



THE STANDARD OF CARE.

Medication

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Purpose: Our purpose is to protect the public by promoting safe nursing practice.

Pub. No. 49044

ISBN 978-1-77116-168-8

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First published November 1996 as Medication Administration Standards. Reprinted January 2000, October 2000, October 2002. Revised June 2003 as *Medication Standards* (2003) (ISBN 1-894557-33-6), Reprinted as Medication January 2004, December 2005. Revised June 2008 as Medication, Revised 2008, Updated June 2009 (ISBN 1-897308-46-9). Updated November 2011 for Bill 179 changes. Revised January 1,2014 for Dispensing. Revised 2015 as Medication. Advance copy circulated March 2015, practice standard in effect May 5, 2015. Revised April 2017 for changes to Ontario Regulation 275/94 (General) under the *Nursing Act, 1991*. (ISBN 978-1-77116-078-0) Updated June 2022 to update links to references. Updated June 2023 to replace *Authorizing Mechanisms* with the *Scope of Practice* standard. Updated January 2019 to reflect current Health Canada Adverse Drug Reaction information. Updated December 2023 to reflect Registered Nurse (RN) prescribing regulations.

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Ce fascicule existe en français sous le titre : Norme d'exercice : Médicaments

Introduction

The purpose of the Medication practice standard is to outline nurses' accountabilities when engaging in medication practices, such as administration, dispensing, medication storage, inventory management and disposal.

Three principles outline the expectations related to medication practices that promote public protection:

- authority
- competence
- safety

All medications have potential to cause harm and nurses are accountable to comply with all relevant legislation, applicable employer policies and the standards of practice of the profession, when engaging in medication practices.

This practice standard applies to all nurses. Registered Nurses (RNs), Registered Practical Nurses (RPNS) and Nurse Practitioners (NPs). In addition, NPs are accountable for the expectations outlined in the *Nurse Practitioner* practice standard, and RNs with prescribing authority are accountable for the expectations outlined in the *Registered Nurse (RN) Prescribing* practice standard.

Bolded terms are defined in the glossary.

Authority

Nurses must have the necessary authority to perform medication practices.¹

RNs² and RPNs require an order for a medication practice when:

- **a controlled act** is involved
- administering a prescription medication³
- it is required by legislation that applies to a practice setting⁴

Nurses:

- accept orders that are
- ✓ clear
- ✓ complete
- ✓ appropriate
- must have a therapeutic nurse-client relationship when engaging in all medication practices, including administering by injection or inhalation, or dispensing a medication
- must only dispense or administer by injection or inhalation medication for therapeutic purposes

When a nurse receives a medication order that is unclear, incomplete or inappropriate, the nurse must not perform the medication practice. Instead, the nurse must follow up with a prescriber in a timely manner.

Orders for medication can be direct orders, which apply to one client, or directives, which apply to more than one client. Orders for controlled substances must be direct orders.

¹ For a list of authorized providers - see the *Scope of Practice* standard.

² RNs with prescribing authority may dispense or administer by injection or inhalation a medication that they are authorized to prescribe without an order from another authorized provider.

³ Medications requiring a prescription can be found in the Health Canada Drug Product Database.

⁴ For example, for client treatments and diagnostic procedures, the *Public Hospitals Act*, regulation 965 requires an order from an identified practitioner, such as a Nurse Practitioner or a physician.

Competence

Nurses ensure that they have the knowledge, skill and judgment needed to perform medication practices safely.

Nurses:

- seek information from the client about their medication history
- collaborate with the client in making decisions about the plan of care in relation to medication practices
- provide education to the client regarding their medication and strategies for minimizing risk
- ensure their medication practices are evidenceinformed
- assess the appropriateness of the medication practice by considering the client, the medication and the environment
- know the limits of their own knowledge, skill and judgement and get help as needed
- do not perform medication practices they are not competent to perform

Safety

Nurses promote safe care and contribute to a culture of safety within their practice environments, when involved in medication practices.⁵

Nurses:

- must administer medication using the route of administration indicated by the prescriber
- must dispense the medication directly to the client or their representative

- must document and retain a copy of the information recorded on the container in which the medication was dispensed and include the information in the client's health record (see Appendix A: Information required on labels for dispensed medication)
- must be satisfied the medication has not and will not expire before the date on which the client is expected to take the last of the medication
- must have reasonable grounds to believe the medication has been obtained and stored safely and in accordance with any applicable legislation
- must promote and/or implement strategies to minimize the risk of misuse, addiction and drug diversion
- must take appropriate action to prevent, resolve or minimize the risk of harm to a client from a medication error or adverse reaction
- must report medication errors, near misses or adverse reactions in a timely manner
- must comply with applicable legislative requirements, employer policies and standards of practice related to the secure and appropriate storage, transportation, documentation and waste of all medication (including controlled substances)
- must not engage in conduct that results, directly or indirectly, in a personal or financial benefit that conflicts with their professional or ethical duty to a client as a result of dispensing a medication
- must collaborate in the development, implementation and evaluation of system approaches that support safe medication practices within the health care team.

⁵ See subsection 16(1) of O. Reg. 275/94.

Appendix A: Information required on labels for dispensed medication

Labels for dispensed medication must include⁶ the following:

- i. identification number, if applicable
- ii. dispensing nurse's name and title along with the name and title of the prescriber, if the nurse is not the prescriber
- iii. name, address and telephone number of the place from which the medication is dispensed
- iv. identification of the medication, as to its name, its strength (where applicable) and, if available, its manufacturer
- v. quantity of the medication dispensed
- vi. date the medication is dispensed
- vii. expiry date of the medication, if applicable
- viii. name of the client for whom the medication is dispensed
- ix. directions for use

The dispensing nurse must retain a copy of the information recorded on the container as part of the client's health record.

⁶ See subsection 18(5)6 & 18(5)7 of O. Reg 275/94.

Glossary

Adverse drug reaction: As defined in the Food and Drug Regulations, a noxious and unintended response to a drug, which occurs at doses normally used or tested for the diagnosis, treatment or prevention of a disease or the modification of an organic function

Adverse reaction: As defined in the Natural Health Products Regulations, a noxious and unintended response to a natural health product that occurs at any dose used or tested for the diagnosis, treatment or prevention of a disease or for modifying an organic function

Controlled acts: Acts that could cause harm if performed by those who do not have the knowledge, skill and judgment to perform them. These activities are listed in the *Regulated Health Professions Act*, 1991 (CNO, 2014).

Dispensing: To select, prepare and transfer stock medication for one or more prescribed medication doses to a client or the client's representative for administration at a later time

Drug diversion: When controlled substances are intentionally transferred from legitimate distribution and dispensing channels (National Opioid Use Guideline Group, 2010)

Evidence-informed: Practice that is based on successful strategies that improve client outcomes and are derived from a combination of various sources of evidence, including client perspective, research, national guidelines, policies, consensus statements, expert opinion and quality improvement data (CNO, 2014)

Medication error: Any preventable event that may cause or lead to inappropriate medication use or patient harm while the medication is in the control of the health care professional, patient, or consumer. Such events may be related to professional practice, health care products, procedures, and systems, including prescribing; order communication; product labeling, packaging and nomenclature; compounding; dispensing; distribution; administration; education; monitoring and use (National Coordinating Council for Medication Error Reporting and Prevention, 2014).

Near miss: An event, situation, or error that took place but was captured before reaching the patient (ISMP, 2009)

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