

**PB 65 of 2018**

**National Health (Pharmaceutical Benefits – early supply) Amendment Instrument 2018 (No. 7)**

*National Health Act 1953*

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I, LISA LA RANCE, Assistant Secretary, Pricing and PBS Policy Branch, Technology Assessment and Access Division, Department of Health, delegate of the Minister for Health, make this Instrument under subsection 84AAA(2) of the *National Health Act 1953*.

Dated 27 JULY 2018

**LISA LA RANCE**

Assistant Secretary

Pricing and PBS Policy Branch

Technology Assessment and Access Division

Department of Health

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1. **Name of Instrument**
2. This Instrument is the *National Health (Pharmaceutical Benefits – early supply) Amendment Instrument 2018 (No. 7)*.
3. This Instrument may also be cited as PB 65 of 2018.
4. **Commencement**

This Instrument commences on 1 August 2018.

1. **Amendment of *National Health (Pharmaceutical Benefits—early supply) Instrument 2015* (PB 120 of 2015)**

Schedule 1 amends the *National Health (Pharmaceutical Benefits—early supply) Instrument 2015* (PB 120 of 2015).

**Schedule 1 Amendments**

1. **Schedule 1, after entry for Ibrutinib**

*insert:*

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| Ibrutinib | Capsule 140 mg | 20 | 120 | 5 |  |

1. **Schedule 1, entry for Levodopa with carbidopa and entacapone**
   1. omit from the column headed “Form”: Tablet 100 mg‑25 mg‑200 mg

*substitute:* Tablet 100 mg‑25 mg (as monohydrate)‑200 mg

* 1. omit from the column headed “Form”: Tablet 125 mg‑31.25 mg-200 mg

substitute: Tablet 125 mg‑31.25 mg (as monohydrate)‑200 mg

* 1. omit from the column headed “Form”: Tablet 150 mg‑37.5 mg -200 mg

substitute: Tablet 150 mg‑37.5 mg (as monohydrate)‑200 mg

* 1. omit from the column headed “Form”: Tablet 200 mg‑50 mg-200 mg

substitute: Tablet 200 mg‑50 mg (as monohydrate)‑200 mg

* 1. omit from the column headed “Form”: Tablet 50 mg‑12.5 mg-200 mg

substitute: Tablet 50 mg‑12.5 mg (as monohydrate)‑200 mg

* 1. omit from the column headed “Form”: Tablet 75 mg‑18.75 mg-200 mg

substitute: Tablet 75 mg‑18.75 mg (as monohydrate)‑200 mg

1. **Schedule 1, entry for Methyldopa**

*omit from the column headed “Form”:* Tablet 250 mg

*substitute:* Tablet 250 mg (as sesquihydrate)

1. **Schedule 1, after entry for Pirfenidone**

*insert:*

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| Pirfenidone | Tablet 267 mg | 20 | 270 | 5 |  |
| Pirfenidone | Tablet 801 mg | 20 | 90 | 5 |  |

1. **Schedule 1, entry for Pramipexole**
2. omit from the column headed “Form”: Tablet (extended release) containing pramipexole hydrochloride 1.5 mg

*substitute:* Tablet (extended release) containing pramipexole dihydrochloride monohydrate 1.5 mg

1. omit from the column headed “Form”: Tablet (extended release) containing pramipexole hydrochloride 2.25 mg

substitute: Tablet (extended release) containing pramipexole dihydrochloride monohydrate 2.25 mg

1. omit from the column headed “Form”: Tablet (extended release) containing pramipexole hydrochloride 3 mg

substitute: Tablet (extended release) containing pramipexole dihydrochloride monohydrate 3 mg

1. omit from the column headed “Form”: Tablet (extended release) containing pramipexole hydrochloride 3.75 mg

substitute: Tablet (extended release) containing pramipexole dihydrochloride monohydrate 3.75 mg

1. omit from the column headed “Form”: Tablet (extended release) containing pramipexole hydrochloride 4.5 mg

substitute: Tablet (extended release) containing pramipexole dihydrochloride monohydrate 4.5 mg