R203-24
January 21, 2026

INFORMATIONAL STATEMENT

The informational statement required by NRS 233B.066 numerically conforms to the subsections of the statute as follows:

1. EXPLANATION OF THE NEED FOR THE ADOPTED REGULATION

The regulation sets forth the requirements necessary for intermediate care facilities or skilled nursing facilities to obtain a license from the Board of Pharmacy. The purpose of the regulation is to ensure the safe acquisition, storage, handling, and administration of controlled substances and dangerous drugs in these facilities. The regulation also sets forth the qualifications, authority, and duties of the owners and contract employees of these facilities as it relates to prescription drugs

2. A DESCRIPTION OF HOW PUBLIC COMMENT WAS SOLICITED, A SUMMARY OF PUBLIC RESPONSE, AND AN EXPLANATION HOW OTHER INTERESTED PERSONS MAY OBTAIN A COPY OF THE SUMMARY.

The Board solicited comment on the proposed amendment by (1) posting notice, with links to the full text of the proposed amendment, to the LCB Administrative Regulation Notices webpage, (2) posting a copy of the full text of the proposed changes to the Board's website as part of the Board Hearing materials, (3) posting notice to the Nevada Public Notice website, operated by the Department of Administration, with a link back to a full text of the proposed amendment on the Board's website, and (4) posting notices and agendas in numerous public locations per NRS Chapter 233B.

The Board also solicited comment from representatives of relevant industry associations that Board Staff deemed likely to have an interest in the proposed amendment. The Board further provided time for public comment at the workshop(s) concerning the proposed amendment. The Board received no public response.

Parties interested in obtaining a copy of the summary of the comments solicited should contact Board Coordination at teamBC@pharmacy.nv.gov or call Darlene Nases at (775) 850-1440 ext. 120.

3. THE NUMBER OF PERSONS WHO: (A) ATTENDED EACH HEARING; (B) TESTIFIED AT EACH HEARING; AND (C) SUBMITTED TO THE AGENCY WRITTEN STATEMENTS.

The number of persons who attended the hearing was: 71
The number of persons who testified at the hearing was: -0-
The number of agency submitted statements was: -0-
The name of persons who testified at the hearing: -0-

4. A DESCRIPTION OF HOW COMMENT WAS SOLICITED FROM AFFECTED BUSINESSES, A SUMMARY OF THEIR RESPONSE, AND AN EXPLANATION HOW OTHER INTERESTED PERSONS MAY OBTAIN A COPY OF THE SUMMARY.

The Board solicited comment on the proposed amendment by (1) posting notice, with links to the full text of the proposed amendment, to the LCB Administrative Regulation Notices webpage, (2) posting a copy of the full text of the proposed changes to the Board's website as part of the Board Hearing materials, (3) posting notice to the Nevada Public Notice website, operated by the Department of Administration, with a link back to a full text of the proposed amendment on the Board's website, and (4) posting notices and agendas in numerous public locations per NRS Chapter 233B.

The Board also solicited comment from representatives of relevant industry associations that Board Staff deemed likely to have an interest in the proposed amendment. Furthermore, the Board provided time for public comment at the workshop(s) concerning the proposed amendment. The Board received no public response.

Parties interested in obtaining a copy of the summary of the comments solicited should contact Board Coordination at teamBC@pharmacy.nv.gov or call Darlene Nases at (775) 850-1440 ext. 120.

5. IF THE REGULATION WAS ADOPTED WITHOUT CHANGING ANY PART OF THE PROPOSED REGULATION, A SUMMARY OF THE REASONS FOR ADOPTING THE REGULATION WITHOUT CHANGE.

This regulation meets the requirements and received a unanimous vote from the Nevada State Board of Pharmacy board members for adoption with no changes.

6. THE ESTIMATED ECONOMIC EFFECT OF THE REGULATION ON THE BUSINESS WHICH IT IS TO REGULATE AND ON THE PUBLIC. THESE MUST BE STATED SEPARATELY, AND IN EACH CASE MUST INCLUDE:

A) BOTH ADVERSE AND BENEFICIAL EFFECTS.

There should be no adverse economic impact from this regulation amendment on the regulated entities or on the public. The regulation amendment will have a beneficial effect on the regulated entities and on the public by having ensuring that controlled substances and dangerous drugs are safely, acquired, stored, handled and administered at intermediate care facilities or skilled nursing facilities.

B) BOTH IMMEDIATE AND LONG-TERM EFFECTS.

Both the immediate and long-term economic effects on regulated entities and on the public will be beneficial as the proposed regulation ensures the safe acquisition, storage, handling and administration of controlled substances and dangerous drugs at intermediate care facilities or skilled nursing facilities.

7. THE ESTIMATED COST TO THE AGENCY FOR ENFORCEMENT OF THE PROPOSED REGULATION.

The cost to the Board for enforcement of the proposed regulation cannot be determined at this time since it will be dependent upon the number of applicants for registration/licensure.

8. A DESCRIPTION OF ANY REGULATIONS OF OTHER STATE OR GOVERNMENT AGENCIES WHICH THE PROPOSED REGULATION OVERLAPS OR DUPLICATES AND A STATEMENT EXPLAINING WHY THE DUPLICATION OR OVERLAPPING IS NECESSARY. IF THE REGULATION OVERLAPS OR DUPLICATES A FEDERAL REGULATION, THE NAME OF THE REGULATING FEDERAL AGENCY.

The Board of Pharmacy is not aware of any similar regulations of other state or government agencies that the proposed regulation overlaps or duplicates.

9. IF THE REGULATION INCLUDES PROVISIONS WHICH ARE MORE STRINGENT THAN A FEDERAL REGULATION WHICH REGULATES THE SAME ACTIVITY, A SUMMARY OF SUCH PROVISIONS.

The regulation does not contain provisions which are more stringent than a federal regulation which regulates the same activity.

10. IF THE REGULATION PROVIDES A NEW FEE OR INCREASES AN EXISTING FEE, THE TOTAL ANNUAL AMOUNT THE AGENCY EXPECTS TO COLLECT AND THE MANNER IN WHICH THE MONEY WILL BE USED.

The regulation amendment will impose a registration and renewal fee for the intermediate care and skill nursing facilities who wishes to acquire, store, handle and administer controlled substances and dangerous drugs to patients at their facility. The revenue generated from the fee will partially offset the costs of regulatory enforcement of this regulation incurred by the Board of Pharmacy, which will include application review, facility inspection, and issuance of license.