



#### Australian Government

Department of Health and Ageing Therapeutic Goods Administration

# National Coordinating Committee on Therapeutic Goods (NCCTG)

The National Coordinating Committee on Therapeutic Goods (NCCTG) was established by Order of the Federal Executive Council on 17 March 1971. The 1971 Order was revoked in October 1986 to facilitate the establishment of NCCTG as a committee of the Australian Health Ministers' Advisory Council (AHMAC).

# NCCTG terms of reference

The Terms of Reference of the NCCTG are:

- a. to promote a coordinated and harmonised approach to the regulation and controls over therapeutic goods and poisons
- b. to provide policy guidance to the National Drugs and Poisons Schedule Committee
- c. to contribute to projects relevant to the regulation and control of therapeutic goods and poisons
- d. to share knowledge on emerging and important matters relevant to the regulation of therapeutic goods and poisons
- e. to report and make recommendations to the Australian Health Ministers' Advisory Council, via the Clinical/Technical and Ethical Principal Committee, as necessary.

# NCCTG servicing

The Committee is serviced by the Therapeutic Goods Administration, a Division of the Australian Government Department of Health and Ageing.

# NCCTG membership

The Committee is chaired by the National Manager of the Therapeutic Goods Administration and there is one other Australian Government appointed member (Branch Director of the Office of Devices, Blood and Tissues of the Therapeutic Goods Administration). One member from each State and Territory health authority is appointed by the relevant Minister responsible for the health portfolio of that State or Territory. One member is nominated by the New Zealand Ministry of Health. A delegate may attend meetings in the absence of a member. At the discretion of the chair, a member may invite an observer.

#### Member terms

There are no fixed terms for members. Neither sitting fees nor any other benefits are paid to members. Members receive fares and travelling allowance for attending meetings from their various health authorities.

### **NCCTG Members**

Dr Rohan Hammett (Chair)

National Manager

Therapeutic Goods Administration

#### Dr Larry Kelly

Office Head, Office of Devices, Blood and Tissues Therapeutic Goods Administration

## Mr Bruce Battye (Acting Member)

Acting Chief Pharmacist NSW Health Department

#### Mr Andrew Petrie

Director, Medicines Infrastructure & Support Queensland Health

#### Mr Keith Moyle

Manager, Drugs and Poisons Section Department of Human Services, Victoria

## Mr Murray Patterson

Chief Pharmacist

Department of Health, Western Australia

## Ms Mary Sharpe (Acting Member)

**Chief Pharmacist** 

Department of Health and Human Services, Tasmania

### Ms Elizabeth Hender (Acting Member)

Pharmaceutical Services

Drug and Alcohol Services, South Australia (DASSA)

#### Ms Gay Lavery

**Poisons Control** 

Department of Health and Families, Northern Territory

#### Ms Jane Strang

**Chief Pharmacist** 

**ACT Health** 

#### Dr Susan Martindale

Medsafe

Ministry of Health, New Zealand

#### Mr P K Harrison (Secretary)

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# NCCTG reports & reviews

- A discussion paper on regulation of extemporaneously prepared medicines in non-hospital pharmacies
  - <a href="http://www.tga.gov.au/meds/extempcomp2.htm">http://www.tga.gov.au/meds/extempcomp2.htm</a> April 2008
- Review of the need for further regulation of extemporaneous compounding <a href="http://www.tga.gov.au/meds/extempcomp.htm">http://www.tga.gov.au/meds/extempcomp.htm</a> September 2005
- Report to Health Ministers on a cost-benefit analysis of pharmacist only (S3) and pharmacy medicines (S2) and risk-based evaluation of the standards
  <a href="http://www.tga.gov.au/meds/s2s3report.htm">http://www.tga.gov.au/meds/s2s3report.htm</a>
  August 2005

Publication content last updated: 17 February 2009 This information last reviewed: 17 February 2009 Web page last updated: 18 June 2009

URL: http://www.tga.gov.au/docs/html/ncctg.htm