

151.212 LABEL OF PRESCRIPTION DRUG CONTAINERS.

Subdivision 1. **Prescription drugs.** Drugs dispensed pursuant to a prescription shall bear a label permanently affixed to the immediate container in which the drug is dispensed and which is received by the purchaser. The label shall bear the name of the manufacturer or distributor of the finished dosage form of the drug and all other information required by law and by rules of the board.

Subd. 2. **Controlled substances.** (a) In addition to the requirements of subdivision 1, when the use of any drug containing a controlled substance, as defined in chapter 152, or any other drug determined by the board, either alone or in conjunction with alcoholic beverages, may impair the ability of the user to operate a motor vehicle, the board shall require by rule that notice be prominently set forth on the label or container. Rules promulgated by the board shall specify exemptions from this requirement when there is evidence that the user will not operate a motor vehicle while using the drug.

(b) In addition to the requirements of subdivision 1, whenever a prescription drug containing an opiate is dispensed to a patient for outpatient use, the pharmacy or practitioner dispensing the drug must prominently display on the label or container a notice that states "Caution: Opioid. Risk of overdose and addiction."

Subd. 3. **Veterinary drugs.** Drugs dispensed, sold, or distributed in any manner pursuant to the order of a licensed veterinarian shall bear a label permanently affixed to the container in which the drug is dispensed and which is received by the purchaser. The label shall bear the name of the manufacturer or distributor of the finished dosage form of the drug and all other information required by law and the rules of the board.

History: 1969 c 933 s 12; 1975 c 101 s 3; 1975 c 356 s 1; 1976 c 338 s 5; 1985 c 248 s 70; 1988 c 550 s 13,14; 1Sp2017 c 6 art 12 s 1