
ASSEMBLY BILL NO. 95—ASSEMBLYMEN SPIEGEL, KIRKPATRICK, BOBZIEN, DIAZ, CARLTON; AIZLEY, ELLIOT ANDERSON, BENITEZ-THOMPSON, CARRILLO, COHEN, DONDERO LOOP, EISEN, ELLISON, HARDY, HEALEY, HICKEY, HOGAN, MARTIN, OHRENSCHALL, PIERCE AND STEWART

FEBRUARY 13, 2013

JOINT SPONSORS: SENATORS PARKS; SPEARMAN AND WOODHOUSE

Referred to Committee on Commerce and Labor

SUMMARY—Revises provisions governing prescription labels. (BDR 54-648)

FISCAL NOTE: Effect on Local Government: No.
Effect on the State: No.

~

EXPLANATION – Matter in *bolded italics* is new; matter between brackets ~~omitted material~~ is material to be omitted.

AN ACT relating to pharmacy; requiring, with limited exceptions, a pharmacist or practitioner to indicate on a prescription label if a generic drug has been substituted for a drug prescribed by brand name; and providing other matters properly relating thereto.

Legislative Counsel's Digest:

1 Existing law authorizes a pharmacist or practitioner to indicate on the label of a
2 prescription that a generic drug has been substituted for a drug prescribed by brand
3 name unless the indication is prohibited by the practitioner who prescribed the
4 drug. (NRS 639.2587) This bill requires such an indication in every circumstance
5 where a generic drug is substituted for a drug prescribed by brand name unless the
6 person for whom the drug is dispensed elects not to have such an indication written
7 or typed on the label.



THE PEOPLE OF THE STATE OF NEVADA, REPRESENTED IN
SENATE AND ASSEMBLY, DO ENACT AS FOLLOWS:

- 1 **Section 1.** NRS 639.2587 is hereby amended to read as
2 follows:
3 639.2587 If a generic drug is substituted for a drug prescribed
4 by brand name, the pharmacist or practitioner ~~†~~
5 ~~1. Shall note† shall:~~
6 **1. Note** the name of the manufacturer, packer or distributor of
7 the drug actually dispensed on the prescription; and
8 2. ~~†Unless prohibited by the practitioner, may indicate†~~
9 **Indicate** the substitution by writing or typing on the label the words
10 “substituted for ~~††~~,” **or substantially similar language**, following
11 the generic name and preceding the brand name of the drug ~~††~~
12 **unless the person for whom the drug is dispensed elects not to**
13 **have such an indication written or typed on the label.**
14 **Sec. 2.** This act becomes effective on July 1, 2013.

