxis of endocarditis or for treatment of Clostridium difficile following metronidazole failure and the prescription is endorsed accordingly. Inj 500 mg	3.58	1	✓ Mylan
94	FLUCONAZOLE Powder for oral suspension 10 mg per ml – Special Authority see SA1359 1148 – Retail pharmacy	34.56	35 ml	✓ Diflucan
	SA1359 1148 Special Authority for Subsidy Initial application – (Systemic candidiasis) from any relevant practitioner. Approvals valid for 6 weeks for applications meeting the following criteria: Both: 1. Patient requires prophylaxis for, or treatment of systemic candidiasis; and			

continued...

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$

Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 July 2013 (continued)

continued...

2. Patient is unable to swallow capsules.

Initial application – (Immunocompromised) from any relevant practitioner. Approvals valid for 6 months for applications meeting the following criteria:

All of the following:

1. Patient is immunocompromised; and
2. Patient is at moderate to high risk of invasive fungal infection; and
3. Patient is unable to swallow capsules.

Renewal – (**Systemic candidiasis**) from any relevant practitioner. Approvals valid for 6 weeks for applications meeting the following criteria:

Both:

1. Patient requires prophylaxis for, or treatment of systemic candidiasis; and
2. Patient is unable to swallow capsules.

Renewal – (Immunocompromised) from any relevant practitioner. Approvals valid for 6 month for applications meeting the following criteria:

All of the following:

1. Patient remains immunocompromised; and
2. Patient remains at moderate to high risk of invasive fungal infection; and
3. Patient is unable to swallow capsules.

98	LAMIVUDINE – Special Authority see SA1360 0032– Retail pharmacy			
	Tab 100 mg	32.50	28	✓ Zetlam
	Oral liq 5 mg per ml	90.00	240 ml	✓ Zeffix

► SA1360 0032 Special Authority for Subsidy

Initial application only from a gastroenterologist, infectious disease specialist, paediatrician or general physician **or on the recommendation of a gastroenterologist, infectious disease specialist, paediatrician or general physician.** Approvals valid for 1 year for applications meeting the following criteria:

Any of the following:

1. All of the following:

- 1.1. HBsAg positive for more than 6 months; and
- 1.1.2. HBsAg positive or HBV DNA positive defined as $> 100,000$ copies per ml by quantitative PCR at a reference laboratory; and
- 1.1.3. ALT greater than twice upper limit of normal or bridging fibrosis or cirrhosis (Metavir stage 3 or 4 or equivalent) on liver histology or clinical/radiological evidence of cirrhosis; or

21 HBV DNA positive cirrhosis prior to liver transplantation; or

32 HBsAg positive and have had a liver, kidney, heart, lung or bone marrow transplant; or

43 Hepatitis B virus naïve patient who has received a liver transplant from an anti-HBc (Hepatitis B core antibody) positive donor; or

4 Hepatitis B surface antigen (HbsAg) positive patient who is receiving chemotherapy for a malignancy, or high dose steroids (at least 20mg/day for at least 7 days) or who has received such treatment within the previous two months; or

5 Hepatitis B surface antigen positive patient who is receiving anti tumour necrosis factor treatment; or

6 Hepatitis B core antibody (anti-HBc) positive patient who is receiving rituximab plus high dose steroids (e.g. R-CHOP).

2. All of the following:

- 2.1. No continuing alcohol abuse or intravenous drug use; and
- 2.2. Not coinfected with HCV or HDV; and
- 2.3. Neither ALT nor AST greater than 10 times upper limit of normal; and
- 2.4. No history of hypersensitivity to lamivudine; and
- 2.5. No previous lamivudine therapy with genotypically proven lamivudine resistance.

continued...

Patients pay a manufacturer's surcharge when the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

continued...

Renewal only from a gastroenterologist, infectious disease specialist, paediatrician or general physician **or on the recommendation of a gastroenterologist, infectious disease specialist, paediatrician or general physician.**

Approvals valid for 2 years for applications meeting the following criteria:

Any of the following:

Renewal for patients who have maintained continuous treatment and response to lamivudine

1. All of the following:

- 1.1. Have maintained continuous treatment with lamivudine; and
- 1.2. Most recent test result shows continuing biochemical response (normal ALT); and
- 1.3. HBV DNA < 100,000 copies per ml by quantitative PCR at a reference laboratory.

Renewal when given in combination with adefovir dipivoxil for patients with cirrhosis and resistance to lamivudine

2. All of the following

- 2.1. lamivudine to be used in combination with adefovir dipivoxil; and
- 2.2. patient is cirrhotic; and
Documented resistance to lamivudine, defined as:
- 2.3. patient has raised serum ALT (> 1 x ULN); and
- 2.4. patient has HBV DNA greater than 100,000 copies per mL, or viral load = 10 fold over nadir; and
- 2.5. detection of M204I or M204V mutation.

Renewal when given in combination with adefovir dipivoxil for patients with resistance to adefovir dipivoxil

3. All of the following

- 3.1. lamivudine to be used in combination with adefovir dipivoxil; and
Documented resistance to adefovir, defined as:
- 3.2. patient has raised serum ALT (> 1 x ULN); and
- 3.3. patient has HBV DNA greater than 100,000 copies per mL, or viral load = 10 fold over nadir; and
- 3.4. detection of N236T or A181T/V mutation

98 ENTECAVIR – Special Authority see **SA1361 0977** – Retail pharmacy
Tab 0.5 mg 400.00 30 ✓ **Baraclude**

➔ **SA1361 0977** Special Authority for Subsidy

Initial application only from a gastroenterologist or infectious disease specialist. Approvals valid without further renewal unless notified for applications meeting the following criteria:

All of the following:

- 1 Patient has confirmed Hepatitis B infection (HBsAg positive for more than 6 months); and
- 2 Patient is Hepatitis B nucleoside analogue treatment-naïve; and
- 3 Entecavir dose 0.5 mg/day; and
- 4 Either:
 - 4.1 ALT greater than upper limit of normal; or
 - 4.2 Bridging fibrosis (**Metavir stage 3 or greater or moderate fibrosis**) or cirrhosis on liver histology; and
- 5 Either:
 - 5.1 HBeAg positive; or
 - 5.2 patient has ≥ 2,000 IU HBV DNA units per ml and fibrosis (Metavir stage 2 or greater) on liver histology; and
- 6 No continuing alcohol abuse or intravenous drug use; and
- 7 Not co-infected with HCV, HIV or HDV; and
- 8 Neither ALT nor AST greater than 10 times upper limit of normal; and
- 9 No history of hypersensitivity to entecavir; and
- 10 No previous documented lamivudine resistance (either clinical or genotypic).

Notes:

- Entecavir should be continued for 6 months following documentation of complete HBeAg seroconversion (defined as loss of HBeAg plus appearance of anti-HBe plus loss of serum HBV DNA) for patients who were HBeAg positive prior to commencing this agent. This period of consolidation therapy should be extended to 12 months in patients with advanced fibrosis (Metavir Stage F3 or F4).
- Entecavir should be taken on an empty stomach to improve absorption

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

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Schedule page ref

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Changes to Restrictions - effective 1 July 2013 (continued)

- 100 TENOFOVIR DISOPROXIL FUMARATE – Subsidy by endorsement; can be waived by Special Authority see **SA1362** ~~1047~~
Endorsement for treatment of HIV/AIDS: Prescription is deemed to be endorsed if tenofovir disoproxil fumarate is co-prescribed with another anti-retroviral subsidised under Special Authority **SA1364** ~~1025~~ and the prescription is annotated accordingly by the Pharmacist or endorsed by the prescriber.
Note: Tenofovir disoproxil fumarate prescribed under endorsement for the treatment of HIV/AIDS is included in the count of up to 4 subsidised antiretrovirals for the purposes of Special Authority **SA1364** ~~1025~~.
Tab 300 mg 531.00 30 ✓ **Viread**

► **SA1362** ~~1047~~ Special Authority for Waiver of Rule

Initial application - (Chronic Hepatitis B) Only from a gastroenterologist, infectious disease specialist or general physician. Approvals valid without further renewal, unless notified, for applications meeting the following criteria:
Any of the following

1. Patient has confirmed Hepatitis B infection (HBsAg positive for more than 6 months); and
 - 1.1. All of the following
 - 1.1.1. Patient has had previous lamivudine, adefovir or entecavir therapy; and
 - 1.1.2. HBV DNA greater than 20,000 IU/mL or increased = 10 fold over nadir; and
 - 1.1.3. Any of the following:
 - 1.1.3.1. Lamivudine resistance - detection of M204I/V mutation; or
 - 1.1.3.2. Adefovir resistance - detection of A181T/V or N236T mutation; or
 - 1.1.3.3. Entecavir resistance - detection of relevant mutations including I169T, L180M T184S/A/I/L/G/C/M, S202C/G/I, M204V or M250I/V mutation; or
2. Patient is either listed or has undergone liver transplantation for HBV; or
3. **Patient has decompensated cirrhosis with a Mayo score >20.**

Initial application - (Pregnant, **active hepatitis B**) only from a gastroenterologist, infectious disease physician or general physician. Approvals valid for **12 months** ~~4 months~~ for applications meeting the following criteria:
Both:

- 1 **Patient is HBsAg positive and pregnant; and**
- 2 ~~Either:~~
 - 2-1 HBV DNA > 20,000 IU/mL and ALT > ULN; or
 - 2-2 HBV DNA > 100 million IU/mL and ALT normal.

Renewal - (Subsequent Pregnancy **or breastfeeding, active hepatitis B**) only from a gastroenterologist, infectious disease specialist or general physician. Approvals valid for **12 months** ~~4 months~~ for applications meeting the following criteria:

- Both:
- 1 Patient is HBsAg positive and pregnant or breastfeeding; and
 - 2 ~~Either:~~
 - 2-1 HBV DNA > 20,000 IU/mL and ALT > ULN; or
 - 2-2 HBV DNA > 100 million IU/mL and ALT normal.

Initial application - (Pregnant, **prevention of vertical transmission**) only from a gastroenterologist, infectious disease specialist or general physician. Approvals valid for **6 months** ~~4 months~~ for applications meeting the following criteria:

- Both:
- 1 Patient is HBsAg positive and pregnant; and
 - 2 ~~Either:~~
 - 2-1 HBV DNA > 20,000 IU/mL and ALT > ULN; or
 - 2-2 HBV DNA > ~~100~~ **20** million IU/mL and ALT normal.

Renewal - (Subsequent pregnancy, prevention of vertical transmission) only from a gastroenterologist, infectious disease specialist or general physician. Approvals valid for **6 months** ~~4 months~~ for applications meeting the following criteria:

continued...

Changes to Restrictions - effective 1 July 2013 (continued)

continued...

Both:

1 Patient is HBsAg positive and pregnant; and

2 ~~Either:~~

2.1 HBV DNA > 20,000 IU/mL and ALT > ULN; or

2.2 HBV DNA > 100 20 million IU/mL and ALT normal.

100	VALACICLOVIR – Special Authority see SA1363 0957 – Retail pharmacy Tab 500 mg	102.72	30	✓ Valtrex
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▶ **SA1363 0957** Special Authority for Subsidy

Initial application – (recurrent genital herpes) from any medical practitioner. Approvals valid for 12 months where the patient has genital herpes with 2 or more breakthrough episodes in any 6 month period while treated with aciclovir 400 mg twice daily.

Renewal – (recurrent genital herpes) from any medical practitioner. Approvals valid for 12 months where the treatment remains appropriate and the patient is benefiting from treatment.

Initial application – (ophthalmic zoster) from any medical practitioner. Approvals valid without further renewal unless notified where the patient has previous history of ophthalmic zoster and the patient is at risk of vision impairment.

Initial application – (CMV prophylaxis) from any medical practitioner. Approvals valid for 3 months where the patient has undergone organ transplantation.

Initial application – (immunocompromised patients) from any medical practitioner. Approvals valid without further renewal unless notified for applications meeting the following criteria:

All of the following:

1 Patients is immunocompromised; and

2 Patient has herpes zoster; and

3 Valaciclovir is to be given for a maximum of 7 days per course.

102 ANTIRETROVIRALS

▶ **SA1364 1025** Special Authority for Subsidy

Initial application – (Confirmed HIV/AIDS) only from a named specialist. Approvals valid without further renewal unless notified for applications meeting the following criteria:

Both:

1 Confirmed HIV infection; and

2 Any of the following:

2.1 Symptomatic patient; or

2.2 Patient aged 12 months and under; or

2.3 Both:

2.3.1 Patient aged 1 to 5 years; and

2.3.2 Any of the following:

2.3.2.1 CD4 counts < 1000 cells/mm³; or

2.3.2.2 CD4 counts < 0.25 × total lymphocyte count; or

2.3.2.3 Viral load counts > 100000 copies per ml; or

2.4 Both:

2.4.1 Patient aged 6 years and over; and

2.4.2 CD4 counts < ~~350~~500 cells/mm³.

Notes: Tenofovir disoproxil fumarate prescribed under endorsement for HIV/AIDS is included in the count of up to 4 subsidised antiretrovirals.

Subsidies for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals.

continued...

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 July 2013 (continued)

continued...

Renewal — (Confirmed HIV/AIDS) only from a named specialist. Approvals valid without further renewal unless notified where the treatment remains appropriate and the patient is benefiting from treatment.

Initial application — (Prevention of maternal transmission) only from a named specialist. Approvals valid for 1 year for applications meeting the following criteria:

Either:

- 1 Prevention of maternal foetal transmission; or
- 2 Treatment of the newborn for up to eight weeks.

Notes: Tenofovir disoproxil fumarate prescribed under endorsement for HIV/AIDS is included in the count of up to 4 subsidised antiretrovirals. Subsidies for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals. Some antiretrovirals are unapproved or contraindicated for this indication. Practitioners prescribing these medications should exercise their own skill, judgement, expertise and discretion, and make their own prescribing decisions with respect to the use of a Pharmaceutical for an indication for which it is not approved or contraindicated.

Initial application – (post-exposure prophylaxis following non-occupational exposure to HIV) only from a named specialist.

Approvals valid for 4 weeks for applications meeting the following criteria:

Both:

- 1 Treatment course to be initiated within 72 hours post exposure; and
- 2 Either:
 - 2.1 Patient has had unprotected receptive anal intercourse with a known HIV positive person; or
 - 2.2 Patient has shared intravenous injecting equipment with a known HIV positive person; **or**
 - 2.3 **Patient has had non-consensual intercourse and the clinician considers that the risk assessment indicates prophylaxis is required**

Notes: Tenofovir disoproxil fumarate prescribed under endorsement for HIV/AIDS is included in the count of up to 4 subsidised antiretrovirals. Subsidies for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals.

Renewal – (second or subsequent post-exposure prophylaxis) only from a named specialist. Approvals valid for 4 weeks for applications meeting the following criteria:

Both:

- 1 Treatment course to be initiated within 72 hours post exposure; and
- 2 Either:
 - 2.1 Patient has had unprotected receptive anal intercourse with a known HIV positive person; or
 - 2.2 Patient has shared intravenous injecting equipment with a known HIV positive person; or
 - 2.3 **Patient has had non-consensual intercourse and the clinician considers that the risk assessment indicates prophylaxis is required**

Initial application – (Percutaneous exposure) only from a named specialist. Approvals valid for 6 weeks where the patient has percutaneous exposure to blood known to be HIV positive.

Notes: Tenofovir disoproxil fumarate prescribed under endorsement for HIV/AIDS is included in the count of up to 4 subsidised antiretrovirals. Subsidies for a combination of up to four antiretroviral medications. The combination of a protease inhibitor and low-dose ritonavir given as a booster (either as part of a combination product or separately) will be counted as one protease inhibitor for the purpose of accessing funding to antiretrovirals.

Renewal – (Second or subsequent percutaneous exposure) only from a named specialist. Approvals valid for 6 weeks where the patient has percutaneous exposure to blood known to be HIV positive.

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

- 105 Guidelines for the use of interferon in the treatment of hepatitis C:
Physicians considering treatment of patients with hepatitis C should discuss cases with a gastroenterologist or an infectious disease physician. All subjects undergoing treatment require careful monitoring for side effects. Patients should be otherwise fit.
Hepatocellular carcinoma should be excluded by ultrasound examination and alpha-fetoprotein level.
Criteria for Treatment
- a) Diagnosis
- Anti-HCV positive on at least two occasions with a positive PCR for HCV-RNA and preferably confirmed by a supplementary RIBA test; or
 - PCR-RNA positive for HCV on at least 2 occasions if antibody negative; or
 - Anti-HCV positive on at least two occasions with a positive supplementary RIBA test with a negative PCR for HCV RNA but with a liver biopsy consistent with 2(b) following.
- ~~b) Establishing Active Chronic Liver Disease~~
- ~~- Confirmed HCV infection and serum ALT/AST levels measured on at least three occasions over six months averaging > 1.5 x upper limit of normal. (ALT is the preferable enzyme); or~~
 - ~~- Liver biopsy showing significant inflammatory activity (active hepatitis) with or without cirrhosis. This is not a necessary requirement for those patients with coagulopathy. (Some patients have active disease on histology with normal transaminase enzymes).~~
- Exclusion Criteria
- a) Autoimmune liver disease. (Interferon may exacerbate autoimmune liver disease as well as other autoimmune diseases such as thyroid disease).
 - b) Pregnancy.
 - c) Neutropenia (<2.0 x 10⁹) and/or thrombocytopenia.
 - d) Continuing alcohol abuse and/or continuing intravenous drug users.
- Dosage
The current recommended dosage is 3 million units of interferon alpha-2a or interferon alpha-2b administered subcutaneously three times a week for 52 weeks (twelve months).
- Exit Criteria
The patient's response to interferon treatment should be reviewed at either three or four months. Interferon treatment should be discontinued in patients who do not show a substantial reduction (50%) in their mean pre-treatment ALT level at this stage.

107 PEGYLATED INTERFERON ALPHA-2A – Special Authority see **SA1365** ~~1134~~ – Retail pharmacy
See prescribing guideline

Inj 135 mcg prefilled syringe	362.00	1	✓Pegasys
	1,448.00	4	✓Pegasys
Inj 180 mcg prefilled syringe	450.00	1	✓Pegasys
	1,800.00	4	✓Pegasys
Inj 135 mcg prefilled syringe × 4 with ribavirin tab 200 mg × 112	1,799.68	1 OP	✓Pegasys RBV Combination Pack
Inj 135 mcg prefilled syringe × 4 with ribavirin tab 200 mg × 168.....	1,975.00	1 OP	✓Pegasys RBV Combination Pack
Inj 180 mcg prefilled syringe × 4 with ribavirin tab 200 mg × 112.....	2,059.84	1 OP	✓Pegasys RBV Combination Pack
Inj 180 mcg prefilled syringe × 4 with ribavirin tab 200 mg × 168	2,190.00	1 OP	✓Pegasys RBV Combination Pack

continued...

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
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Changes to Restrictions - effective 1 July 2013 (continued)

continued...

▶ **SA1365** ~~134~~ Special Authority for Subsidy

Initial application – (chronic hepatitis C - genotype 1, 4, 5 or 6 infection or co-infection with HIV **or genotype 2 or 3 post liver transplant**) from any specialist. Approvals valid for 18 months for applications meeting the following criteria:

Both:

1 **Any of the following: Either:**

1.1 Patient has chronic hepatitis C, genotype 1, 4, 5 or 6 infection; or

1.2 Patient has chronic hepatitis C and is co-infected with HIV; ~~and~~ **or**

1.3 Patient has chronic hepatitis C genotype 2 or 3 and has received a liver transplant; and

2 Maximum of 48 weeks therapy.

Notes:

- Consider stopping treatment if there is absence of a virological response (defined as at least a 2-log reduction in viral load) following 12 weeks of treatment since this is predictive of treatment failure.

- Consider reducing treatment to 24 weeks if serum HCV RNA level at Week 4 is undetectable by sensitive PCR assay (less than 50 IU/ml) AND Baseline serum HCV RNA is less than 400,000 IU/ml

Initial application – (chronic hepatitis C - genotype 2 or 3 infection without co-infection with HIV) from any specialist.

Approvals valid for 12 months for applications meeting the following criteria:

Both:

1 Patient has chronic hepatitis C, genotype 2 or 3 infection; and

2 Maximum of 6 months therapy.

Initial application – (Hepatitis B) only from a gastroenterologist, infectious disease specialist or general physician.

Approvals valid for 18 months for applications meeting the following criteria:

All of the following:

1 Patient has confirmed Hepatitis B infection (HBsAg positive for more than 6 months); and

2 Patient is Hepatitis B treatment-naive; and

3 ALT > 2 times Upper Limit of Normal; and

4 HBV DNA < 10 log₁₀ IU/ml; and

5 Either:

5.1 HBeAg positive; or

5.2 serum HBV DNA ≥ 2,000 units/ml and significant fibrosis (≥ Metavir Stage F2 **or moderate fibrosis**);
and

6 Compensated liver disease; and

7 No continuing alcohol abuse or intravenous drug use; and

8 Not co-infected with HCV, HIV or HDV; and

9 Neither ALT nor AST > 10 times upper limit of normal; and

10 No history of hypersensitivity or contraindications to pegylated interferon; and

11 Maximum of 48 weeks therapy.

Notes:

- Approved dose is 180 mcg once weekly.

- The recommended dose of Pegylated Interferon-alpha 2a is 180 mcg once weekly.

- In patients with renal insufficiency (calculated creatinine clearance less than 50ml/min), Pegylated Interferon-alpha 2a dose should be reduced to 135 mcg once weekly.

- In patients with neutropaenia and thrombocytopaenia, dose should be reduced in accordance with the datasheet guidelines.

- Pegylated Interferon-alpha 2a is not approved for use in children.

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

119	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE Inj 1%, 5 ml ampoule – Up to 25 inj available on a PSO.....	35.60 6.90	50 25	✓Xylocaine ✓Lidocaine-Clarix
130	SUMATRIPTAN Inj 12 mg per ml, 0.5 ml cartridge – Maximum of 10 inj per prescription	13.80	2 OP	✓Arrow-Sumatriptan
131	HYOSCINE (SCOPOLAMINE) – Special Authority see SA1387 0939 – Retail pharmacy Patch 1.5 mg	11.95	2	✓Scopoderm TTS

▶ SA1387 0939 Special Authority for Subsidy

Initial application from any relevant practitioner. Approvals valid for 1 year for applications meeting the following criteria:

All of the following:

Either:

- 1 Control of intractable nausea, vomiting, or inability to swallow saliva in the treatment of malignancy or chronic disease **where the patient cannot tolerate or does not adequately respond to oral anti-nausea agents; or and**
- 2 **Control of clozapine-induced hypersalivation where trials of at least two other alternative treatments have proven ineffective.**
- 2—Patient cannot tolerate or does not adequately respond to oral anti-nausea agents; and
- 3—The applicant must specify the underlying malignancy or chronic disease.

Renewal from any relevant practitioner. Approvals valid for 1 year where the treatment remains appropriate and the patient is benefiting from treatment.

166	ADALIMUMAB – Special Authority see SA1371 1156 – Retail pharmacy Inj 20 mg per 0.4 ml prefilled syringe	1,799.92	2	✓Humira
	Inj 40 mg per 0.8 ml prefilled pen	1,799.92	2	✓HumiraPen
	Inj 40 mg per 0.8 ml prefilled syringe	1,799.92	2	✓Humira

Note: Only the new criteria is listed below existing criteria remains unchanged.

▶ SA1371 1156 Special Authority for Subsidy

Initial application - (juvenile idiopathic arthritis) only from a named specialist or rheumatologist. Approvals valid for 4 months for applications meeting the following criteria:

Either:

1 Both:

1.1 The patient has had an initial Special Authority approval for etanercept for juvenile idiopathic arthritis (JIA); and

1.2 Either:

1.2.1 The patient has experienced intolerable side effects from etanercept; or

1.2.2 The patient has received insufficient benefit from etanercept to meet the renewal criteria for etanercept for juvenile idiopathic arthritis; or

2 All of the following:

2.1 To be used as an adjunct to methotrexate therapy or monotherapy where use of methotrexate is limited by toxicity or intolerance; and

2.2 Patient diagnosed with JIA; and

2.3 Patient has had severe active polyarticular course JIA for 6 months duration or longer; and

2.4 Patient has tried and not responded to at least three months of oral or parenteral methotrexate (at a dose of 10-20 mg/m² weekly or at the maximum tolerated dose) in combination with either oral corticosteroids (prednisone 0.25 mg/kg or at the maximum tolerated dose) or a full trial of serial intra-articular corticosteroid injections; and

2.5 Both:

2.5.1 Either:

continued...

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Changes to Restrictions - effective 1 July 2013 (continued)

continued...

- 2.5.1.1 Patient has persistent symptoms of poorly-controlled and active disease in at least 20 swollen, tender joints; or
- 2.5.1.2 Patient has persistent symptoms of poorly-controlled and active disease in at least four joints from the following: wrist, elbow, knee, ankle, shoulder, cervical spine, hip; and
- 2.5.2 Physician's global assessment indicating severe disease.

Initial application – (fistulising Crohn's disease) only from a gastroenterologist. Approvals valid for 6 months for applications meeting the following criteria:

- 1 Patient has confirmed Crohn's disease; and
- 2 Either
 - 2.1 Patient has one or more complex externally draining enterocutaneous fistula(e); or
 - 2.2 Patient has one or more rectovaginal fistula(e); and
- 3 A Baseline Fistula Assessment has been completed and is no more than 1 month old at the time of application; and
- 4 The patient will be assessed for response to treatment after 4 months' adalimumab treatment (see Note).
Note: a maximum of 4 months' adalimumab will be subsidised on an initial Special Authority approval for fistulising Crohn's disease.

Renewal – (juvenile idiopathic arthritis) only from a named specialist, rheumatologist or Practitioner on the recommendation of a named specialist or rheumatologist. Approvals valid for 6 months for applications meeting the following criteria:

All of the following:

- 1 Either:
 - 1.1 Applicant is a named specialist or rheumatologist; or
 - 1.2 Applicant is a Practitioner and confirms that a named specialist or rheumatologist has provided a letter, email or fax recommending that the patient continues with adalimumab treatment; and
- 2 Subsidised as an adjunct to methotrexate therapy or monotherapy where use of methotrexate is limited by toxicity or intolerance; and
- 3 Either:
 - 3.1 Following 3 to 4 months' initial treatment, the patient has at least a 50% decrease in active joint count and an improvement in physician's global assessment from baseline; or
 - 3.2 On subsequent reapplications, the patient demonstrates at least a continuing 30% improvement inactive joint count and continued improvement in physician's global assessment from baseline.

Renewal – (fistulising Crohn's disease) only from a gastroenterologist or Practitioner on the recommendation of a gastroenterologist. Approvals valid for 6 months meeting the following criteria:

Both:

- 1 Either:
 - 1.1 Applicant is a gastroenterologist; or
 - 1.2 Applicant is a Practitioner and confirms that a gastroenterologist has provided a letter, email or fax recommending that the patient continues with adalimumab treatment; and
- 2 Either:
 - 2.1 The number of open draining fistulae have decreased from baseline by at least 50%; or
 - 2.2 There has been a marked reduction in drainage of all fistula(e) from baseline as demonstrated by a reduction in the Fistula Assessment score, together with less induration and patient-reported pain.

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 July 2013 (continued)

- 175 BEE VENOM ALLERGY TREATMENT – Special Authority see **SA1368 0053** – Retail pharmacy
- | | | | |
|---|--------|------|--------|
| Maintenance kit - 6 vials 120 mcg freeze dried venom,
6 diluent 1.8 ml | 285.00 | 1 OP | ✓Albay |
| Treatment kit - Inj 1 vial 550 mcg freeze dried venom,
1 diluent 9 ml, 3 diluent 1.8 ml..... | 285.00 | 1 OP | ✓Albay |
- ➔ **SA1368 0053** Special Authority for Subsidy
Initial application only from a relevant specialist. Approvals valid for 2 years for applications meeting the following criteria:
Both:
1 RAST or skin test positive; and
2 Patient has had severe generalised reaction to the sensitising agent.
Renewal only from a relevant specialist. Approvals valid for 2 years where the treatment remains appropriate and the patient is benefiting from treatment.
- 175 WASP VENOM ALLERGY TREATMENT – Special Authority see **SA1367 0053** – Retail pharmacy
- | | | | |
|--|--------|------|--------|
| Treatment kit (Paper wasp venom) - 1 vial 550 mcg freeze
dried polister venom, 1 diluent 9 ml, 1 diluent 1.8 ml | 285.00 | 1 OP | ✓Albay |
| Treatment kit (Yellow jacket venom) - 1 vial 550 mcg freeze
dried vespula venom, 1 diluent 9 ml, 1 diluent 1.8 ml | 285.00 | 1 OP | ✓Albay |
- ➔ **SA1367 0053** Special Authority for Subsidy
Initial application only from a relevant specialist. Approvals valid for 2 years for applications meeting the following criteria:
Both:
1 RAST or skin test positive; and
2 Patient has had severe generalised reaction to the sensitising agent.
Renewal only from a relevant specialist. Approvals valid for 2 years where the treatment remains appropriate and the patient is benefiting from treatment.
- 196 CARBOHYDRATE
- ➔ **SA1373 1001** Special Authority for Subsidy
Initial application – (Cystic fibrosis or **kidney disease renal failure**) only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 3 years for applications meeting the following criteria:
Either:
1 cystic fibrosis; or
2 chronic **kidney disease renal failure** or continuous ambulatory peritoneal dialysis (CAPD) patient
Initial application – (Indications other than cystic fibrosis or renal failure) only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:
Any of the following:
1 cancer in children; or
2 cancers affecting alimentary tract where there are malabsorption problems in patients over the age of 20 years; or
3 **faltering growth in an infant/child; or failure to thrive; or growth deficiency; or**
4 bronchopulmonary dysplasia; or
5 premature and post premature infant; or
6 inborn errors of metabolism; or
7 **for use as a component in a modular formula.**

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Changes to Restrictions - effective 1 July 2013 (continued)

196 CARBOHYDRATE AND FAT

➔ ~~SA1376 1091~~ Special Authority for Subsidy

Initial application – (Cystic fibrosis) only from a dietitian, relevant specialist or vocationally registered general practitioner.

Approvals valid for 3 years for applications meeting the following criteria:

Both:

- 1 infant **or child** aged four years or under; and
- 2 cystic fibrosis.

Initial application – (Indications other than cystic fibrosis) only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Both:

- 1 Infant **or child** aged four years or under; and
- 2 Any of the following:
 - 2.1 cancer in children; or
 - 2.2 **faltering growth; or**
~~failure to thrive; or~~
~~growth deficiency; or~~
 - 2.3 bronchopulmonary dysplasia; or
 - 2.4 premature and post premature infants.

197 FAT

➔ ~~SA1374 1092~~ Special Authority for Subsidy

Initial application — (Indications other than inborn errors of metabolism) only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Any of the following:

- 1 **faltering growth in an infant/child; or**
~~failure to thrive where other high-calorie products are inappropriate or inadequate; or~~
~~growth deficiency; or~~
- 2 bronchopulmonary dysplasia; or
- 3 fat malabsorption; or
- 4 lymphangiectasia; or
- 5 short bowel syndrome; or
- 6 infants with necrotising enterocolitis; or
- 7 biliary atresia; **or**
- 8 **for use in a ketogenic diet; or**
- 9 **chyle leak; or**
- 10 **acites; or**
- 11 **for use as a component in a modular formula.**

198 PROTEIN

➔ ~~SA1375 1093~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Either:

Any of the following:

- 1 protein losing enteropathy; or
- 2 high protein needs (eg burns); **or**
- 3 **for use as a component in a modular formula.**

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 July 2013 (continued)

198 RESPIRATORY PRODUCTS

▶ ~~SA1094~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year where the patient has CORD and hypercapnia, **defined as a CO2 value exceeding 55 mmHg.**

199 FAT MODIFIED PRODUCTS

▶ ~~SA1381~~ ~~1096~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Either:

Any of the following:

- 1 Patient has metabolic disorders of fat metabolism; or
- 2 Patient has ~~chylolthorax~~ **a chyle leak; or**
- 3 **Modified as a modular feed for adults.**

199 HIGH PROTEIN PRODUCTS

▶ ~~SA1378~~ ~~1097~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Both:

- 1 Anorexia and weight loss; and

2 Either:

- 2.1 ~~decompensating liver disease without encephalopathy; or~~
- 2.2 ~~protein losing gastro-enteropathy~~

Either:

- 1 **decompensating liver disease without encephalopathy; or**
- 2 **protein losing gastro-enteropathy.**

200 PAEDIATRIC PRODUCTS

▶ ~~SA1379~~ ~~1224~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Both:

- 1 Child is aged one to ten years; and
- 2 Any of the following:
 - 2.1 the child is being fed via a tube or a tube is to be inserted for the purposes of feeding; or
 - 2.2 any condition causing malabsorption; or
~~failure to thrive; or~~
 - 2.3 **faltering growth in an infant/child; or**
 - 2.4 increased nutritional requirements; or
 - 2.5 **the child is being transitioned from TPN or tube feeding to oral feeding.**

200 PAEDIATRIC PRODUCTS FOR CHILDREN WITH CHRONIC RENAL FAILURE

▶ ~~SA1099~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 3 years where the patient is a child (up to 18 years) with **acute or chronic kidney disease** ~~renal failure~~.

201 RENAL PRODUCTS

▶ ~~SA1101~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 3 years where the patient has acute or chronic ~~renal failure~~ **kidney disease**.

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 July 2013 (continued)

202 SPECIALISED AND ELEMENTAL PRODUCTS

▶ ~~SA1377~~ ~~†102~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

Any of the following:

- 1 malabsorption; or
- 2 short bowel syndrome; or
- 3 enterocutaneous fistulas; or
~~pancreatitis.~~

4 eosinophilic oesophagitis; or

5 inflammatory bowel disease; or

6 patients with multiple food allergies requiring enteral feeding.

203 RENAL ORAL FEED ~~†~~KCAL/ML **2** KCAL/ML – Special Authority see SA1101 – Hospital pharmacy [HP3]

Liquid..... 3.80 237 ml OP ✓ **Suplena**

208 ADULT PRODUCTS HIGH CALORIE

▶ ~~SA1195~~ Special Authority for Subsidy

Note: Only the criteria that have been amended are shown.

Initial application — (Indications other than cystic fibrosis) only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 1 year for applications meeting the following criteria:

All of the following:

- 1 Any of the following:
 - 1.1 any condition causing malabsorption; or
~~failure to thrive; or~~
 - 1.2 **faltering growth in an infant/child; or**
 - 1.3 increased nutritional requirements; or
 - 1.4 fluid restricted; and
- 2 other lower calorie products have been tried; and
- 3 patient has substantially increased metabolic requirements or is fluid restricted.

212 EXTENSIVELY HYDROLYSED FORMULA – Special Authority see **SA1380** ~~†220~~ – Hospital pharmacy [HP3]

Powder 15.21 450 g OP ✓ **Pepti Junior Gold
Karicare Aptamil**

▶ ~~SA1380~~ ~~†220~~ Special Authority for Subsidy

Initial application only from a dietitian, relevant specialist or vocationally registered general practitioner. Approvals valid for 6 months for applications meeting the following criteria:

Any of the following:

- 1 Both:
 - 1.1 Cows milk formula is inappropriate due to severe intolerance or allergy to its protein content; and
 - 1.2 Either:
 - 1.2.1 Soy milk formula has been trialed without resolution of symptoms; or
 - 1.2.2 Soy milk formula is considered clinically inappropriate or contraindicated; or
- 2 Severe malabsorption; or
- 3 Short bowel syndrome; or
- 4 Intractable diarrhea; or
- 5 Biliary atresia; or
- 6 Cholestatic liver diseases causing malabsorption; or
~~Chylous ascite; or~~
~~Chyl thorax; or~~
- 7 Cystic fibrosis; or
- 8 Proven fat malabsorption; or
- 9 Severe intestinal motility disorders causing significant malabsorption; or
- 10 Intestinal failure.

Patients pay a manufacturer's surcharge when
the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 June 2013

29	ACARBOSE – Brand switch fee payable (Pharmacode 2433257) – see page 177 for details * Tab 50 mg 9.82 90 ✓ Accarb * Tab 100 mg 15.83 90 ✓ Accarb		
58	METOLAZONE – Special Authority see SA1323 – Retail pharmacy Tab 5 mg CBS 50 ✓ Zaroxolyn ^{\$29} 1 ✓ Metolazone ^{\$29}		
	▶ SA1323 Special Authority for Subsidy Initial application from any relevant practitioner. Approvals valid without further renewal unless notified where used for applications meeting the following criteria: the treatment of patients with refractory heart failure who are intolerant or have not responded to loop diuretics and/or loop-thiazide combination therapy. Either: 1) For the treatment of heart failure in patients who are intolerant or have not responded to ACE inhibitors and/or angiotensin receptor blockers; or 2) For the treatment of heart failure, in patients in whom treatment with ACE inhibitors and/or angiotensin receptor blockers is not tolerated due to renal impairment.		
72	ETHINYLLOESTRADIOL WITH LEVONORGESTREL * Tab 20 mcg with levonorgestrel 100 mcg and 7 inert tab – Up to 84 tab available on a PSO – Brand switch fee payable (Pharmacode 2427958) – see page 177 for details 2.95 84 ✓ Ava 20 ED		
98	LAMIVUDINE – Special Authority see SA0832 – Retail pharmacy – Brand switch fee payable (Pharmacode 2433257) – see page 177 for details Tab 100 mg 32.50 28 ✓ Zetlam		
118	ENTACAPONE – Brand switch fee payable (Pharmacode 2433249) – see page 177 for details ▲ Tab 200 mg 47.92 100 ✓ Entapone		
131	METOCLOPRAMIDE HYDROCHLORIDE * Tab 10 mg – For metoclopramide oral liquid formulation refer, page 188 3.95 100 ✓ Metamide		
182	CHLORAMPHENICOL Eye drops 0.5% 1.20 10 ml OP ✓ Chlorafast Funded for use in the ear* Indications marked with* are Unapproved Indications.		
182	EYE PREPARATIONS Eye preparations are only funded for use in the eye, unless explicitly stated otherwise . The exception is pilocarpine eye drops 1%, 2% and 4% which are subsidised for oral use pursuant to the Standard Formulae.		
184	PILOCARPINE Eye drops 4% - Subsidised for oral use pursuant to the Standard Formulae 7.99 15 ml OP ✓ Isopto Carpine		

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Restrictions - effective 1 May 2013

52 PERINDOPRIL

Perindopril will be funded to the level of the ex-manufacturer price listed in the Schedule for patients who were taking these ACE inhibitors for the treatment of congestive heart failure prior to 1 June 1998. The prescription must be endorsed accordingly. We recommend that the words used to indicate eligibility are "certified condition" or an appropriate description of the patient such as "congestive heart failure", "CHF", "congestive cardiac failure" or "CCF". Definition of Congestive Heart Failure At the request of some prescribers the PTAC Cardiovascular subcommittee has provided a definition of congestive heart failure for the purposes of the funding of the manufacturer's surcharge: "Clinicians should use their clinical judgement. Existing patients would be eligible for the funding of the surcharge if the patient shows signs and symptoms of congestive heart failure, and requires or has in the past required concomitant treatment with a diuretic. The definition could also be considered to include patients post myocardial infarction with an ejection fraction of less than 40%."

* Tab 2 mg – Higher subsidy of \$18.50 per 30 tab
with Endorsement.....

3.75 30
(18.50)

✓ Apo-Perindopril
Coversyl

* Tab 4 mg – Higher subsidy of \$25.00 per 30 tab
with Endorsement.....

4.80 30
(25.00)

✓ Apo-Perindopril
Coversyl

52 TRANDOLAPRIL

Higher subsidy by endorsement is available Trandolapril will be funded to the level of the ex-manufacturer price listed in the Schedule for patients who were taking these ACE inhibitors **trandolapril** for the treatment of congestive heart failure prior to 1 June 1998. The prescription must be endorsed accordingly. We recommend that the words used to indicate eligibility are "certified condition" or an appropriate description of the patient such as "congestive heart failure", "CHF", "congestive cardiac failure" or "CCF". Definition of Congestive Heart Failure At the request of some prescribers the PTAC Cardiovascular subcommittee has provided a definition of congestive heart failure for the purposes of the funding of the manufacturer's surcharge: "Clinicians should use their clinical judgement. Existing patients would be eligible for the funding of the surcharge if the patient shows signs and symptoms of congestive heart failure, and requires or has in the past required concomitant treatment with a diuretic. The definition could also be considered to include patients post myocardial infarction with an ejection fraction of less than 40%." **For the purposes of this endorsement, congestive heart failure includes patients post myocardial infarction with an ejection fraction of less than 40%. Patients who started on trandolapril after 1 June 1998 are not eligible for full subsidy by endorsement.**

* Cap 1 mg – Higher subsidy of \$18.67 per 28 cap
with Endorsement.....

3.06 28
(18.67)

Gopten

* Cap 2 mg – Higher subsidy of \$27.00 per 28 cap
with Endorsement.....

4.43 28
(27.00)

Gopten

57 DILTIAZEM HYDROCHLORIDE

* Cap long-acting 120 mg

– Brand switch fee payable (Pharmacode 2437775).....

31.83 500

✓ Apo-Diltiazem CD

* Cap long-acting 180 mg

– Brand switch fee payable (Pharmacode 2437775).....

47.67 500

✓ Apo-Diltiazem CD

* Cap long-acting 240 mg

– Brand switch fee payable (Pharmacode 2437775).....

63.58 500

✓ Apo-Diltiazem CD

Patients pay a manufacturer's surcharge when
the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Restrictions - effective 1 May 2013 (continued)

91	CIPROFLOXACIN – Subsidy by endorsement 1) Subsidised only if: a) Patient has: i) microbiologically confirmed and clinically significant pseudomonas infection; or ii) prostatitis; or iii) pyelonephritis; or iv) gonorrhoea; b) Prescription or PSO is written by, or on the recommendation of, an infectious disease physician or a clinical microbiologist; and 2) The prescription or PSO is endorsed accordingly. Recommended for patients with the any of the following: i) microbiologically confirmed and clinically significant pseudomonas infection; or ii) prostatitis; or iii) pyelonephritis; or iv) gonorrhoea; or			
	Tab 250 mg – Up to 5 tab available on a PSO	2.20	28	✓ Cipflox
	Tab 500 mg – Up to 5 tab available on a PSO	3.00	28	✓ Cipflox
		10.71	100	✓ Cipflox
	Tab 750 mg	5.15	28	✓ Cipflox
		5.52	30	✓ Ciprofloxacin Rex
92	CLINDAMYCIN Cap hydrochloride 150 mg – Maximum of 4 cap per prescription; can be waived by endorsement – Retail pharmacy-Specialist... 9.90 Specialist must be an infectious disease physician or a clinical microbiologist Inj phosphate 150 mg per ml, 4 ml – Retail pharmacy-Specialist... 160.00 Prescriptions must be written by, or on the recommendation of, an infectious disease physician or a clinical microbiologist.		16	✓ Clindamycin ABM
			10	✓ Dalacin C
94	ITRACONAZOLE Cap 100 mg – Subsidy by endorsement 4.25 Funded for tinea vesicolor where topical treatment has not been successful and diagnosis has been confirmed by mycology, or for tinea unguium where terbinafine has not been successful in eradication or the patient is intolerant to terbinafine and diagnosis has been confirmed by mycology and the prescription is endorsed accordingly. Can be waived by endorsement - Retail pharmacy - Specialist. Specialist must be an infectious disease physician, clinical microbiologist, clinical immunologist or dermatologist.		15	✓ Itrazole
96	ISONIAZID – Retail pharmacy-Specialist a) No patient co-payment payable b) Prescriptions must be written by, or on the recommendation of, an internal medicine physician, paediatrician, clinical microbiologist, dermatologist or public health physician			
	* Tab 100 mg	20.00	100	✓ PSM
	* Tab 100 mg with rifampicin 150 mg	90.04	100	✓ Rifinah
	* Tab 150 mg with rifampicin 300 mg	179.57	100	✓ Rifinah
119	LIGNOCAINE HYDROCHLORIDE LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE Viscous soln 2%..... 55.00 Inj 1%, 5 ml – Up to 5 inj available on a PSO 35.00 Inj 2%, 5 ml – Up to 5 inj available on a PSO 23.00 6.90 Inj 1%, 20 ml – Up to 5 inj available on a PSO 20.00 Inj 2%, 20 ml – Up to 5 inj available on a PSO 15.00 2.40		200 ml 50 50 25 5 5 1	✓ Xylocaine Viscous ✓ Xylocaine ✓ Xylocaine ✓ Lidocaine-Clarix ✓ Xylocaine ✓ Xylocaine ✓ Lidocaine-Clarix

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Subsidy and Manufacturer's Price

Effective 1 July 2013

28	INSULIN LISPRO WITH INSULIN LISPRO PROTAMINE (↓ subsidy) ▲ Inj lispro 25% with insulin lispro protamine 75% 100 u per ml, 3 ml.....	42.66	5	✓ Humalog Mix 25
	▲ Inj lispro 50% with insulin lispro protamine 50% 100 u per ml, 3 ml	42.66	5	✓ Humalog Mix 50
38	ISPAGUHULA (PHYLLIUM) HUSK – Only on a prescription (↓ subsidy) * Powder for oral soln	5.51	500 g OP	✓ Konsyl-D
39	SODIUM CITRATE WITH SODIUM LAURYL SULPHOACETATE – Only on a prescription (↓ subsidy) Enema 90 mg with sodium lauryl sulphoacetate 9 mg per ml, 5 ml.....	19.95	50	✓ Micolette
51	TERAZOSIN (↓ subsidy) * Tab 1 mg	0.50	28	✓ Arrow
	* Tab 2 mg	0.45	28	✓ Arrow
	* Tab 5 mg	0.68	28	✓ Arrow
51	CILAZAPRIL (↓ subsidy) * Tab 0.5 mg	2.00	90	✓ Zapril
	* Tab 2.5 mg	4.31	90	✓ Zapril
	* Tab 5 mg	6.98	90	✓ Zapril
57	CLONIDINE HYDROCHLORIDE (↓ subsidy) * Tab 25 mcg	13.47	100	✓ Dixarit
58	SPIRONOLACTONE (↓ subsidy) * Tab 25 mg	3.65	100	✓ Spirotone
	* Tab 100 mg	11.80	100	✓ Spirotone
60	EZETIMIBE – Special Authority see SA1045 – Retail pharmacy (↓ subsidy) Tab 10 mg	34.43	30	✓ Ezetrol
60	EZETIMIBE WITH SIMVASTATIN – Special Authority see SA1046 – Retail pharmacy (↓ subsidy) Tab 10 mg with simvastatin 10 mg.....	36.68	30	✓ Vytorin
	Tab 10 mg with simvastatin 20 mg.....	38.70	30	✓ Vytorin
	Tab 10 mg with simvastatin 40 mg.....	41.40	30	✓ Vytorin
	Tab 10 mg with simvastatin 80 mg.....	45.45	30	✓ Vytorin
65	FUSIDIC ACID (↑ subsidy) Oint 2%	3.45	15 g OP	✓ Foban
	a) Maximum of 15 g per prescription b) Only on a prescription c) Not in combination			
78	LEVONORGESTREL (↓ subsidy) * Tab 750 mcg.....	3.50	2	✓ Next Choice
78	MEDROXYPROGESTERONE ACETATE (↓ subsidy) * Inj 150 mg per ml, 1 ml syringe – Up to 5 inj available on a PSO.....	7.00	1	✓ Depo-Provera

Patients pay a manufacturer's surcharge when
the Manufacturer's Price is greater than the Subsidy

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Subsidy and Manufacturer's Price - effective 1 July 2013 (continued)

92	CLINDAMYCIN (↓ subsidy) Inj phosphate 150 mg per ml, 4 ml – Retail pharmacy-Specialist	100.00	10	✓ Dalacin C
99	ACICLOVIR (↓ subsidy) * Tab dispersible 200 mg	1.78	25	✓ Lovir
	* Tab dispersible 400 mg	5.98	56	✓ Lovir
	* Tab dispersible 800 mg	6.64	35	✓ Lovir
119	TETRABENAZINE (↓ subsidy) Tab 25 mg	118.00	112	✓ Motetis
119	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE (↓ subsidy) Inj 2%, 5 ml – Up to 5 inj available on a PSO	13.80	50	✓ Xylocaine
	Inj 2%, 20 ml – Up to 5 inj available on a PSO	12.00	5	✓ Xylocaine
120	DIHYDROCODEINE TARTRATE (↓ subsidy) Tab long-acting 60 mg	13.64	60	✓ DHC Continus
122	MORPHINE SULPHATE (↓ subsidy) a) Only on a controlled drug form b) No patient co-payment payable c) Safety medicine; prescriber may determine dispensing frequency			
	Tab long-acting 10 mg	1.95	10	✓ Arrow-Morphine LA
	Tab long-acting 30 mg	2.98	10	✓ Arrow-Morphine LA
	Tab long-acting 60 mg	5.75	10	✓ Arrow-Morphine LA
	Tab long-acting 100 mg	6.45	10	✓ Arrow-Morphine LA
122	MORPHINE TARTRATE (↑ subsidy) a) Only on a controlled drug form b) No patient co-payment payable c) Safety medicine; prescriber may determine dispensing frequency			
	Inj 80 mg per ml, 1.5 ml	35.60	5	✓ Hospira
	Inj 80 mg per ml, 5 ml	107.67	5	✓ Hospira
124	SERTRALINE (↓ subsidy) * Tab 50 mg	3.64	90	✓ Arrow-Sertraline
	* Tab 100 mg	6.28	90	✓ Arrow-Sertraline
125	VENLAFAXINE – Special Authority see SA1061 – Retail pharmacy (↓ subsidy)			
	Tab 37.5 mg	7.84	28	✓ Arrow-Venlafaxine XR
	Tab 75 mg	13.94	28	✓ Arrow-Venlafaxine XR
	Tab 150 mg	17.08	28	✓ Arrow-Venlafaxine XR
	Tab 225 mg	27.14	28	✓ Arrow-Venlafaxine XR
	Cap 37.5 mg	8.71	28	✓ Efexor XR
	Cap 75 mg	17.42	28	✓ Efexor XR
	Cap 150 mg	21.35	28	✓ Efexor XR

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Subsidy and Manufacturer's Price - effective 1 July 2013 (continued)

130	SUMATRIPTAN (↓ subsidy)			
	Tab 50 mg	1.19	4	✓ Arrow-Sumatriptan
		29.80	100	✓ Arrow-Sumatriptan
	Tab 100 mg	1.10	2	✓ Arrow-Sumatriptan
		54.80	100	✓ Arrow-Sumatriptan
	Inj 12 mg per ml, 0.5 ml cartridge			
	– Maximum of 10 inj per prescription	13.80	2 OP	✓ Arrow-Sumatriptan
140	NITRAZEPAM – Safety medicine; prescriber may determine dispensing frequency (↑ subsidy)			
	Tab 5 mg	4.98	100	✓ Nitrados
	‡ Safety cap for extemporaneously compounded oral liquid preparations.			
145	NALTREXONE HYDROCHLORIDE – Special Authority see SA0909 – Retail pharmacy (↓ subsidy)			
	Tab 50 mg	79.00	30	✓ Naltraccord
149	METHOTREXATE (↓ subsidy)			
	* Inj 25 mg per ml, 2 ml – PCT – Retail pharmacy-Specialist	20.20	5	✓ Hospira
	* Inj 25 mg per ml, 20 ml – PCT – Retail pharmacy-Specialist	27.78	1	✓ Hospira
151	DOCETAXEL – PCT only – Specialist (↓ subsidy)			
	Inj 20 mg per ml, 1 ml	48.75	1	✓ Taxotere
	Inj 20 mg per ml, 4 ml	195.00	1	✓ Taxotere
	Inj 1 mg for ECP	2.63	1 mg	✓ Baxter
153	TEMOZOLOMIDE – Special Authority see SA1063 – Retail pharmacy (↓ subsidy)			
	Cap 5 mg	8.00	5	✓ Temaccord
	Cap 20 mg	36.00	5	✓ Temaccord
	Cap 100 mg	175.00	5	✓ Temaccord
	Cap 250 mg	410.00	5	✓ Temaccord
154	VINCISTINE SULPHATE (↓ subsidy)			
	Inj 1 mg per ml, 1 ml – PCT			
	– Retail pharmacy-Specialist	64.80	5	✓ Hospira
	Inj 1 mg per ml, 2 ml – PCT			
	– Retail pharmacy-Specialist	69.60	5	✓ Hospira
	Inj 1 mg for ECP – PCT only – Specialist	9.45	1 mg	✓ Baxter
166	BACILLUS CALMETTE-GUERIN (BCG) VACCINE – PCT only – Specialist (↓ subsidy)			
	Subsidised only for bladder cancer.			
	Inj 2-8 × 100 million CFU	149.37	1	✓ OncoTICE
178	IPRATROPIUM BROMIDE (↓ subsidy)			
	Nebuliser soln, 250 mcg per ml, 1 ml			
	– Up to 40 neb available on a PSO	3.26	20	✓ Univent
	Nebuliser soln, 250 mcg per ml, 2 ml			
	– Up to 40 neb available on PSO	3.37	20	✓ Univent

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Changes to Subsidy and Manufacturer's Price - effective 1 June 2013

53	AMIODARONE HYDROCHLORIDE (↓ subsidy) Inj 50 mg per ml, 3 ml ampoule – Up to 6 inj available on a PSO.....	22.80	6	✓ Cordarone-X
72	MALATHION WITH PERMETHRIN AND PIPERONYL BUTOXIDE (↓ subsidy) Spray 0.25% with permethrin 0.5% and piperonyl butoxide 2%.....	11.15	90 g OP	✓ Para Plus
87	METYRAPONE (↑ subsidy) Cap 250 mg – Retail pharmacy-Specialist	520.00	50	✓ Metopirone
88	PRAZQUANTEL (↑ subsidy) Tab 600 mg	68.00	8	✓ Biltricid
193	METHYLCELLULOSE (↑ subsidy) Powder	36.95	100 g	✓ MidWest

Effective 1 May 2013

52	PERINDOPRIL (↑ subsidy) Tab 2 mg	3.75 (18.50)	30	Coversyl
	Tab 4 mg	4.80 (25.00)	30	Coversyl
66	CALAMINE (↑ price) a) Only on a prescription b) Not in combination Crm, aqueous, BP	1.77 (3.80)	100 g	Home Essential
78	LEVONORGESTREL (↓ subsidy) * Tab 1.5 mg	3.50	1	✓ Postinor-1
	a) Maximum of 2 tab per prescription b) Up to 5 tab available on a PSO			
120	CODEINE PHOSPHATE (↓ subsidy) – Safety medicine; prescriber may determine dispensing frequency Tab 15 mg	4.75	100	✓ PSM
	Tab 30 mg	5.80	100	✓ PSM
	Tab 60 mg	12.50	100	✓ PSM
132	AMISULPRIDE (↓ subsidy) – Safety medicine; prescriber may determine dispensing frequency Tab 100 mg	6.22	30	✓ Solian
	Tab 200 mg	21.92	60	✓ Solian
	Tab 400 mg	44.52	60	✓ Solian
	Oral liq 100 mg per ml	52.50	60 ml	✓ Solian

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to General Rules

Effective 1 July 2013

- 11 ~~"Assessed Pharmaceuticals" means the list of Pharmaceuticals set out in Section H Part III of the Schedule, that have been or are being assessed by PHARMAC.~~
- 12 **"Optional Pharmaceuticals" means the list of National Contract Pharmaceuticals set out in Section H Part III of the Schedule.**
- 12 ~~"Hospital Pharmaceuticals" means National Contract Pharmaceuticals, DV Pharmaceuticals, Discretionary Community Supply Pharmaceuticals and Assessed Pharmaceuticals.~~ **the list of Pharmaceuticals set out in Section H Part II of the Schedule which includes some National Contract Pharmaceuticals.**
- 12 "Community Pharmaceutical" means a Pharmaceutical listed in Section A to G and Section I of the Pharmaceutical Schedule that is subsidised by the Funder from the Pharmaceutical Budget for use in the community.
- 12 ~~"Discretionary Community Supply Pharmaceutical" means the list of Pharmaceuticals set out in Section H Part IV of the Schedule, which may be funded by a DHB Hospital from its own budget for use in the community.~~
- 12 ~~"Hospital Pharmaceuticals in the Community (HPC)" means the pathway under the Named Patient Pharmaceutical Assessment policy to allow District Health Board hospitals to fund a medicine for a patient in the community if this is more affordable for the DHB than paying for the treatment that would otherwise need to be provided.~~
- 14 **"Practitioner"** means a Doctor, a Dentist, a Dietitian, a Midwife, a Nurse Prescriber or an Optometrist or a Pharmacist as those terms are defined in the Pharmaceutical Schedule.
- 16 **"Unlisted Pharmaceutical" means a Pharmaceutical that is within the scope of a Hospital Pharmaceutical, but is not listed in Section H Part II.**
- 19 ~~3.6 Pharmacists' prescriptions~~
The following apply to every prescription written by a Pharmacist:
- 3.6.1 ~~Prescriptions written by a Pharmacist for a Community Pharmaceutical will only be subsidised where they are for the CareSens, CareSens N and CareSens N POP blood glucose diagnostic meters and annotated appropriately.~~
- 3.6.2 ~~The prescribing and dispensing of blood glucose diagnostic meters by Pharmacists must be in accordance with regulations 41 and 42 of the Medicines Regulations 1984.~~

Effective 1 May 2013

- 18 3.3 Original Packs, Certain Antibiotics and Unapproved Medicines
- 3.3.2 If a Community Pharmaceutical is either:
- the liquid oral form of an antibiotic to which a diluent must be added by the Contractor at the time of dispensing; or
 - an unapproved medicine supplied under Section 29 of the Medicines Act 1981 **excluding any medicine listed as Cost, Brand, Source of Supply**, and it is prescribed or ordered by a Practitioner in an amount that does not coincide with the amount contained in one or more standard packs of that Community Pharmaceutical, Subsidy will be paid for the amount prescribed or ordered by the Practitioner in accordance with either clause 3.1 or clause 3.3 of the Schedule, and for the balance of any pack or packs from which the Community Pharmaceutical has been dispensed. At the time of dispensing the Contractor must keep a record of the quantity discarded. To ensure wastage is reduced, the Contractor should reduce the amount dispensed to make it equal to the quantity contained in a whole pack where:

continued...

Patients pay a manufacturer's surcharge when the Manufacturer's Price is greater than the Subsidy
50

§29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Changes to General Rules – effective 1 May 2013 (continued)

continued...

- a) the difference the amount dispensed and the amount prescribed by the Practitioner is less than 10% (eg; if a prescription is for 105 mls then a 100 ml pack would be dispensed); and
- b) in the reasonable opinion of the Contractor the difference would not affect the efficacy of the course of treatment prescribed by the Practitioner.

Note: For the purposes of audit and compliance it is an act of fraud to claim wastage and then use the wastage amount for any subsequent prescription.

Effective 1 April 2013

13 "Hospital Pharmacy-Specialist" means that the Community Pharmaceutical is not eligible for Subsidy unless it is supplied by a hospital or pharmacy contracted to the Funder to dispense as a hospital pharmacy to an Outpatient either:

- a) on a Prescription signed by a Specialist, or
- b) where the treatment with the Community Pharmaceutical has been recommended by a Specialist, on the Prescription of a practitioner which is either:
 - i) endorsed with the words "recommended by [name of specialist and year of authorisation]" and signed by the Practitioner, or
 - ii) **endorsed with the word 'protocol' which means "initiated in accordance with DHB hospital approved protocol"**,
 - iii) annotated by the dispensing pharmacist, following verbal confirmation from the Practitioner of the name of the Specialist and date of recommendation, with the words "recommended by [name of specialist and date of authorisation], confirmed by [practitioner]". Where the Contractor has an electronic record of such an Endorsement or Annotation from a previous prescription for the same Community Pharmaceutical written by a prescriber for the same patient, they may annotate the prescription accordingly.

"As recommended by a Specialist" to be interpreted as **either**:

- 1)
 - a) follows a substantive consultation with an appropriate Specialist;
 - b) the consultation to relate to the Patient for whom the Prescription is written;
 - c) consultation to mean communication by referral, telephone, letter, facsimile or email;
 - d) except in emergencies consultation to precede annotation of the Prescription; and
 - e) both the specialist and the General Practitioner must keep a written record of the consultation; **or**
- 2) **treatment with the Community Pharmaceutical has been initiated in accordance with a DHB hospital approved protocol**

15 "Retail Pharmacy-Specialist" means that the Community Pharmaceutical is only eligible for Subsidy if it is either:

- a) supplied on a Prescription or Practitioner's Supply Order signed by a Specialist, or,
- b) in the case of treatment recommended by a Specialist, supplied on a Prescription or Practitioner's Supply Order and either:
 - i) endorsed with the words "recommended by [name of Specialist and year of authorisation]" and signed by the Practitioner, or
 - ii) **endorsed with the word 'protocol' which means "initiated in accordance with DHB hospital approved protocol", or**
 - iii) Annotated by the dispensing pharmacist, following verbal confirmation from the Practitioner of the name of the Specialist and date of recommendation, with the words "recommended by [name of specialist and year of authorisation], confirmed by [practitioner]". Where the Contractor has an electronic record of such an Endorsement or Annotation from a previous prescription for the same Community Pharmaceutical written by a prescriber for the same patient, they may annotate the prescription accordingly.

"As recommended by a Specialist" to be interpreted as **either**:

- 1)
 - a) follows a substantive consultation with an appropriate Specialist;
 - b) the consultation to relate to the Patient for whom the Prescription is written;

continued...

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ **fully subsidised**

Changes to General Rules – effective 1 May 2013 (continued)

continued...

- c) consultation to mean communication by referral, telephone, letter, facsimile or email;
 - d) except in emergencies consultation to precede annotation of the Prescription; and
 - e) both the Specialist and the General Practitioner must keep a written record of consultation; **or**
- 2) treatment with the Community Pharmaceutical has been initiated in accordance with a DHB hospital approved protocol.**

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Changes to Brand Name

Effective 1 July 2013

209	FOOD THICKENER – Special Authority see SA1106 – Hospital pharmacy [HP3] Powder	7.25	380 g OP	✓ Aptamil Feed Thickener Feed Thickener Karicare Aptamil
212	EXTENSIVELY HYDROLYSED FORMULA – Special Authority see SA1380 – Hospital pharmacy [HP3] Powder	15.21	450 g OP	✓ Gold Pepti Junior Pepti Junior Gold Karicare Aptamil

Effective 1 June 2013

209	FOOD THICKENER – Special Authority see SA1106 – Hospital pharmacy [HP3] Powder	7.25	380 g OP	✓ Karicare Food Thickener Aptamil Feed Thickener
212	EXTENSIVELY HYDROLYSED FORMULA – Special Authority see SA1380 – Hospital pharmacy [HP3] Powder	15.21	450 g OP	✓ Pepti Junior Gold Aptamil Gold Pepti Junior

Effective 1 May 2013

85	LEVOTHYROXINE Tab 50 mcg	1.71	28	✓ Goldshield Mercury Pharma
	‡ Safety cap for extemporaneously compounded oral liquid preparations. Tab 100 mcg.....	1.78	28	✓ Goldshield Mercury Pharma
	‡ Safety cap for extemporaneously compounded oral liquid preparations.			

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

Changes to Section I

Effective 1 May 2013

224	INFLUENZA VACCINE – Hospital pharmacy [Xpharm] Inj	90.00	10	✓ Fluarix ✓ Fluvax
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A) is available each year for patients who meet the following criteria, as set by PHARMAC:

- a) all people 65 years of age and over;
- b) people under 65 years of age **with who**:
 - i) **have** the following cardiovascular disease:
 - 1) ischaemic heart disease,
 - 2) congestive heart disease,
 - 3) rheumatic heart disease,
 - 4) congenital heart disease, or
 - 5) cerebo-vascular disease;
 - ii) **have** the following chronic respiratory disease:
 - 1) asthma, if on a regular preventative therapy, or
 - 2) other chronic respiratory disease with impaired lung function;
 - ~~iii) are children aged four and under who have been hospitalised for respiratory illness or have a history of significant respiratory illness.~~
 - ~~iii) **have** diabetes;~~
 - ~~iv) **have** chronic renal disease;~~
 - ~~v) **have** any cancer, excluding basal and squamous skin cancers if not invasive;~~
 - ~~vi) **have any of the following other conditions**:

 - a) autoimmune disease,
 - b) immune suppression,
 - c) HIV,
 - d) transplant recipients,
 - e) neuromuscular and CNS diseases,
 - f) haemoglobinopathies, or
 - g) **are** children on long term aspirin; **or**~~
 - ~~vii) **are pregnancy pregnant**.~~
- c) people under 18 years of age living within the boundaries of the Canterbury District Health Board
- d) are children aged four and under who have been hospitalised for respiratory illness or have a history of significant respiratory illness**

Unless meeting other the criteria above, the following conditions are excluded from funding:

- a) asthma not requiring regular preventative therapy,
- b) hypertension and/or dyslipidaemia without evidence of end-organ disease.

B) Doctors are the only Contractors entitled to claim payment from the Funder for the supply of influenza vaccine to patients eligible under the above criteria for subsidised immunisation and they may only do so in respect of the influenza vaccine listed in the Pharmaceutical Schedule.

C) Individual DHBs may fund patients over and above the above criteria. The claiming process for these additional patients should be determined between the DHB and Contractor.

D) Stock of the seasonal influenza vaccine is typically available from February until late July with suppliers being required to ensure supply until at least 30 June. Exact start and end dates for each season will be notified each year.

Changes to Section E

Effective 1 July 2013

214	BLOOD GLUCOSE DIAGNOSTIC TEST METER ✓Meter 1
214	BLOOD KETONE DIAGNOSTIC TEST METER ✓Meter 1
214	BLOOD GLUCOSE DIAGNOSTIC TEST STRIP ✓Test strip 50 strip
214	CYPROTERONE ACETATE WITH ETHINYLLOESTRADIOL ✓Tab 2 mg with ethinylloestradiol 35 mcg and 7 inert tabs 84
216	KETONE BLOOD BETA-KETONE ELECTRODES ✓Test strip 10 strip
216	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE ✓Inj 1%, 5 ml 25 5

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

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Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
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Delisted Items

Effective 1 July 2013

27	PANTOPRAZOLE * Inj 40 mg	6.50	1	✓Pantocid IV
49	DEXTROSE WITH ELECTROLYTES Soln with electrolytes.....	6.60 6.75	1,000 ml OP	✓Pedialyte – Fruit ✓Pedialyte – Plain
52	QUINAPRIL * Tab 5 mg	1.15	30	✓Accupril
	* Tab 10 mg	1.55	30	✓Accupril
	* Tab 20 mg	2.11	30	✓Accupril
55	PROPRANOLOL * Tab 10 mg	3.55	100	✓Cardinol
66	CALAMINE a) Only on a prescription b) Not in combination Crm, aqueous, BP	1.77 (3.80)	100 g	Home Essential
81	METHYLPREDNISOLONE SODIUM SUCCINATE – Retail pharmacy-Specialist Inj 500 mg	18.00	1	✓Solu-Medrol
118	LEVODOPA WITH CARBIDOPA * Tab 100 mg with carbidopa 25 mg – For levodopa with carbidopa oral liquid formulation refer, page 188.....	20.00	100	✓Sinemet
	* Tab long-acting 200 mg with carbidopa 50 mg	47.50	100	✓Sinemet CR
	* Tab 250 mg with carbidopa 25 mg	40.00	100	✓Sinemet
	Note – new presentations of Sinemet and Sinemet CR were listed 1 January 2013.			
127	GABAPENTIN Cap 100 mg	7.16	100	✓Nupentin
	Cap 300 mg	11.50	100	✓Nupentin
	Note – the Nupentin capsules in the blister pack are delisted. The Nupentin capsules in bottles will remain listed as fully funded.			
186	PHARMACY SERVICES Brand switch fee.....	4.33	1 fee	✓BSF Alphapharm ✓BSF Nevirapine Alphapharm ✓BSF Caresens II ✓BSF Caresens N ✓BSF Caresens N POP
212	AMINO ACID FORMULA – Special Authority see SA1219 below – Hospital pharmacy [HP3] Powder	53.00	400 g OP	✓Neocate

Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Delisted Items - effective 1 June 2013

53	AMIODARONE HYDROCHLORIDE Inj 50 mg per ml, 3 ml ampoule – Up to 6 inj available on a PSO.....	60.84	10	✓ Cordarone-X
59	BEZAFIBRATE *Tab 200 mg.....	9.70	90	✓ Fibalip
104	STAVUDINE [D4T] – Special Authority see SA1025 – Retail pharmacy Cap 30 mg	377.80	60	✓ Zerit
109	DICLOFENAC SODIUM *Tab EC 25 mg..... *Tab EC 50 mg	1.63 1.60 (2.13)	50 50	✓ Diclofenac Sandoz Diclofenac Sandoz
130	DOMPERIDONE *Tab 10 mg – For domperidone oral liquid formulation refer, page 188.....	3.25 (11.99)	100	Motilium
176	PROMETHAZINE HYDROCHLORIDE *‡ Oral liq 5 mg per 5 ml	2.79 (3.10)	100 ml	Promethazine Winthrop Elixir
186	PHARMACY SERVICES *Brand switch fee	4.33	1 fee	✓ BSF Accarb ✓ BSF Ava 20 ED ✓ BSF Entapone ✓ BSF Zetlam

Effective 1 May 2013

24	CALCIUM CARBONATE WITH AMINOACETIC ACID * Tab 420 mg with aminoacetic acid 180 mg – Higher subsidy of \$6.30 per 100 tab with Endorsement.....	3.00 (6.30)	100	Titralac
Additional subsidy by endorsement is available for pregnant women. The prescription must be endorsed accordingly.				
40	MICONAZOLE Oral gel 20 mg per g	4.95 (8.70)	40 g OP	Daktarin
47	RIVAROXABAN – Special Authority see SA1066 – Retail pharmacy Tab 10 mg	306.00	30	✓ Xarelto

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Check your Schedule for full details
Schedule page ref

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Delisted Items - effective 1 May 2013 (continued)

57	DILTIAZEM HYDROCHLORIDE				
	Cap long-acting 120 mg	1.91			
		(4.34)		Cardizem CD	
	Cap long-acting 180 mg	2.86			
		(6.50)		Cardizem CD	
	Cap long-acting 240 mg	3.81			
		(8.67)		Cardizem CD	
63	SILDENAFIL – Special Authority see SA1293 – Retail pharmacy				
	Tab 25 mg	39.00	4	✓ Viagra	
	Tab 50 mg	43.50	4	✓ Viagra	
	Tab 100 mg – For sildenafil oral liquid formulation refer, page 179	47.00	4	✓ Viagra	
72	CALCIPOTRIOL				
	Oint 50 mcg per g	20.20	30 g OP	✓ Daivonex	
	Soln 50 mcg per ml	33.79	60 ml OP	✓ Daivonex	
89	AZITHROMYCIN – Maximum of 5 days treatment per prescription; can be waived by endorsement For Endorsement, patient has either: i) Received a lung transplant and requires treatment or prophylaxis for bronchiolitis obliterans syndrome *; or ii) Cystic fibrosis and has chronic infection with Pseudomonas aeruginosa or Pseudomonas related gram negative organisms * Indications marked with * are Unapproved Indications				
	Tab 500 mg – Up to 8 tab available on a PSO	1.25	2 OP	✓ Arrow-Azithromycin	
113	PAMIDRONATE DISODIUM				
	Inj 3 mg per ml, 10 ml	16.00	1		
		(37.50)		Pamisol	
	Inj 6 mg per ml, 10 ml	32.00	1		
		(75.00)		Pamisol	
	Inj 9 mg per ml, 10 ml	48.00	1		
		(112.50)		Pamisol	
185	TYLOXAPOL				
	* Eye drops 0.25%	8.63	15 ml OP	✓ Enuclene	
212	AMINO ACID FORMULA – Special Authority see SA1219 – Hospital pharmacy [HP3]				
	Powder (tropical)	53.00	400 g OP	✓ Neocate Advance	

Check your Schedule for full details
Schedule page ref

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Items to be Delisted

Effective 1 August 2013

178	SALBUTAMOL				
	‡ Oral liq 2 mg per 5 ml	1.20	90 ml	✓Broncolin ^{S29}	
		1.99	150 ml	✓Ventolin	
186	PHARMACY SERVICES - May only be claimed once per patient				
	* Brand switch fee.....	4.33	1 fee	✓BSF Apo-Diltiazem CD	

Effective 1 October 2013

45	SODIUM TETRADECYL SULPHATE				
	* Inj 0.5% 2 ml	23.20	5		
		(51.00)		Fibro-vein	
	* Inj 1% 2 ml	25.00	5		
		(55.00)		Fibro-vein	
57	CLONIDINE HYDROCHLORIDE				
	* Tab 25 mcg.....	13.47	100	✓Dixarit	
78	LEVONORGESTREL				
	* Tab 750 mcg.....	3.50	2	✓Next Choice	
88	CEFOXITIN SODIUM – Retail pharmacy-Specialist – Subsidy by endorsement				
	Only if prescribed for dialysis or cystic fibrosis patient and the prescription is endorsed accordingly.				
	Inj 1 g.....	55.00	5	✓Mayne	
89	CEFUROXIME SODIUM				
	Inj 250 mg – Maximum of 3 inj per prescription; can be				
	waived by endorsement	20.97	10	✓Mayne	
	Waiver by endorsement must state that the prescription is for dialysis or cystic fibrosis patient.				
	Inj 1.5 g – Retail pharmacy-Specialist – Subsidy by				
	endorsement	2.65	1	✓Mylan	
		4.04		✓Zinacef	
	Only if prescribed for dialysis or cystic fibrosis patient and the prescription is endorsed accordingly.				
92	FUSIDIC ACID				
	Inj 500 mg sodium fusidate per 10 ml – Retail pharmacy-				
	Specialist – Subsidy by endorsement.....	12.87	1		
		(17.80)		Fucidin	
	Only if prescribed for a dialysis or cystic fibrosis patient and the prescription is endorsed accordingly.				
130	SUMATRIPTAN				
	Tab 50 mg	1.19	4	✓Arrow-Sumatriptan	
	Tab 100 mg	1.10	2	✓Arrow-Sumatriptan	
184	HOMATROPINE HYDROBROMIDE				
	* Eye drops 2%	7.18	15 ml OP	✓Isopto Homatropine	
186	PHARMACY SERVICES				
	* Brand switch fee	4.33	1 fee	✓BSF Arrow-Quinapril	

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

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Check your Schedule for full details Schedule page ref	Subsidy (Mnfr's price) \$	Per	Brand or Generic Mnfr ✓ fully subsidised
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Items to be Delisted - effective 1 November 2013

181	SODIUM CROMOGLYCATE Nasal spray, 4%	15.85	22 ml OP	✓ Rex
211	AMINOACID FORMULA WITHOUT PHENYLALANINE – Special Authority see SA1108 – Hospital pharmacy [HP3] Sachets (tropical)	324.00	30	✓ Phlexy 10

Effective 1 December 2013

31	INSULIN SYRINGES, DISPOSABLE WITH ATTACHED NEEDLE – Maximum of 100 dev per prescription * Syringe 0.3 ml with 31 g × 8 mm needle	13.00	100	✓ ABM
75	CONDOMS * 53 mm extra strength – Up to 144 dev available on a PSO	1.11 13.36	12 144	✓ Gold Knight ✓ Gold Knight
193	METHYLCELLULOSE Powder	14.00	100 g	✓ ABM

Effective 1 January 2014

41	ASCORBIC ACID a) No more than 100 mg per dose b) Only on a prescription * Tab 100 mg	13.80	500	✓ Vitala-C
41	VITAMIN B COMPLEX * Tab, strong, BPC	4.70	500	✓ B-PlexADE
42	VITAMINS * Tab (BPC cap strength)	8.00	1,000	✓ MultiADE
74	MAGNESIUM SULPHATE * Paste	2.98 (4.90)	80 g	PSM
83	OESTROGENS – See prescribing guideline Conjugated, equine tab 300 mcg	3.01 (11.48)	28	Premarin
	Conjugated, equine tab 625 mcg	4.12 (11.48)	28	Premarin
Note: The old Pharmacodes are being delisted; Pharmacodes 2427478 and 2427486 will remain fully funded.				
90	AMOXYCILLIN Drops 125 mg per 1.25 ml	4.00	30 ml OP	✓ Ospamox Paediatric Drops
92	LINCOMYCIN – Retail pharmacy-Specialist Prescriptions must be written by, or on the recommendation of, an infectious disease physician or a clinical microbiologist Inj 300 mg per ml, 2 ml	80.00	5	✓ Lincocin

Patients pay a manufacturer's surcharge when the Manufacturer's Price is greater than the Subsidy
60

S29 Unapproved medicine supplied under Section 29
‡ safety cap reimbursed **Sole Subsidised Supply**

Check your Schedule for full details
Schedule page ref

Subsidy
(Mnfr's price)
\$ Per

Brand or
Generic Mnfr
✓ fully subsidised

Items to be Delisted - effective 1 January 2014 (continued)

126	GABAPENTIN – Special Authority see SA1071 – Retail pharmacy ▲ Cap 400 mg	14.75	100	✓ Nupentin
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Note: This is the blister pack presentation only. The Nupentin capsules in the bottle will remain fully funded.

▲ Three months supply may be dispensed at one time if endorsed "certified exemption" by the prescriber.

* Three months or six months, as applicable, dispensed all-at-once

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