

Therapeutic Goods Information (Information about Advisory Committee Meetings) Specification 2014

*Therapeutic Goods Act 1989*

I, JOHN SKERRITT, a delegate of the Minister for Health, make this Specification under subsection 61(5D) of the *Therapeutic Goods Act 1989*.

Dated

11th April 2014

(Signed by)

**JOHN SKERRITT**

Delegate of the Minister for Health

1 Name of Specification

This Specification is the *Therapeutic Goods Information (Information about Advisory Committee Meetings) Specification 2014.*

2 Commencement

This Specification commences on the day after it is registered.

3 Definitions

In this Specification:

***Act*** means the *Therapeutic Goods Act 1989*.

***Committee*** means any of the committees established under Divisions 1-1EB of Part 6 of the Regulations.

***Regulations*** means the *Therapeutic Goods Regulations 1990*.

***TGA*** means the Therapeutic Goods Administration, which is part of the Department of Health.

4 Therapeutic goods information

The kinds of therapeutic goods information mentioned in Schedule 1 are specified under subsection 61(5D) of the Act, for the purposes of subsection 61(5C) of the Act.

Schedule 1 Specified kinds of therapeutic goods information

(section 4)

The following kinds of therapeutic goods information, being information relating to a Committee:

Note: The following specified kinds of therapeutic goods information may be released by the Secretary to the public under subsection 61(5C) of the Act.

Information about the nature or content of any discussion about matters or items considered by the Committee at a Committee meeting, or that summarises that discussion.

**Note**

1. All legislative instruments and compilations are registered on the Federal Register of Legislative Instruments kept under the *Legislative Instruments Act 2003.* See http://www.frli.gov.au