653—13.1(148,272C) Standards of practice—packaging, labeling and records of prescription drugs dispensed by a physician.

13.1(1) A physician shall dispense a prescription drug only in a container that meets the requirements of the Poison Prevention Packaging Act of 1970, 15 U.S.C. Sections 1471-1476 (2001), unless otherwise requested by the patient, and of Section 502G of the Federal Food, Drug and Cosmetic Act, 21 U.S.C. Section 301 et seq. (2001).

13.1(2) A label affixed to a container in which a prescription drug is dispensed by a physician shall include:

a. The name and address of the physician.

b. The name of the patient.

c. The date dispensed.

d. The directions for administering the prescription drug and any cautionary statement deemed appropriate by the physician.

e. The name and strength of the prescription drug in the container.

13.1(3) The provisions of subrules 13.1(1) and 13.1(2) shall not apply to packaged drug samples.

13.1(4) Physicians must document all prescription drugs dispensed to patients, including required label information. Noting such information on the patient's chart or record maintained by the physician is sufficient.

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