
ASSEMBLY BILL NO. 113—ASSEMBLYMAN HOGAN

PREFILED JANUARY 23, 2009

Referred to Committee on Commerce and Labor

SUMMARY—Revises provisions relating to the information specified on the label of the container for a prescription drug. (BDR 54-86)

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

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

AN ACT relating to prescription drugs; revising provisions relating to the information specified on the label of the container for a prescription drug; and providing other matters properly relating thereto.

Legislative Counsel’s Digest:

1 Existing law provides that, subject to certain exceptions, the expiration date of
2 a drug or medicine dispensed pursuant to a prescription must be included on the
3 label of the container in which the drug or medicine is dispensed. (NRS 639.2801)
4 **Section 1** of this bill: (1) removes the authority of a dispensing practitioner to
5 provide on the label an expiration date which is earlier than the expiration date
6 provided by the manufacturer of the drug or medicine; and (2) requires storage
7 conditions for the prescription drug or medicine to be stated on the label under
8 certain circumstances.

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

1 **Section 1.** NRS 639.2801 is hereby amended to read as
2 follows:
3 639.2801 Unless specified to the contrary in writing on the
4 prescription by the prescribing practitioner, all prescriptions filled
5 by any practitioner must be dispensed in a container to which is
6 affixed a label or other device which clearly shows:
7 1. The date.



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- 1 2. The name, address and prescription serial number of the
- 2 practitioner who filled the prescription.
- 3 3. The names of the prescribing practitioner and of the person
- 4 for whom prescribed.
- 5 4. The number of dosage units.
- 6 5. The symptom or purpose for which the drug is prescribed, if
- 7 included by the practitioner pursuant to NRS 639.2352.
- 8 6. Specific directions for use given by the prescribing
- 9 practitioner.
- 10 7. The expiration date of the effectiveness of the drug or
- 11 medicine dispensed, if that information is included on the original
- 12 label of the manufacturer of that drug or medicine. If the ~~expiration~~
- 13 ~~date specified by the manufacturer is not less than 1 year after the~~
- 14 ~~date of dispensing, the practitioner may use a date that is 1 year after~~
- 15 ~~the date of dispensing as the expiration date.] *validity of the*~~
- 16 *manufacturer's expiration date is dependent on specified storage*
- 17 *conditions, these conditions must be stated on the label.*
- 18 8. The proprietary or generic name of the drug or medicine as
- 19 written by the prescribing practitioner.
- 20 9. The strength of the drug or medicine.
- 21 ↪ The label must contain the warning:

22
23 Caution: Do not use with alcohol or nonprescribed drugs
24 without consulting the prescribing practitioner.

25 **Sec. 2.** This act becomes effective on January 1, 2010.

