

The Pharmacy Board of Australia's codes, guidelines and policies help to clarify the Board's expectations on a range of issues for pharmacists. Below is a quick reference guide to specific Board guidance. All codes, guidelines and policies can be accessed at [www.pharmacyboard.gov.au/Codes-Guidelines.aspx](http://www.pharmacyboard.gov.au/Codes-Guidelines.aspx)

### Guidelines on compounding of medicines

- 1 When to compound medicines
- 2 Competence to undertake compounding
- 3 Quality assurance
- 4 Facilities, equipment, working environments, materials and support staff
- 5 Formulation considerations
- 6 Assigning expiry dates to compounded medicines
- 7 Batch preparation
- 8 Managing risks that may lead to injury
- 9 Documentation
- 10 Packing and labelling requirements
- 11 Counselling and provision of information on compounded medicines
- 12 Supporting informed patient choice
- 13 Reporting of adverse events
- 14 Advertising
- 15 Reference texts and other sources of information relevant to compounding

Also refer to:

- *Background on the regulation of supply and manufacture of medicines including medicines compounded by pharmacists*
- *FAQ*
- *Joint statement on compounded medicines – Pharmacy Board of Australia and Medical Board of Australia*
- *Consumer fact sheet*

### Guidelines for proprietor pharmacists

- 1 Proprietors to maintain an active interest in how the pharmacy business is conducted
- 2 Proprietor pharmacists cannot delegate their professional obligations
- 3 Responsibilities of proprietor pharmacists

Also refer to:

- *Fact sheet- Registration type required by proprietor pharmacists*

### Guidelines on continuing professional development

- 1 Development of a continuing professional development plan and undertaking self-directed professional development
- 2 Accredited and non-accredited CPD
- 3 Range of activities
- 4 Records of CPD undertaken
- 5 Temporary absence from practice

Also refer to:

- *Registration standard: Continuing professional development*
- *FAQ*

### Guidelines on dose administration aids and staged supply of dispensed medicines

- 1 Dose administration aids
  - 1.1 Packaging of DAAs
  - 1.2 Labelling of DAAs
  - 1.3 Checking of DAAs
  - 1.4 Records of DAA packaging
  - 1.5 Packing oral cytotoxic and other hazardous medicines into DAAs
  - 1.6 Automated and semi-automated dose packaging systems
  - 1.7 Packing by a third party
- 2 Staged supply

### Guidelines for mandatory notifications

### Social media policy

### Code of conduct for pharmacists

### Guidelines for dispensing of medicines

- 1 The dispensing process
- 2 Dispensing precaution - safety of prescriptions
- 3 Dispensing multiple repeat prescriptions at one time
- 4 Hard-copy prescriptions that are copied and transmitted electronically
- 5 Internet, mail-order dispensing and other indirect supply of medicines
- 6 Incident records
- 7 Labelling of dispensed medicines
- 8 Counselling patients about prescribed medicines
- 9 Privacy and confidentiality
- 10 Dispensing errors and near misses
- 11 Pharmacists' workloads
- 12 Dispensary assistants/dispensary technicians and hospital pharmacy technicians
- 13 Patients' rights to choose where to access medicines
- 14 Return of unwanted medicines

### Guidelines on practice-specific issues

- 1 Reference texts for pharmacists
- 2 Drugs of abuse
- 3 Pseudoephedrine
- 4 Supply of Schedule 2 poisons (Pharmacy Medicines) and Schedule 3 poisons (Pharmacist Only Medicines)
- 5 Complementary and alternative medicines
- 6 Allied health, and complementary and alternative therapy when practised by other persons in the pharmacy
- 7 Screening and risk assessment
- 8 Raw materials not approved for human use in medicines
- 9 Supply of tobacco and alcohol products

Also refer to:

- *Guideline 1 (List of reference texts for pharmacists)*

### Guidelines for advertising regulated health services

Also refer to: *Advertising fact sheet and FAQ*