



College of Pharmacists of Manitoba

200 Tache Avenue, Winnipeg, Manitoba R2H 1A7

Phone (204) 233-1411 | Fax: (204) 237-3468

E-mail: info@cphm.ca | Website: www.cphm.ca

Practice Direction Sale of NAPRA Schedule 3 Drugs (non-prescription, pharmacy only sales)

1.0 Scope and Objective:

1.1 Expected Outcome

This document is a practice direction of Council concerning the sale of NAPRA “Schedule 3 of the Manual” Drugs through the authority of The Pharmaceutical Regulations to *The Pharmaceutical Act* and *The Pharmaceutical Act*.

1.2 Document Jurisdiction (Area of Practice)

Schedule 3 drugs can only be distributed from a licensed community pharmacy as described under section 84(4) of the regulations.

1.3 Regulatory Authority Reference

Schedule 3 drugs may be sold in a self-selection area of the pharmacy under the direct supervision of a licensed pharmacist who is available to assist the patient or designate in medication selection.

2.0 Practice Direction

- 2.1 A licensed pharmacist must be available and accessible to a person who needs to make self-selection of a Schedule 3 drug.
- 2.2 A licensed pharmacist should take the necessary steps to enter into dialogue with, or provide information to, the patient or the designate who:
 - a) requests help in selecting a Schedule 3 drug;
 - b) appears to be having difficulty in selecting a Schedule 3 drug;
 - c) is observed to be making frequent or repeat purchases of a Schedule 3 drug or purchasing a quantity that is therapeutically inappropriate; or
 - d) is recognized as someone who presents a risk from the use of a Schedule 3

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To protect the health and well being of the public by ensuring and
promoting safe, patient-centred and progressive pharmacy practice.*

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drug.

2.3 When engaging in a dialogue, the licensed pharmacist should gather specific information such as:

- 2.3.1 the condition or symptom(s) to be treated;
- 2.3.2 any previous history of complaint given as well as length of present symptoms;
- 2.3.3 current and relevant information regarding disease state(s), allergies and/or sensitivities;
- 2.3.4 current medications; and/or
- 2.3.5 other medications or therapies previously tried.

2.4 Dialogue must occur in a confidential manner.

2.5 The licensed pharmacist will enable the patient to make a choice and will discuss:

- 2.5.1 if recommending a drug therapy, directions for proper use and length of therapy, common adverse effects, and expected response or outcome or benefit(s);
- 2.5.2 non-drug treatments, if any;
- 2.5.3 follow-up with the licensed pharmacist or another health care professional if there is no improvement or symptoms change or worsen;
- 2.5.4 the need for referral to another health care professional if the condition or symptom(s) are deemed to be serious in nature; if unsure of the diagnosis or if the situation cannot be appropriately treated with non-prescription drugs; and,
- 2.5.5 the need for further dialogue with the patient directly (if a designate is involved or if a drug product is being delivered).

2.6 A licensed pharmacist will document the patient interaction and any recommendation(s) in the patient's health record if deemed appropriate.

2.7 A licensed pharmacist may refuse the sale of a Schedule 3 Drug.

3.0 Compliance Adjudication

All documentation must be readily accessible and open to regulatory review.

4.0 Appendices

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Not applicable

A Practice Direction is a written statement of a regulatory position made by Council for the purposes of giving direction to members and owners about the conduct of their practice or pharmacy operations.

A Practice Direction carries similar legal weight to a Regulation under the Act and compliance by all Manitoba pharmacists and pharmacy license holders is expected.

The process for development, consultation, implementation, appeal and review is been published on the College website.

Development Source:	Standards of Practice Committee
Regulatory Reference:	Sec 84(4), <i>The Pharmaceutical Regulations</i>
Consultation Close:	September 17, 2013
Authorized by Council:	September 30, 2013
Effective Date:	January 1, 2014
Revised:	
Review Due:	

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