

Pharmaceutical Management Agency
New Zealand
Pharmaceutical Schedule

Section H Update

for Hospital Pharmaceuticals

February 2022

The logo for PHARMAC, featuring the word "PHARMAC" in a large, bold, sans-serif font, with "TE PĀTAKA WHAIORANGA" in a smaller, all-caps, sans-serif font below it. The text is centered within a white circle that overlaps a background of stylized, wavy, concentric lines in shades of gray and white.

PHARMAC
TE PĀTAKA WHAIORANGA

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Summary of decisions

EFFECTIVE 1 FEBRUARY 2022

- Baricitinib (Olumiant) tab 2 mg and 4 mg – new listing
 - Betahistine dihydrochloride (Serc) tab 16 mg – new listing and transfer of HSS
 - Betahistine dihydrochloride (Vergo 16) tab 16 mg – removal of HSS and to be delisted 1 July 2022
 - Bortezomib inj 2.5 mg vial – presentation to be delisted 1 August 2022
 - Calcium lactate gluconate with calcium carbonate (e.g. Calcium-Sandoz Forte) tab eff 2.94 g with calcium carbonate 0.3 g (500 mg elemental) – chemical name change
 - Casirivimab and imdevimab (Ronapreve) inj 120 mg per ml casirivimab, 11.1 ml vial (1) and inj 120 mg per ml imdevimab, 11.1 ml vial (1), 1 treatment pack – new listing
 - Clonidine hydrochloride (Clonidine Teva) tab 25 mcg – new listing
 - Megestrol acetate (Megace) tab 160 mg – new listing and to be delisted 1 February 2023
 - Mivacurium chloride (Mivacron) inj 2 mg per ml, 5 ml ampoule – brand to be delisted 1 August 2022
 - Mivacurium chloride (Mivacron) inj 2 mg per ml, 10 ml ampoule – brand delisted 1 February 2022
 - Oxycodone hydrochloride (HamelN) inj 10 mg per ml, 1 ml and 2 ml ampoule and inj 50 mg per ml, 1 ml ampoule – new listing and addition of PSS
 - Oxycodone hydrochloride (Oxynorm) inj 10 mg per ml, 1 ml and 2 ml ampoule and inj 50 mg per ml, 1 ml ampoule – to be delisted 1 July 2022
 - Oxazepam (Ox-Pam) tab 10 mg and 15 mg – brand to be delisted 1 April 2022
 - Nicotine oral spray 1 mg per dose (e.g. Nicorette QuickMist Mouth Spray) and soln for inhalation 15 mg cartridge (e.g. Nicorette Inhalator) – amended restriction criteria
 - Paediatric oral feed 1 kcal/ml (Pediasure (chocolate), Pediasure (Strawberry) and Pediasure (Vanilla)) liquid 2.8 g protein, 11.2 g carbohydrate and 5 g fat per 100 ml, bottle – presentation description change
 - Paediatric oral feed 1 kcal/ml (Pediasure (Vanilla)) liquid 2.8 g protein, 11.2 g carbohydrate and 5 g fat per 100 ml, can – presentation description change
 - Paediatric oral feed 1.5 kcal/ml (e.g. Pediasure Plus) liquid 4.2 g, 16.7 g carbohydrate and 7.5 g fat per 100 ml, 500 ml bottle – new listing
 - Sunitinib (Sunitinib Pfizer) cap 12.5 mg, 25 mg and 50 mg – new listing and addition of PSS
 - Sunitinib (Sutent) cap 12.5 mg, 25 mg and 50 mg – to be delisted 1 July 2022
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Summary of decisions – effective 1 February 2022 (continued)

- Tocilizumab (Actemra) inj 20 mg per ml, 4 ml vial, 10 ml vial and 20 ml vial
– amended restriction criteria

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

Effective 1 February 2022

ALIMENTARY TRACT AND METABOLISM

20	CALCIUM LACTATE GLUCONATE WITH CALCIUM CARBONATE (chemical name change) Tab eff 2.94 g with calcium carbonate 0.3 g (500 mg elemental)			<i>e.g. Calcium-Sandoz Forte</i>
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CARDIOVASCULAR SYSTEM

47	CLONIDINE HYDROCHLORIDE (new listing) Tab 25 mcg.....	36.50	112	Clonidine Teva
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MUSCULOSKELETAL SYSTEM

106	MIVACURIUM CHLORIDE (brand delisting) Inj 2 mg per ml, 5 ml ampoule	33.92	5	Mivacron
	Inj 2 mg per ml, 10 ml ampoule	67.17	5	Mivacron
	Note – Mivacron inj 2 mg per ml, 5 ml ampoule to be delisted on 1 August 2022 and 10 ml ampoule delisted 1 February 2022 (brand only).			

NERVOUS SYSTEM

109	BROMOCRIPTINE (delist delayed) → Tab 2.5 mg – Restricted: For continuation only Note – this delist delayed from 1 March 2022 to 1 September 2022.			
116	XYCODONE HYDROCHLORIDE (brand change and addition of PSS) Inj 10 mg per ml, 1 ml ampoule – 5% DV Jul-22 to 2024	5.82	5	Hameln
	Inj 10 mg per ml, 2 ml ampoule – 5% DV Jul-22 to 2024	11.49	5	Hameln
	Inj 50 mg per ml, 1 ml ampoule – 5% DV Jul-22 to 2024	22.92	5	Hameln
	Note – OxyNorm inj 10 mg per ml, 1 ml ampoule, 2 ml ampoule and inj 50 mg per ml, 1 ml ampoule to be delisted 1 July 2022.			
122	BETAHISTINE DIHYDROCHLORIDE (brand change and transfer of HSS) Tab 16 mg – 1% DV Feb-22 to 2023	4.62	100	Serc
	Note – Vergo 16 tab 16 mg to be delisted 1 July 2022.			
127	OXAZEPAM (brand delisting) Tab 10 mg.....	6.17	100	Ox-Pam
	Tab 15 mg.....	8.53	100	Ox-Pam
	Note – Ox-Pam tab 10 mg and 15 mg brand to be delisted 1 April 2022.			

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 February 2022 (continued)

133	NICOTINE (amended restriction criteria) → Oral spray 1 mg per dose			<i>e.g. Nicorette QuickMist Mouth Spray e.g. Nicorette Inhalator</i>
	→ Soln for inhalation 15 mg cartridge			
	Restricted Initiation Any of the following:			
	1 For perioperative use in patients who have a 'nil by mouth' instruction; or			
	2 For use within mental health inpatient units; or			
	3 Patient would be admitted to a mental health inpatient unit, but is unable to due to COVID-19 self-isolation requirement; or			
	34 For acute use in agitated patients who are unable to leave the hospital facilities.			

ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

138	BORTEZOMIB (presentation delisting) → Inj 2.5 mg vial Note – Bortezomib inj 2.5 mg vial presentation to be delisted 1 August 2022.			
149	SUNITINIB (brand change and addition of PSS) → Cap 12.5 mg – 5% DV Jul-22 to 2024..... 208.38	28		Sunitinib Pfizer
	→ Cap 25 mg – 5% DV Jul-22 to 2024..... 416.77	28		Sunitinib Pfizer
	→ Cap 50 mg – 5% DV Jul-22 to 2024..... 694.62	28		Sunitinib Pfizer
	Note – Sutent cap 12.5 mg, 25 mg and 50 mg to be delisted 1 July 2022.			
152	MEGESTROL ACETATE – Restricted: For continuation only (new listing) → Tab 160 mg..... 48.80	30		Megace
	Note – Megace tab 16 mg to be delisted 1 February 2023.			

	Price (ex man. Excl. GST) \$ Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 February 2022 (continued)

174	CASIRIVIMAB AND IMDEVIMAB (new listing) → Inj 120 mg per ml casirivimab, 11.1 ml vial (1) and inj 120 mg per ml imdevimab, 11.1 ml vial (1).....	0.00	1	Ronapreve
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Restricted

Initiation – mild to moderate COVID-19-hospitalised patients

Limited to 2 weeks

All of the following:

- 1 Patient has confirmed (or probable) COVID-19; and
- 2 Patient is an in-patient in hospital with mild to moderate disease severity*; and
- 3 Patient's symptoms started within the last 10 days; and
- 4 Patient is not receiving high flow oxygen or assisted/mechanical ventilation; and
- 5 Any of the following:
 - 5.1 Age > 50; or
 - 5.2 BMI > 30; or
 - 5.3 Patient is Māori or Pacific ethnicity; or
 - 5.4 Patient is at increased risk of severe illness from COVID-19, excluding pregnancy, as described on the Ministry of Health website (see Notes); and
- 6 Either:
 - 6.1 Patient is unvaccinated; or
 - 6.2 Patient is seronegative where serology testing is readily available or strongly suspected to be seronegative where serology testing is not readily available; and
- 7 Casirivimab and imdevimab is to be administered at a maximum dose of no greater than 2,400 mg.

Note:* Mild to moderate disease severity as described on the [Ministry of Health Website](#)

** (<https://www.health.govt.nz/our-work/diseases-and-conditions/covid-19-novel-coronavirus/covid-19-information-specific-audiences/covid-19-advice-higher-risk-people>)

Initiation – Treatment of profoundly immunocompromised patients

Limited to 2 weeks

All of the following:

- 1 Patient has confirmed (or probable) COVID-19; and
- 2 The patient is in the community (treated as an outpatient) with mild to moderate disease severity*; and
- 3 Patient is profoundly immunocompromised** and is at risk of not having mounted an adequate response to vaccination against COVID-19 or is unvaccinated; and
- 4 Patient's symptoms started within the last 10 days; and
- 5 Patient is not receiving high flow oxygen or assisted/mechanical ventilation; and
- 6 Casirivimab and imdevimab is to be administered at a maximum dose of no greater than 2,400 mg.

Note:* Mild to moderate disease severity as described on the [Ministry of Health Website](#)

** Examples include B-cell depletive illnesses or patients receiving treatment that is B-Cell depleting.

	Price (ex man. Excl. GST) \$	Per	Brand or Generic Manufacturer
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Changes to Section H Part II – effective 1 February 2022 (continued)

200	TOCILIZUMAB (amended restriction criteria – affected criteria shown only)			
	→ Inj 20 mg per ml, 4 ml vial	220.00	1	Actemra
	→ Inj 20 mg per ml, 10 ml vial	550.00	1	Actemra
	→ Inj 20 mg per ml, 20 ml vial	1,100.00	1	Actemra

Restricted

Indication – moderate to severe COVID-19*

Therapy limited to 1 dose

All of the following:

- 1 Patient has confirmed (or probable) COVID-19; and
- 2 Oxygen saturation of <92% on room air, or requiring supplemental oxygen; and
- ~~3 Patient has significantly increased laboratory markers of systemic inflammation (eg CRP, PCT or ferritin); and~~
- ~~34 Patient is receiving adjunct systemic corticosteroids, or systemic corticosteroids are contraindicated; and~~
- ~~45 Tocilizumab is to be administered at doses no greater than 8mg/kg IV for a maximum of one dose; and~~
- 5 Tocilizumab is not to be administered in combination with baricitinib.**

Note: Indications marked with * are unapproved indications.

211	BARICITINIB (new listing)			
	→ Tab 2 mg.....	0.00	28	Olumiant
	→ Tab 4 mg.....	0.00	28	Olumiant

Restricted

Initiation – moderate to severe COVID-19*

Limited to 14 days treatment

All of the following:

- 1 Patient has confirmed (or probable) COVID-19*; and
- 2 Oxygen saturation of <92% on room air, or requiring supplemental oxygen; and
- 3 Patient is receiving adjunct systemic corticosteroids, or systemic corticosteroids are contraindicated; and
- 4 Baricitinib is to be administered at doses no greater than 4 mg daily for up to 14 days; and
- 5 Baricitinib is not to be administered in combination with tocilizumab.

Note: Indications marked with * are unapproved indications.

SPECIAL FOODS

251	PAEDIATRIC ORAL FEED 1 KCAL/ML (presentation description change)			
	→ Liquid 4-22.8 g protein, +6-711.2 g carbohydrate and 7-55 g fat per 100 ml, bottle	1.07	200 ml	Pediasure (Chocolate) Pediasure (Strawberry) Pediasure (Vanilla)
	→ Liquid 4-22.8 g protein, +6-711.2 g carbohydrate and 7-55 g fat per 100 ml, can	1.34	250 ml	Pediasure (Vanilla)
251	PAEDIATRIC ORAL FEED 1.5 KCAL/ML (new listing)			
	→ Liquid 4.2 g protein, 16.7 g carbohydrate and 7.5 g fat per 100 ml, 500 ml bottle			<i>e.g. Pediasure Plus</i>

→ Restriction

(Brand) indicates a brand example only. It is not a contracted product.

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New Zealand
Permit No. 478



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