

Pharmaceutical Management Agency  
New Zealand  
Pharmaceutical Schedule

# Section H Update

for Hospital Pharmaceuticals

January 2022

The logo for PHARMAC, featuring the word "PHARMAC" in a bold, uppercase, sans-serif font, with "TE PĀTAKA WHAIORANGA" in a smaller, uppercase, sans-serif font below it. The logo is centered within a white circle that overlaps a large, stylized graphic of white and grey wavy lines on a grey background.

PHARMAC  
TE PĀTAKA WHAIORANGA

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## Summary of decisions

EFFECTIVE 1 JANUARY 2022

- Aflibercept (Eylea) inj 40 mg per ml, 0.1 ml vial – amended restriction criteria
- Bisacodyl (Pharmacy Health) tab 5 mg – new listing and addition of PSS
- Bisacodyl (Lax-tabs) tab 5 mg – to be delisted 1 June 2022
- Bupivacaine hydrochloride with adrenaline inj 2.5 mg with adrenaline 1:200,000, 10 ml ampoule – new presentation listing
- Bupivacaine hydrochloride with fentanyl (Bupafen NRFit) inj 1.25 mg with fentanyl 2 mcg per ml, 15 ml syringe – delisted 1 January 2022
- Colchicine (Colgout) tab 500 mcg – price increase
- Corticorelin inj 100 mcg vial – chemical name change
- Docusate sodium with sennosides (Laxsol) tab 50 mg with sennosides 8 mg – price increase
- Doxazosin (Doxazosin Clinect) tab 2 mg and 4 mg – new listing
- Doxazosin (Apo-Doxazosin) tab 2 mg and 4 mg – to be delisted 1 September 2022
- Eplerenone (Inspra) tab 25 mg and 50 mg – price increase and addition of PSS
- Gelatine, succinylated (Gelifusine) inj 4%, 500 ml bag – price increase
- Glycine (B Braun) irrigation soln 1.5%, 3,000 ml bag – price increase
- Hydroxychloroquine (Plaquenil) tab 200 mg – price increase
- Lidocaine [lignocaine] hydrochloride with adrenaline inj 1% with adrenaline 1:1,000,000, 20 ml vial – new presentation listing
- Lisinopril (Ethics Lisinopril) tab 5 mg, 10 mg and 20 mg – price increase
- Losartan potassium with hydrochlorothiazide (Arrow-Losartan & Hydrochlorothiazide) tab 50 mg with hydrochlorothiazide 12.5 mg – price increase
- Megestrol acetate (Apo-Megestrol) tab 160 mg – to be delisted 1 May 2022
- Octreotide (Max Health) inj 50 mcg per ml, 100 mcg per ml, 500 mcg per ml, 1 ml ampoule – new listing and addition of PSS
- Octreotide (DBL Octreotide) inj 50 mcg per ml, 100 mcg per ml, 500 mcg per ml, 1 ml ampoule – to be delisted 1 June 2022
- Oxybutynin (Alchemy Oxybutynin) tab 5 mg – new listing
- Oxycodone hydrochloride (Oxycodone Sandoz) tab controlled-release 5 mg, 10 mg, 20 mg, 40 mg and 80 mg – price increase and addition of PSS

## Summary of decisions – effective 1 January 2022 (continued)

- Pancreatic enzyme cap pancreatin 150 mg (amylase 8,000 Ph Eur U, lipase 10,000 Ph Eur U, total protease 600 Ph Eur U) (Creon 10000) and cap pancreatin 300 mg (amylase 18,000 Ph Eur U, lipase 25,000 Ph Eur U, total protease 1,000 Ph Eur U) (Creon 25000) – addition of PSS
- Perindopril (Coversyl) tab 2 mg and 4 mg – new listing and addition of PSS
- Phenoxymethylpenicillin [Penicillin V] (Cilicaine VK) cap 250 mg (Pharmacode 2602865) – new Pharmacode listing
- Phenoxymethylpenicillin [Penicillin V] (Cilicaine VK) cap 250 mg (Pharmacode 2048841) – Pharmacode to be delisted 1 July 2022
- Pneumococcal (PCV13) conjugate vaccine (Prevenar 13) Inj 30.8 mcg of pneumococcal polysaccharide serotypes 1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A, 19F and 23F in 0.5 ml syringe, 1 and 10 inj pack – amended restriction criteria
- Ranibizumab inj 10 mg per ml, 0.23 ml vial and 0.3 ml vial – amended restriction criteria
- Sodium chloride (B Braun) irrigation soln 0.9%, 3,000 ml bag – price increase
- Sodium chloride (BD PosiFlush) inj 0.9%, 3 ml, 5 ml and 10 ml syringe, non-sterile pack – price increase
- Teicoplanin (Targocid) inj 400 mg vial – new listing and addition of PSS
- Teicoplanin (Teicoplanin Mylan) inj 400 mg vial – to be delisted 1 June 2022
- Water (B Braun) irrigation soln, 3,000 ml bag – price increase

|  |  | Price<br>(ex man. Excl. GST)<br>\$ Per | Brand or<br>Generic<br>Manufacturer |
|--|--|--|-------------------------------------|
|--|--|--|-------------------------------------|

## Section H changes to Part II

Effective 1 January 2022

### ALIMENTARY TRACT AND METABOLISM

|    |  |       |     |                        |
|----|--|-------|-----|------------------------|
| 12 | PANCREATIC ENZYME (addition of PSS)<br>Cap pancreatin 150 mg (amylase 8,000 Ph Eur U,<br>lipase 10,000 Ph Eur U,<br>total protease 600 Ph Eur U) – <b>5% DV Jun-22 to 2024</b> ..... | 34.93 | 100 | <b>Creon 10000</b>     |
|    | Cap pancreatin 300 mg (amylase 18,000 Ph Eur U,<br>lipase 25,000 Ph Eur U,<br>total protease 1,000 Ph Eur U) – <b>5% DV Jun-22 to 2024</b> ....                                      | 94.38 | 100 | <b>Creon 25000</b>     |
| 14 | DOCUSATE SODIUM WITH SENNOSIDES (↑ price)<br>Tab 50 mg with sennosides 8 mg .....  | 4.20  | 200 | Laxsol                 |
| 15 | BISACODYL (brand change and addition of PSS)<br>Tab 5 mg – <b>5% DV Jun-22 to 2024</b> .....   | 5.80  | 200 | <b>Pharmacy Health</b> |
|    | Note – Lax-Tabs tab 5 mg to be delisted 1 June 2022.   |       |     |                        |

### BLOOD AND BLOOD FORMING ORGANS

|    |  |        |     |              |
|----|--|--------|-----|--------------|
| 40 | SODIUM CHLORIDE (↑ price)<br>→ Inj 0.9%, 3 ml syringe, non-sterile pack..... | 168.00 | 480 | BD PosiFlush |
|    | → Inj 0.9%, 5 ml syringe, non-sterile pack.....                              | 169.92 | 480 | BD PosiFlush |
|    | → Inj 0.9%, 10 ml syringe, non-sterile pack.....                             | 177.60 | 480 | BD PosiFlush |

### CARDIOVASCULAR SYSTEM

|    |  |        |     |   |
|----|--|--------|-----|---|
| 40 | GELATINE, SUCCINYLATED (↑ price)<br>Inj 4%, 500 ml bag .....   | 129.00 | 10  | Gelofusine                              |
| 42 | LISINAPRIL (↑ price)<br>Tab 5 mg.....  | 17.50  | 90  | Ethics Lisinopril                       |
|    | Tab 10 mg.....   | 17.50  | 90  | Ethics Lisinopril                       |
|    | Tab 20 mg.....   | 17.50  | 90  | Ethics Lisinopril                       |
| 42 | PERINDOPRIL (new listing and addition of PSS)<br>Tab 2 mg – <b>5% DV Jan-22 to 2024</b> .....            | 1.58   | 30  | <b>Coversyl</b>                         |
|    | Tab 4 mg – <b>5% DV Jan-22 to 2024</b> .....   | 2.95   | 30  | <b>Coversyl</b>                         |
| 43 | LOSARTAN POTASSIUM WITH HYDROCHLOROTHIAZIDE (↑ price)<br>Tab 50 mg with hydrochlorothiazide 12.5 mg..... | 15.25  | 30  | Arrow-Losartan &<br>Hydrochlorothiazide |
| 43 | DOXAZOSIN (brand change)<br>Tab 2 mg.....  | 17.35  | 500 | Doxazosin Clinect                       |
|    | Tab 4 mg.....  | 20.94  | 500 | Doxazosin Clinect                       |
|    | Note – Apo-Doxazosin tab 2 mg and 4 mg to be delisted 1 September 2022.                                  |        |     |   |

|  | Price<br>(ex man. Excl. GST)<br>\$ | Per | Brand or<br>Generic<br>Manufacturer |
|--|------------------------------------|-----|-------------------------------------|
|--|------------------------------------|-----|-------------------------------------|

## Changes to Section H Part II – effective 1 January 2022 (continued)

|    |  |       |                |
|----|--|-------|----------------|
| 48 | EPLERENONE (↑ price and addition of PSS) |       |                |
|    | → Tab 25 mg – 5% DV Jun-22 to 2024 ..... | 18.50 | 30             |
|    | → Tab 50 mg – 5% DV Jun-22 to 2024 ..... | 25.00 | 30             |
|    |  |       | <b>Inspira</b> |
|    |  |       | <b>Inspira</b> |

## GENITO-URINARY SYSTEM

|    |  |      |                    |
|----|--|------|--------------------|
| 66 | OXYBUTYNIN – Restricted: For continuation only (new listing) |      |                    |
|    | → Tab 5 mg.....  | 5.42 | 100                |
|    |  |      | Alchemy Oxybutynin |

## HORMONE PREPARATIONS

|    |   |  |  |
|----|---|--|--|
| 71 | <del>CORTICORELIN</del> <del>CORTICOTRORELIN</del> (OVINE) (chemical name change) |  |  |
|    | Inj 100 mcg vial  |  |  |

## INFECTIONS

|    |   |      |                     |
|----|---|------|---------------------|
| 81 | PHENOXYMETHYLPENICILLIN [PENICILLIN V] (Pharmacode change)                                    |      |                     |
|    | Cap 250 mg – 5% DV Jan-22 to 2024 .....   | 3.84 | 50                  |
|    | Note – this listing is for Pharmacode 2602865. Pharmacode 2048841 to be delisted 1 July 2022. |      | <b>Cilicaine VK</b> |

|    |  |       |                 |
|----|--|-------|-----------------|
| 85 | TEICOPLANIN (brand change and addition of PSS)                       |       |                 |
|    | → Inj 400 mg vial – 5% DV Jun-22 to 2024 .....                       | 49.95 | 1               |
|    | Note – Teicoplanin Mylan inj 400 mg vial to be delisted 1 June 2022. |       | <b>Targocid</b> |

## MUSCULOSKELETAL SYSTEM

|     |                              |      |           |
|-----|------------------------------|------|-----------|
| 101 | HYDROXYCHLOROQUINE (↑ price) |      |           |
|     | → Tab 200 mg.....            | 8.78 | 100       |
|     |                              |      | Plaquenil |

|     |                      |       |         |
|-----|----------------------|-------|---------|
| 106 | COLCHICINE (↑ price) |       |         |
|     | Tab 500 mcg.....     | 10.06 | 100     |
|     |                      |       | Colgout |

## NERVOUS SYSTEM

|     |  |  |  |
|-----|--|--|--|
| 111 | BUPIVACAINE HYDROCHLORIDE WITH ADRENALINE (new presentation listing) |  |  |
|     | Inj 2.5 mg per ml with adrenaline 1:200,000, 10 ml ampoule           |  |  |

|     |   |       |                      |
|-----|---|-------|----------------------|
| 113 | BUPIVACAINE HYDROCHLORIDE WITH FENTANYL (delisting)   |       |                      |
|     | Inj 1.25 mg with fentanyl 2 mcg per ml, 15 ml syringe .....   | 36.00 | 5                    |
|     | Note – Bupafen NRFit inj 1.25 mg with fentanyl 2 mcg per ml, 15 ml syringe delisted 1 January 2022. |       | <b>Bupafen NRFit</b> |

|     |   |  |  |
|-----|---|--|--|
| 113 | LIDOCAINE [LIGNOCAINE] HYDROCHLORIDE WITH ADRENALINE (new presentation listing) |  |  |
|     | Inj 1% with adrenaline 1:100,000, 20 ml vial                                    |  |  |

|     |   |       |                         |
|-----|---|-------|-------------------------|
| 116 | OXYCODONE HYDROCHLORIDE (↑ price and addition of PSS)     |       |                         |
|     | Tab controlled-release 5 mg – 5% DV Jun-22 to 2024 .....  | 2.69  | 20                      |
|     | Tab controlled-release 10 mg – 5% DV Jun-22 to 2024 ..... | 2.69  | 20                      |
|     | Tab controlled-release 20 mg – 5% DV Jun-22 to 2024 ..... | 3.49  | 20                      |
|     | Tab controlled-release 40 mg – 5% DV Jun-22 to 2024 ..... | 5.49  | 20                      |
|     | Tab controlled-release 80 mg – 5% DV Jun-22 to 2024 ..... | 12.99 | 20                      |
|     |   |       | <b>Oxycodone Sandoz</b> |
|     |   |       | <b>Oxycodone Sandoz</b> |
|     |   |       | <b>Oxycodone Sandoz</b> |
|     |   |       | <b>Oxycodone Sandoz</b> |
|     |   |       | <b>Oxycodone Sandoz</b> |

→ Restriction

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

|  | Price<br>(ex man. Excl. GST)<br>\$ Per | Brand or<br>Generic<br>Manufacturer |
|--|--|-------------------------------------|
|--|--|-------------------------------------|

## Changes to Section H Part II – effective 1 January 2022 (continued)

### ONCOLOGY AGENTS AND IMMUNOSUPPRESSANTS

|  |  |    |                   |
|--|--|----|-------------------|
| 152  | MEGESTROL ACETATE – Restricted: For continuation only (delisting)<br>➔ Tab 160 mg.....63.53                                  | 30 | Apo-Megestrol     |
| Note – Apo-Megestrol tab 160 mg to be delisted 1 May 2022.   |  |    |                   |
| 153  | OCTREOTIDE (brand change and addition of PSS)<br>Inj 50 mcg per ml, 1 ml ampoule<br>– <b>5% DV Jun-22 to 2024</b> .....27.58 | 5  | <b>Max Health</b> |
|  | Inj 100 mcg per ml, 1 ml ampoule<br>– <b>5% DV Jun-22 to 2024</b> .....32.71   | 5  | <b>Max Health</b> |
|  | Inj 500 mcg per ml, 1 ml ampoule<br>– <b>5% DV Jun-22 to 2024</b> .....113.10  | 5  | <b>Max Health</b> |
| Note – DBL Octreotide inj 50 mcg per ml, 100 mcg per ml and 500 mcg per ml, 1 ml ampoule to be delisted 1 June 2022.                         |  |    |                   |
| 172  | AFLIBERCEPT (amended restriction criteria – amended criteria shown only)<br>➔ Inj 40 mg per ml, 0.1 ml vial .....1,250.00    | 1  | Eylea             |
| Restricted   |  |    |                   |
| Initiation - Wet Age Related Macular Degeneration  |  |    |                   |
| Ophthalmologist <b>or nurse practitioner</b>   |  |    |                   |
| <i>Re-assessment required after 3 months</i>   |  |    |                   |
| Either:  |  |    |                   |
| 1. All of the following:   |  |    |                   |
| 1.1. Any of the following:   |  |    |                   |
| 1.1.1. Wet age-related macular degeneration (wet AMD); or  |  |    |                   |
| 1.1.2. Polypoidal choroidal vasculopathy; or   |  |    |                   |
| 1.1.3. Choroidal neovascular membrane from causes other than wet AMD; and  |  |    |                   |
| 1.2. Either:   |  |    |                   |
| 1.2.1. The patient has developed severe endophthalmitis or severe posterior uveitis following treatment with bevacizumab; or                 |  |    |                   |
| 1.2.2. There is worsening of vision or failure of retina to dry despite three intraocular injections of bevacizumab four weeks apart; and    |  |    |                   |
| 1.3. There is no structural damage to the central fovea of the treated eye; and  |  |    |                   |
| 1.4. Patient has not previously been treated with ranibizumab for longer than 3 months; or   |  |    |                   |
| 2. Either:   |  |    |                   |
| 2.1. Patient has current approval to use ranibizumab for treatment of wAMD and was found to be intolerant to ranibizumab within 3 months; or |  |    |                   |
| 2.2. Patient has previously* (*before June 2018) received treatment with ranibizumab for wAMD and disease was stable while on treatment.     |  |    |                   |
| Continuation - Wet Age Related Macular Degeneration  |  |    |                   |
| Ophthalmologist <b>or nurse practitioner</b>   |  |    |                   |
| <i>Re-assessment required after 12 months</i>  |  |    |                   |
| All of the following:  |  |    |                   |
| 1. Documented benefit must be demonstrated to continue; and  |  |    |                   |
| 2. Patient's vision is 6/36 or better on the Snellen visual acuity score; and  |  |    |                   |
| 3. There is no structural damage to the central fovea of the treated eye.  |  |    |                   |

continued...

| Price<br>(ex man. Excl. GST)<br>\$ | Per | Brand or<br>Generic<br>Manufacturer |
|------------------------------------|-----|-------------------------------------|
|------------------------------------|-----|-------------------------------------|

## Changes to Section H Part II – effective 1 January 2022 (continued)

continued...

Initiation - Diabetic Macular Oedema (DMO)

Ophthalmologist **or nurse practitioner**

*Re-assessment required after 4 months*

All of the following:

1. Patient has centre involving diabetic macular oedema (DMO); and
2. Patient's disease is non responsive to 4 doses of intravitreal bevacizumab when administered 4-6 weekly; and
3. Patient has reduced visual acuity between 6/9 – 6/36 with functional awareness of reduction in vision; and
4. Patient has DMO within central OCT (ocular coherence tomography) subfield > 350 micrometers; and
5. There is no centre-involving sub-retinal fibrosis or foveal atrophy.

Continuation - Diabetic Macular Oedema (DMO)

Ophthalmologist **or nurse practitioner**

*Re-assessment required after 12 months*

All of the following:

1. There is stability or two lines of Snellen visual acuity gain; and
2. There is structural improvement on OCT scan (with reduction in intra-retinal cysts, central retinal thickness, and sub-retinal fluid); and
3. Patient's vision is 6/36 or better on the Snellen visual acuity score; and
4. There is no centre-involving sub-retinal fibrosis or foveal atrophy; and
5. After each consecutive 12 months treatment with aflibercept, patient has retrialled with at least one injection of bevacizumab and had no response.

184 RANIBIZUMAB (amended restriction criteria)

→ Inj 10 mg per ml, 0.23 ml vial

→ Inj 10 mg per ml, 0.3 ml vial

Restricted

Initiation - Wet Age Related Macular Degeneration

Ophthalmologist **or nurse practitioner**

*Re-assessment required after 3 months*

Either:

1. All of the following:
  - 1.1. Any of the following:
    - 1.1.1. Wet age-related macular degeneration (wet AMD); or
    - 1.1.2. Polypoidal choroidal vasculopathy; or
    - 1.1.3. Choroidal neovascular membrane from causes other than wet AMD; and
  - 1.2. Either:
    - 1.2.1. The patient has developed severe endophthalmitis or severe posterior uveitis following treatment with bevacizumab; or
    - 1.2.2. There is worsening of vision or failure of retina to dry despite three intraocular injections of bevacizumab four weeks apart; and
  - 1.3. There is no structural damage to the central fovea of the treated eye; and
  - 1.4. Patient has not previously been treated with aflibercept for longer than 3 months; or
2. Patient has current approval to use aflibercept for treatment of wAMD and was found to be intolerant to aflibercept within 3 months

Continuation - Wet Age Related Macular Degeneration

Ophthalmologist **or nurse practitioner**

*Re-assessment required after 12 months*

All of the following:

1. Documented benefit must be demonstrated to continue; and
2. Patient's vision is 6/36 or better on the Snellen visual acuity score; and
3. There is no structural damage to the central fovea of the treated eye.



|  | Price<br>(ex man. Excl. GST)<br>\$ | Per | Brand or<br>Generic<br>Manufacturer |
|--|------------------------------------|-----|-------------------------------------|
|--|------------------------------------|-----|-------------------------------------|

**Changes to Section H Part II – effective 1 January 2022 (continued)**

**VARIOUS**

|     |   |       |   |         |
|-----|---|-------|---|---------|
| 233 | GLYCINE († price)<br>Irrigation soln 1.5%, 3,000 ml bag .....         | 33.20 | 4 | B Braun |
| 233 | SODIUM CHLORIDE († price)<br>Irrigation soln 0.9%, 3,000 ml bag ..... | 28.80 | 4 | B Braun |
| 233 | WATER († price)<br>Irrigation soln, 3,000 ml bag .....                | 30.95 | 4 | B Braun |

**VACCINES**

|     |  |      |         |                            |
|-----|--|------|---------|----------------------------|
| 257 | PNEUMOCOCCAL (PCV13) CONJUGATE VACCINE (amended restriction criteria – amended criteria shown only)<br>→ Inj 30.8 mcg of pneumococcal polysaccharide serotypes<br>1, 3, 4, 5, 6A, 6B, 7F, 9V, 14, 18C, 19A,<br>19F and 23F in 0.5 ml syringe ..... | 0.00 | 1<br>10 | Prevenar 13<br>Prevenar 13 |
|-----|--|------|---------|----------------------------|

Restricted

Initiation – High risk adults and children 5 years and over

*Therapy limited to 4 doses*

Up to an additional four doses (as appropriate) are funded for (re-)immunisation of patients 5 years and over with HIV, for patients pre or post haematopoietic stem cell transplantation, or chemotherapy; pre- or post splenectomy; functional asplenia, pre- or post- solid organ transplant, renal dialysis, complement deficiency (acquired or inherited), cochlear implants, **intracranial shunts**, **cerebrospinal fluid leaks** or primary immunodeficiency.

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New Zealand  
Permit No. 478



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Te Kāwanatanga o Aotearoa New Zealand Government

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