

PB 65 of 2018

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

National Health Act 1953

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

#### 1 Name of Instrument

- (1) This Instrument is the National Health (Pharmaceutical Benefits early supply) Amendment Instrument 2018 (No. 7).
- (2) This Instrument may also be cited as PB 65 of 2018.

#### 2 Commencement

This Instrument commences on 1 August 2018.

### 3 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

Ibrutin	nib Capsule 140 mg	20	120	5		
Sch	Schedule 1, entry for Levodopa with carbidopa and entacapone					
(a)	omit from the column headed "Form": substitute:	Tablet 100 mg-25 mg-200 mg Tablet 100 mg-25 mg (as monohydrate)-200 mg				
(b)	omit from the column headed "Form": substitute:	Tablet 125 mg-31.25 mg-200 mg Tablet 125 mg-31.25 mg (as monohydrate)-200 mg				
(c)	omit from the column headed "Form": substitute:	Tablet 150 mg-37.5 mg -200 mg Tablet 150 mg-37.5 mg (as monohydrate)-200 mg				
(d)	omit from the column headed "Form": substitute:	Tablet 200 mg-50 mg-200 mg Tablet 200 mg-50 mg (as monohydrate)-200 mg				
(e)	omit from the column headed "Form": substitute:	Tablet 50 mg-12.5 mg-200 mg Tablet 50 mg-12.5 mg (as monohydrate)-200 mg				
(f)	omit from the column headed "Form": substitute:	Tablet 75 mg-18.75 mg-200 mg Tablet 75 mg-18.75 mg (as monohydrate)-200 mg				
Schedule 1, entry for Methyldopa						
omit from the column headed "Form": Tablet 250 mg substitute: Tablet 250 mg (as sesquihydrate)						
Schedule 1, after entry for Pirfenidone						
inser	nsert:					
Pirfen	hidone Tablet 267 mg	20	270	5		
Pirfen	nidone Tablet 801 mg	20	90	5		

(a) omit from the column headed "Form": substitute:
(b) omit from the column headed "Form": substitute:
(c) Tablet (extended release) containing pramipexole dihydrochloride 1.5 mg
(c) Tablet (extended release) containing pramipexole dihydrochloride 2.25 mg
(c) Tablet (extended release) containing pramipexole dihydrochloride 2.25 mg
(c) Tablet (extended release) containing pramipexole dihydrochloride 2.25 mg
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(c)	omit from the column headed "Form": substitute:	Tablet (extended release) containing pramipexole hydrochloride 3 mg Tablet (extended release) containing pramipexole dihydrochloride monohydrate 3 mg
(d)	omit from the column headed "Form": substitute:	Tablet (extended release) containing pramipexole hydrochloride 3.75 mg Tablet (extended release) containing pramipexole dihydrochloride monohydrate 3.75 mg
(e)	omit from the column headed "Form": substitute:	Tablet (extended release) containing pramipexole hydrochloride 4.5 mg Tablet (extended release) containing pramipexole dihydrochloride monohydrate 4.5 mg