

Pharmaceutical Management Agency
New Zealand
Pharmaceutical Schedule

Section H Update

for Hospital Pharmaceuticals

June 2026

The logo for PHARMAC, featuring the word "PHARMAC" in a bold, uppercase, sans-serif font. Below it, the Māori name "TE PĀTAKA WHAIORANGA" is written in a smaller, uppercase, sans-serif font. The logo is centered within a white circle that overlaps a background of stylized, wavy, concentric lines in shades of gray and white.

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Summary of decisions

EFFECTIVE 1 JUNE 2026

- Beta-HCG low sensitivity urine test kit (CheckTop) midstream – price increase
- Colestyramine (Questran Light (Neon)) powder for oral suspension 4 g sachet – new listing
- Cyclizine lactate (Hamelin) inj 50 mg per ml, 1 ml ampoule – price increase
- Doxorubicin hydrochloride (Adriamycin) inj 2 mg per ml, 100 ml vial – new listing
- Droperidol (Droperidol Panpharma) inj 2.5 mg per ml, 1 ml ampoule – new listing
- Efavirenz with emtricitabine and tenofovir disoproxil (TEEVIR (India)) tab 600 mg with emtricitabine 200 mg and tenofovir disoproxil 245 mg (300 mg as a fumarate) – new listing
- Elexacaftor with tezacaftor, ivacaftor and ivacaftor (Trikafta) – amended restriction criteria
 - Tab elexacaftor 100 mg with tezacaftor 50 mg, ivacaftor 75 mg (56) and ivacaftor 150 mg (28)
 - Tab elexacaftor 50 mg with tezacaftor 25 mg, ivacaftor 37.5 mg (56) and ivacaftor 75 mg (28)
 - Oral granules elexacaftor 100 mg with tezacaftor 50 mg, ivacaftor 75 mg (28) and ivacaftor 75 mg (28), sachets
 - Oral granules elexacaftor 80 mg with tezacaftor 40 mg, ivacaftor 60 mg (28) and ivacaftor 59.5mg (28), sachets
- Ezetimibe with simvastatin (Vytorin) tab 10 mg with simvastatin 80 mg – new listing
- Glucose [Dextrose] (HypoPak Glucose) oral soln 15 g per 80 ml sachet – price increase
- Ivacaftor (Kalydeco) tab 150 mg, oral granules 13.4 mg, 25 mg, 50 mg and 75 mg, sachet – amended restriction criteria
- Montelukast (Montelukast Paediatric (Relonchem)) tab 5 mg – amended brand name
- Nimodipine (Nimorhage) tab 30 mg – new listing
- Oestradiol valerate (Progynova) tab 2 mg – new Pharmacode listing
- Paracetamol (Noumed) oral liq 120 mg per 5 ml, 200 ml – new listing and addition of PSS
- Paracetamol (Ethics) oral liq 120 mg per 5 ml, 200 ml – to be delisted 1 November 2026
- Progesterone (Utrogestan) cap 100 mg – new Pharmacode listing
- Ramipril (Ramipril Viatris) tab 2.5 mg – new listing
- Ribociclib (Kisqali) tab 200 mg, 21, 42 and 63 tab pack – new Pharmacode listing
- Risperidone (Risperidone Sandoz) tab 0.5 mg and 2 mg – new listing

Summary of decisions – effective 1 June 2026 (continued)

- Rizatriptan (Rizatriptan Viatrix) tab orodispersible 10 mg – new listing
- Rotavirus oral vaccine (Rotarix) oral susp live attenuated human rotavirus 1,000,000 CCID50 per dose, squeezable tube – delisted 1 June 2026
- Sodium bicarbonate (Sodibic) cap 840 mg – Pharmacode 2085992 delisted 1 June 2026
- Trimethoprim with sulphamethoxazole [co-trimoxazole] (Deprim) oral liq 8 mg with sulphamethoxazole 40 mg per ml, 100 ml – new Pharmacode listing
- Vanzacaftor with tezacaftor and deutivacaftor (Alyftrek) tab vanzacaftor 4 mg with tezacaftor 20 mg and deutivacaftor 50 mg and tab vanzacaftor 10 mg with tezacaftor 50 mg and deutivacaftor 125 mg – amended restriction criteria
- Venetoclax (Venclexta) tab 14 × 10 mg, 7 × 50 mg, 21 × 100 mg, tab 10 mg, tab 50 mg and tab 100 mg – amended restriction criteria
- Water (Multichem) inj 10 ml ampoule – price increase and addition of PSS
- Water (Fresenius Kabi) inj 10 ml ampoule – delisted 1 June 2026

Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Section H changes to Part II

Effective 1 June 2026

ALIMENTARY TRACT AND METABOLISM

9	GLUCOSE [DEXTROSE] (↑ price) Oral soln 15 g per 80 ml sachet	75.00	50	HypoPak Glucose
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BLOOD AND BLOOD FORMING ORGANS

43	WATER (↑ price and addition of PSS) Inj 10 ml ampoule – 5% DV Nov-26 to 2028	9.50	50	Multichem
	Note – Fresenius Kabi inj 10 ml ampoule delisted 1 June 2026.			

43	SODIUM BICARBONATE (delisted) Cap 840 mg	8.52	100	Sodibic
	Note – This delist applies to Pharmacode 2085992, delisted 1 June 2026.			

CARDIOVASCULAR SYSTEM

45	RAMIPRIL (new listing) Tab 2.5 mg	5.50	30	Ramipril Viatris
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50	NIMODIPINE (new listing) Tab 30 mg	344.83	100	Nimorhage
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54	COLESTYRAMINE (new listing) Powder for oral suspension 4 g sachet	61.50	50	Questran Light (Neon)
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54	EZETIMIBE WITH SIMVASTATIN (new listing) Tab 10 mg with simvastatin 80 mg	14.27	30	Vytorin
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GENITO-URINARY SYSTEM

77	PROGESTERONE (new listing) Cap 100 mg	14.85	30	Utrogestan
	Note – This is a new Pharmacode listing 2730340			

HORMONE PREPARATIONS

82	OESTRADIOL VALERATE (new listing) Tab 2 mg	8.24	56	Progynova
	Note – this is a new Pharmacode listing, 2730979.			

Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 June 2026 (continued)

INFECTIONS

98	TRIMETHOPRIM WITH SULPHAMETHOXAZOLE [CO-TRIMOXAZOLE] (new listing) Oral liq 8 mg with sulphamethoxazole 40 mg per ml – 5% DV Aug-25 to 2028	4.95	100 ml	Deprim
	Note – this is a new Pharmacode listing, 2730847.			
106	EFAVIRENZ WITH EMTRICITABINE AND TENOFOVIR DISOPROXIL (new listing) → Tab 600 mg with emtricitabine 200 mg and tenofovir disoproxil 245 mg (300 mg as a fumarate)	106.88	30	TEEVIR (India)

NERVOUS SYSTEM

127	PARACETAMOL (brand change and addition of PSS) Oral liq 120 mg per 5 ml – 5% DV Nov-26 to 2028	3.18	200 ml	Noumed
	Note – Paracetamol (Ethics) oral liq 120 mg per 5 ml to be delisted from 1 November 2026.			
134	RIZATRIPTAN (new listing) Tab orodispersible 10 mg	4.84	30	Rizatriptan Viatrix
135	CYCLIZINE LACTATE († price) Inj 50 mg per ml, 1 ml ampoule	17.86	10	Hameln
135	DROPERIDOL (new listing) Inj 2.5 mg per ml, 1 ml ampoule	43.85	10	Droperidol Panpharma
137	RISPERIDONE (new listing) Tab 0.5 mg.....	4.01	60	Risperidone Sandoz
	Tab 2 mg.....	5.38	60	Risperidone Sandoz

ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

153	DOXORUBICIN HYDROCHLORIDE (new listing) Inj 2 mg per ml, 100 ml vial	164.50	1	Adriamycin
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Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 June 2026 (continued)

160	VENETOCLAX (amended restriction criteria – affected criteria shown only)			
	→ Tab 14 × 10 mg, 7 × 50 mg, 21 × 100 mg.....	1,771.86	42	Venclexta
	→ Tab 10 mg.....	13.68	2	Venclexta
	→ Tab 50 mg.....	239.44	7	Venclexta
	→ Tab 100 mg.....	8,209.41	120	Venclexta

Restricted

Initiation – previously untreated chronic lymphocytic leukaemia in combination with obinutuzumab

Either:

- 1 Individual is currently on treatment with venetoclax and obinutuzumab and met all of the following criteria prior to commencing treatment; or
- 2 All of the following:
 - 2.1 Individual has previously untreated chronic lymphocytic leukaemia; and
 - 2.2 Venetoclax is to be administered with obinutuzumab; and
 - 2.3 Venetoclax is to be used to a maximum dose of 400 mg and for a total of 12 (28 day) cycles*.

Note: *maximum number of cycles refers to 12 cycles of full dose venetoclax, in addition to the initial 5-week dose ramp-up period.

Initiation – previously untreated chronic lymphocytic leukaemia in combination with ibrutinib

Either:

- 1 Individual is currently on treatment with venetoclax and/or ibrutinib and met all of the following criteria prior to commencing treatment; or
- 2 All of the following:
 - 2.1 Individual has previously untreated chronic lymphocytic leukaemia; and
 - 2.2 Venetoclax is to be administered in combination with ibrutinib; and
 - 2.3 Venetoclax is to be used to a maximum dose of 400 mg and for a total of 12 (28 day) cycles*.

Note: *maximum number of cycles refers to 12 cycles of full dose venetoclax, in addition to the initial 5-week dose ramp-up period.

170	RIBOCICLIB (new listing)			
	→ Tab 200 mg.....	1,883.00	21	Kisqali
		3,767.00	42	Kisqali
		5,650.00	63	Kisqali

Note – These are new Pharmacode listings 2728133 (21 pack), 2728141 (42 pack) and 2728168 (63 pack)

RESPIRATORY SYSTEM AND ALLERGIES

270	MONTELUKAST (amended brand name)			
	Tab 5 mg.....	3.10	28	Montelukast Paediatric (Relonchem)

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 June 2026 (continued)

272 ELEXACAFTOR WITH TEZACAFTOR, IVACAFTOR AND IVACAFTOR (amended restriction criteria)

→ Tab elexacaftor 50 mg with tezacaftor 25 mg, ivacaftor 37.5 mg (56) and ivacaftor 75 mg (28)	27,647.39	84	Trikafta
→ Tab elexacaftor 100 mg with tezacaftor 50 mg, ivacaftor 75 mg (56) and ivacaftor 150 mg (28)	27,647.39	84	Trikafta
→ Oral granules elexacaftor 80 mg with tezacaftor 40 mg, ivacaftor 60 mg (28) and ivacaftor 59.5mg (28), sachets	27,647.39	56	Trikafta
→ Oral granules elexacaftor 100 mg with tezacaftor 50 mg, ivacaftor 75 mg (28) and ivacaftor 75 mg (28), sachets	27,647.39	56	Trikafta

Restricted

Initiation

All of the following:

1 Patient has been diagnosed with cystic fibrosis; and

2 Either:

2.1 Patient has two cystic fibrosis-causing mutations in the cystic fibrosis transmembrane regulator (CFTR) gene (one from each parental allele); or

2.2 Patient has a sweat chloride value of at least 60 mmol/L; and

3 Either:

3.1 Patient has a heterozygous or homozygous F508del mutation; or

3.2 Patient has a mutation responsive to elexacaftor/tezacaftor/ivacaftor (see note); and

4 The treatment must be the sole funded CFTR modulator therapy for this condition; and

5 Treatment with elexacaftor/tezacaftor/ivacaftor must be given concomitantly with standard therapy for this condition.

Note: Eligible mutations are listed in the Food and Drug Administration (FDA) Trikafta prescribing information

https://www.accessdata.fda.gov/drugsatfda_docs/label/2025/212273s015lbl.pdf

https://www.accessdata.fda.gov/drugsatfda_docs/label/2026/212273s014lbl.pdf

272 IVACAFTOR (amended restriction criteria)

→ Tab 150 mg	29,386.00	56	Kalydeco
→ Oral granules 13.4 mg, sachet	29,386.00	56	Kalydeco
→ Oral granules 25 mg, sachet	29,386.00	56	Kalydeco
→ Oral granules 50 mg, sachet	29,386.00	56	Kalydeco
→ Oral granules 75 mg, sachet	29,386.00	56	Kalydeco

Restricted

Initiation

All of the following:

1 Patient has been diagnosed with cystic fibrosis; and

2 Either:

2.1 Patient has two cystic fibrosis-causing mutations in the cystic fibrosis transmembrane regulator (CFTR) gene (one from each parental allele); or

2.2 Patients must have a sweat chloride value of at least 60 mmol/L; and

3 ~~Patient must have at least one mutation on the list of CFTR mutations that produce CFTR protein and are known to be responsive to ivacaftor**;~~ and **Patient has a mutation responsive to ivacaftor (see note); and**

4 Treatment with ivacaftor must be given concomitantly with standard therapy for this condition; and

5 The dose of ivacaftor will not exceed one tablet or one sachet twice daily.

Note:** Mutations listed in Table 3 of the Food and Drug Administration (FDA) Ivacaftor prescribing information

https://www.accessdata.fda.gov/drugsatfda_docs/label/2023/203188s038lbl.pdf

Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 June 2026 (continued)

273	VANZACAFTOR WITH TEZACAFTOR AND DEUTIVACAFTOR (amended restriction criteria)			
	→ Tab vanzacaftor 4 mg with tezacaftor 20 mg and deutivacaftor 50 mg	29,029.76	84	Alyftrek
	→ Tab vanzacaftor 10 mg with tezacaftor 50 mg and deutivacaftor 125 mg	29,029.76	56	Alyftrek

Restricted

Initiation

All of the following:

- 1 Patient has been diagnosed with cystic fibrosis; and
- 2 Either:
 - 2.1 Patient has two cystic fibrosis-causing mutations in the cystic fibrosis transmembrane regulator (CFTR) gene (one from each parental allele); or
 - 2.2 Patient has a sweat chloride value of at least 60 mmol/L; and
- 3 Either:
 - 3.1 Patient has a heterozygous or homozygous F508del mutation; or
 - 3.2 Patient has a mutation responsive to vanzacaftor/tezacaftor/deutivacaftor (see note); and
- 4 The treatment must be the sole funded CFTR modulator therapy for this condition; and
- 5 Treatment with vanzacaftor/tezacaftor/deutivacaftor must be given concomitantly with standard therapy for this condition.

Note: Eligible mutations are listed in the in the Food and Drug Administration (FDA) Alyftrek prescribing information

https://www.accessdata.fda.gov/drugsatfda_docs/label/2025/218730s002lbl.pdf

https://www.accessdata.fda.gov/drugsatfda_docs/label/2026/218730s001lbl.pdf

VACCINES

321	ROTAVIRUS ORAL VACCINE (delisted)			
	→ Oral susp live attenuated human rotavirus 1,000,000 CCID50 per dose, squeezable tube	0.00	10	Rotarix
	Note – Rotarix oral susp live attenuated human rotavirus 1,000,000 CCID50 per dose, squeezable tube delisted 1 June 2026.			

OPTIONAL PHARMACEUTICALS

323	BETA-HCG LOW SENSITIVITY URINE TEST KIT (↑ price)			
	Note: For use in abortion services only.			
	Midstream	17.47	1 test	CheckTop

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 May 2026

CARDIOVASCULAR SYSTEM

56	HYDRALAZINE HYDROCHLORIDE (new listing) Inj 20 mg per ml, 1 ml vial	283.00	25	Eugia
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MUSCULOSKELETAL SYSTEM

121	NAPROXEN (new listing) Tab 250 mg.....	3.92	50	Inza
	Tab 500 mg.....	6.89	50	Inza

125	LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE (new listing) Spray 10% – 5% DV Feb-26 to 2028	82.90	50 ml	Xylocaine
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NERVOUS SYSTEM

137	OLANZAPINE (new listing) Tab orodispersible 5 mg	3.93	28	APO-Olanzapine
	Tab orodispersible 10 mg	4.49	28	APO-Olanzapine

ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

152	THIOTEPA (new listing) Inj 100 mg vial.....	1,800.00	1	Thiotepa-Reach
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Te Kāwanatanga o Aotearoa [New Zealand Government](#)

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