ADOPTED REGULATION OF THE

STATE BOARD OF PHARMACY

LCB File No. R016-01

Effective November 1, 2001

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

AUTHORITY: §1, NRS 453.246 and 639.070.

Section 1. NAC 639.512 is hereby amended to read as follows:

639.512 1. This section only applies to Class A and B packaging as defined in the *United States Pharmacopoeia*.

2. Each unit dose of a controlled substance or dangerous drug packaged or repackaged by a pharmacy must contain a label which specifies:

(a) The generic or trade name;

(b) The strength;

(c) The expiration date; and

(d) Where applicable, an internal control number or the lot number of the bulk package.

3. A unit dose of a controlled substance or dangerous drug packaged or repackaged by a pharmacy, including a hospital pharmacy, must be dispensed before the expiration date thereof. For the purposes of this section, "expiration date" means the date **[**~~6~~**]** ***12*** months after the date of the packaging or repackaging of the substance or dangerous drug **.** **[**~~unless a stability study acceptable to the board justifies an expiration date greater than the 6-month period.~~**]** No expiration date may exceed the original manufacturer's expiration date.

4. Each pharmacy must maintain a log containing , with respect to each controlled substance or dangerous drug packaged or repackaged by the pharmacy:

(a) The generic name, trade name and manufacturer;

(b) The strength;

(c) The manufacturer's expiration date;

(d) Where applicable, an internal control number;

(e) The lot number of the bulk packaging;

(f) The date of packaging or repackaging;

(g) The number of doses packaged or repackaged; and

(h) The initials of the pharmacist.