Therapeutic Goods (Classes of Therapeutic Goods) Instrument 2018

I, Larry Kelly, Acting Deputy Secretary, Health Products Regulation Group, a delegate of the Secretary of the Department of Health for the purposes of section 23A of the *Therapeutic Goods Act 1989*, make the following instrument under that section.

Dated 29 March 2018

(Signed by)

LARRY KELLY
Delegate of the Secretary of the Department of Health

1 Name

This instrument is the *Therapeutic Goods (Classes of Therapeutic Goods) Instrument 2018*.

2 Commencement

This instrument commences on the day after it is registered.

3 Definitions

In this instrument:

Act means the *Therapeutic Goods Act 1989*.

complementary medicine has the meaning given by regulation 2 of the Regulations.

OTC medicine has the meaning given by regulation 2 of the Regulations.

Regulations means the *Therapeutic Goods Regulations 1990*.

4 Classes of therapeutic goods

(1) For the purposes of section 23B of the Act, the following are the classes of therapeutic goods:
  (a) prescription and other medicines specified in subsection (2);
  (b) complementary medicines;
  (c) OTC medicines specified in items 1 to 3 of Part 3 of Schedule 10 to the Regulations, other than medicines specified in paragraph (2)(b), (c), (d) or (e);
  (d) other registrable therapeutic goods.

(2) For the purposes of paragraph (1)(a), the following prescription and other medicines are specified:
  (a) prescription and other medicines specified in items 1 to 13 of Part 1 of Schedule 10 to the Regulations;
  (b) medicines containing oral nitrates for the treatment of heart disease;
  (c) nasal corticosteroids;
  (d) metered-dose asthma inhalers;
  (e) transdermal nicotine patches.