**EXPLANATORY STATEMENT**

***National Health Act 1953***

***National Health (Originator Brand) Amendment Determination 2020 (No. 1)***

**PB 9 of 2020**

**Authority**

This legislative instrument is made pursuant to subsection 99ADB(6B) of the *National Health Act 1953* (the Act).

**Purpose**

This legislative instrument amends the *National Health (Originator Brand) Determination 2015* (PB 100 of 2015) (the Principal Instrument) to determine the originator brand of a pharmaceutical item for two drugs new to the Pharmaceutical Benefits Scheme (PBS) F2 formulary on 1 February 2020.

The Principal Instrument determines originator brands of pharmaceutical items that have a drug on the F2 formulary. On meeting certain criteria, drugs move from the F1 formulary (see s 85AB of the Act), or the single brand Combination Drug List (CDL), to F2. All drugs on F2 are subject to price disclosure. This instrument is necessary to implement removal of originator brand data from price disclosure calculations (originator removal) in certain circumstances as set out in the *National Health (Pharmaceutical Benefits) Regulations 2017* (the Regulations). Originator removal will potentially increase price disclosure related price reductions because originator brands tend to maintain higher prices than other brands*.*

**Amendments**

An originator brand determination is set out in Schedule 1 of this amending instrument for two drugs that are moving from F1 and CDL to F2 (ezetimibe with atorvastatin; and tiotropium) on 1 February 2020. Subsection 99ADB(6C) of the Act provides that when deciding whether to determine originator brands the Minister (or delegate) must have regard to whether the brand was on F1 or CDL when it was first determined as a brand of pharmaceutical item under subsection 85(6) of the Act.

The main criteria used to determine these brands as originators were that they were the listed brands of the drug when it was on F1 or CDL.

*Basis for amendments*

Subsection 33(3) of the *Acts Interpretation Act 1901* is relied upon to vary or revoke the determination made under subsection 99ADB(6B) for the medicines affected by this amending instrument.

**Consultation**

The companies with a PBS listed brand of the drugs for which an originator brand has been determined were consulted on the potential originator brand determination.

This instrument commences on 1 February 2020.

This instrument is a legislative instrument for the purposes of the *Legislation Act 2003.*

**Statement of Compatibility with Human Rights**

*Prepared in accordance with Part 3 of the Human Rights (Parliamentary Scrutiny) Act 2011*

***National Health (Originator Brand) Amendment Determination 2020 (No. 1) (PB 9 of 2020)***

This legislative instrument is compatible with the human rights and freedoms recognised or declared in the international instruments listed in section 3 of the *Human Rights (Parliamentary Scrutiny) Act 2011*.

**Overview of the Legislative Instrument**

This legislative instrument amends the *National Health (Originator Brand) Determination 2015* (PB 100 of 2015) (the Principal Instrument) to determine the originator brand of a pharmaceutical item for two drugs new to the Pharmaceutical Benefits Scheme (PBS) F2 formulary on 1 February 2020.

The instrument is necessary to implement removal of originator brand data for price disclosure calculations (originator removal) in certain circumstances as set out in the *National Health (Pharmaceutical Benefits) Regulations 2017* (the Regulations). Originator removal will potentially increase PBS price reductions as originator brands tend to maintain higher prices than other brands. Originator removal means that the Government price would more closely reflect the prices at which generic brands of the medicine are being sold in the market, not the prices of all brands.

The PBS provides Australians with timely, reliable and affordable access to necessary and
cost-effective medicines. The PBS operates under Part VII of the *National Health Act 1953* (the Act) which regulates the listing, prescribing, pricing, charging and payment of subsidies for supply of drugs and medicinal preparations as pharmaceutical benefits. TheRegulations prescribe matters and set out details in relation to the operation of the PBS.

**Human rights implications**

This legislative instrument is compatible with Articles 2 and 12 of the International Covenant on Economic, Social and Cultural Rights by assisting with the progressive realisation by all appropriate means of the right of everyone to the enjoyment of the highest attainable standard of physical and mental health.

The PBS assists with advancement of this human right by providing subsidised access to medicines for Australians.

Originator removal will improve the operation of the PBS by delivering better value for money for PBS medicines through price reductions. This will assist consumers by reducing out-of-pocket costs for some PBS medicines.

**Conclusion**

This legislative instrument is compatible with human rights as it advances the protection of human rights.

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