151.36 DRUGS, MISBRANDING.

A drug shall be deemed to be misbranded:

(1) if its labeling is false or misleading in any particular;

(2) if in package form and not dispensed pursuant to a prescription unless it bears a label containing (a) the name and place of business of the manufacturer, packer, or distributor, (b) a statement of ingredients, and (c) an accurate statement of the net quantity of the contents in terms of weight, measure, or numerical count, provided, however, that under (c) reasonable variations shall be permitted, and exceptions as to small packages shall be allowed in accordance with the federal act;

(3) if any word, statement, or other information required by or under authority of this chapter to appear on the label or labeling is not prominently placed thereon with such conspicuousness (as compared with other words, statements, designs or devices, in the labeling) and in such terms as to render it to be read and understood by the ordinary individual under customary conditions of purchase and use;

(4) if it otherwise fails to meet the labeling requirements of the federal act.

History: *1969 c 933 s 17; 2014 c 285 s 7*