

ASSEMBLY BILL NO. 199—ASSEMBLYMEN SMITH, ATKINSON, HORNE, CONKLIN; BOBZIEN, DALY, DONDERO LOOP, HICKEY, KIRKPATRICK, MASTROLUCA AND OCEGUERA

FEBRUARY 22, 2011

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Referred to Committee on Commerce and Labor

**SUMMARY**—Revises provisions governing the practice of pharmacy. (BDR 54-875)

**FISCAL NOTE:** Effect on Local Government: Increases or Newly Provides for Term of Imprisonment in County or City Jail or Detention Facility.

Effect on the State: Yes.

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

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AN ACT relating to the practice of pharmacy; revising provisions governing the authority of a registered pharmacist to collaborate with a practitioner for the implementation, monitoring and modification of drug therapy; authorizing the State Board of Pharmacy to establish regulations relating to collaborative pharmacy practice; providing a penalty; and providing other matters properly relating thereto.

**Legislative Counsel's Digest:**

Existing law authorizes a registered pharmacist to collaborate with a practitioner to engage in the implementation and modification of drug therapy for a patient at a licensed medical facility or licensed pharmacy. (NRS 639.0124) **Section 1** of this bill prescribes requirements for written guidelines and protocols which must be developed by a pharmacist who collaborates with a practitioner and requires those guidelines and protocols to be approved by the State Board of Pharmacy. **Section 1** also authorizes the written guidelines and protocols to set forth provisions for a pharmacist to implement, monitor and modify the drug therapy of a patient in a setting other than a licensed medical facility or licensed pharmacy.

A person who violates any provision of chapter 639 of NRS governing pharmacists and pharmacies, including any provision of this bill, is guilty of a misdemeanor. (NRS 639.310)

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\* A B 1 9 9 R 1 \*

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

1      **Section 1.** Chapter 639 of NRS is hereby amended by adding  
2 thereto a new section to read as follows:

3      **1. Written guidelines and protocols developed by a registered  
4      pharmacist in collaboration with a practitioner which authorize  
5      the implementation, monitoring and modification of drug therapy:**

6      **(a) May authorize a pharmacist to order and use the findings  
7      of laboratory tests and examinations.**

8      **(b) May provide for implementation, monitoring and  
9      modification of drug therapy for a patient receiving care in a  
10     licensed medical facility or, if developed to ensure continuity of  
11     care for a patient, in any setting outside a medical facility where  
12     the patient is receiving care.**

13      **(c) Must be approved by the Board.**

14      **2. The Board may adopt regulations which:**

15      **(a) Prescribe additional requirements for written guidelines  
16      and protocols developed pursuant to this section; and**

17      **(b) Set forth the process for obtaining the approval of the  
18      Board of such written guidelines and protocols.**

19      **Sec. 2.** (Deleted by amendment.)

20      **Sec. 3.** (Deleted by amendment.)

21      **Sec. 4.** (Deleted by amendment.)

22      **Sec. 5.** (Deleted by amendment.)

23      **Sec. 6.** (Deleted by amendment.)

24      **Sec. 7.** (Deleted by amendment.)

25      **Sec. 8.** (Deleted by amendment.)

26      **Sec. 9.** NRS 639.0124 is hereby amended to read as follows:

27      639.0124 "Practice of pharmacy" includes, but is not limited  
28 to, the:

29      1. Performance or supervision of activities associated with  
30 manufacturing, compounding, labeling, dispensing and distributing  
31 of a drug, including the receipt, handling and storage of  
32 prescriptions and other confidential information relating to patients.

33      2. Interpretation and evaluation of prescriptions or orders for  
34 medicine.

35      3. Participation in drug evaluation and drug research.

36      4. Advising of the therapeutic value, reaction, drug interaction,  
37 hazard and use of a drug.

38      5. Selection of the source, storage and distribution of a drug.

39      6. Maintenance of proper documentation of the source, storage  
40 and distribution of a drug.

41      7. Interpretation of clinical data contained in a person's record  
42 of medication.



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1       8. Development of written guidelines and protocols in  
2 collaboration with a practitioner which are intended for a patient in a  
3 licensed medical facility *or in a setting outside a medical facility*  
4 *where the patient is receiving care* and *which* authorize the  
5 implementation, monitoring and modification of drug therapy. The  
6 written guidelines and protocols ~~[may authorize a pharmacist to  
7 order and use the findings of laboratory tests and examinations.]~~  
8 *must comply with section 1 of this act.*

9       9. Implementation and modification of drug therapy in  
10 accordance with the authorization of the prescribing practitioner for  
11 a patient in a pharmacy in which drugs, controlled substances,  
12 poisons, medicines or chemicals are sold at retail.

13     → The term does not include the changing of a prescription by a  
14 pharmacist or practitioner without the consent of the prescribing  
15 practitioner, except as otherwise provided in NRS 639.2583.

16     **Sec. 10.** NRS 639.230 is hereby amended to read as follows:

17     639.230 1. A person operating a business in this State shall  
18 not use the letters "Rx" or "RX" or the word "drug" or "drugs,"  
19 "prescription" or "pharmacy," or similar words or words of similar  
20 import, without first having secured a license from the Board.

21     2. Each license must be issued to a specific person and for a  
22 specific location and is not transferable. The original license must be  
23 displayed on the licensed premises as provided in NRS 639.150.  
24 The original license and the fee required for reissuance of a license  
25 must be submitted to the Board before the reissuance of the license.

26     3. If the owner of a pharmacy is a partnership or corporation,  
27 any change of partners or corporate officers must be reported to the  
28 Board at such a time as is required by a regulation of the Board.

29     4. Except as otherwise provided in subsection 6, in addition to  
30 the requirements for renewal set forth in NRS 639.180, every person  
31 holding a license to operate a pharmacy must satisfy the Board that  
32 the pharmacy is conducted according to law.

33     5. Any violation of any of the provisions of this chapter by a  
34 managing pharmacist or by personnel of the pharmacy under the  
35 supervision of the managing pharmacist is cause for the suspension  
36 or revocation of the license of the pharmacy by the Board.

37     6. The provisions of this section do not prohibit ~~[a]~~:

38     (a) *A Canadian pharmacy which is licensed by the Board and*  
39 *which has been recommended by the Board pursuant to subsection 4*  
40 *of NRS 639.2328 for inclusion on the Internet website established*  
41 *and maintained pursuant to subsection 9 of NRS 223.560 from*  
42 *providing prescription drugs through mail order service to residents*  
43 *of Nevada in the manner set forth in NRS 639.2328 to 639.23286,*  
44 *inclusive ~~H~~; or*



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1       (b) A registered pharmacist or practitioner from collaborating  
2       in the implementation, monitoring and modification of drug  
3       therapy pursuant to guidelines and protocols approved by the  
4       Board.

5       **Sec. 11.** This act becomes effective upon passage and approval  
6       for the purpose of adopting regulations and on October 1, 2011, for  
7       all other purposes.

(30)



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