PUBLIC HEALTH CODE (EXCERPT)
Act 368 of 1978

333.17705 Definitions; L.

Sec. 17705. (1) "Label" means a display of written, printed, or graphic matter on the immediate container of a drug or device, but does not include package liners. A requirement made by or under authority of this part that a word, statement, or other information appear on the label is not complied with unless the word, statement, or other information appears on the outside container or wrapper of the retail package of the drug or device as displayed for sale or is easily legible through an outside container or wrapper.

(2) "Labeling" means the labels and other written, printed, or graphic matter on a drug or device or its container or wrapper, or accompanying the drug or device.

(3) "License" in addition to the definition in section 16106 means a pharmacy license, drug control license, or a manufacturer, wholesale distributor, or wholesale distributor-broker of drugs or devices license.

History: 1978, Act 368, Eff. Sept. 30, 1978;—Am. 1986, Act 304, Eff. Mar. 31, 1987;—Am. 2020, Act 142, Imd. Eff. July 14, 2020.

Popular name: Act 368