LCB File No. R218-08

PROPOSED REGULATION OF THE STATE BOARD OF PHARMACY

NAC 453.530 Schedule III. (NRS 453.146**,** 639.070**)**

1. Schedule III consists of the drugs and other substances listed in this section, by whatever official, common, usual, chemical or trade name designated.

2. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture or preparation which contains any quantity of the following substances having a stimulant effect on the central nervous system, including their salts, isomers and salts of such isomers, whenever the existence of such salts, isomers and salts of isomers is possible within the specific chemical designation, is hereby enumerated on schedule III, including:

(a) Those compounds, mixtures or preparations in dosage unit form containing any substance listed in schedule II which has a stimulant effect on the central nervous system, which compounds, mixtures or preparations were listed on August 25, 1971, as excepted compounds under the regulations of the Drug Enforcement Administration of the Department of Justice, and any other drug of the same quantitative composition as a drug shown on the list or which is the same except that it contains a lesser quantity of controlled substances;

(b) Benzphetamine;

(c) Chlorphentermine;

(d) Clortermine; or

(e) Phendimetrazine.

➥ For the purposes of this subsection, "isomer" includes the optical, position or geometric isomer.

3. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture or preparation which contains any quantity of the following substances having a depressant effect on the central nervous system is hereby enumerated on schedule III:

(a) Any substance which contains any quantity of a derivative of barbituric acid or any salt thereof;

(b) Chlorhexadol;

(c) Embutramide;

(d) Lysergic acid;

(e) Lysergic acid amide;

(f) Methyprylon;

(g) Sulfondiethylmethane;

(h) Sulfonethylmethane;

(i) Sulfonmethane;

(j) Any compound, mixture or preparation containing amobarbital, secobarbital, pentobarbital or any salt thereof and one or more other active medicinal ingredients, which are not listed in any schedule;

(k) Any suppository dosage form containing amobarbital, secobarbital, pentobarbital, or any salt of any of these drugs approved by the Food and Drug Administration of the United States Department of Health and Human Services for marketing only as a suppository; or

(l) Tiletamine and zolazepam or any salt thereof. (Some trade or other names for a tiletamine-zolazepam combination product: Telzol. Some trade or other names for tiletamine: 2-(ethylamino)-

2-(2-thienyl)-cyclohexanone. Some trade or other names for zolazepam: 4-(2-fluorophenyl)-6,8-dihydro-1,3,8-trimethylpyrazolo-[3,4-e][1,4]-diazepin-7(1H)-one, flupyrazapon).

4. Nalorphine is hereby enumerated on schedule III.

5. Unless specifically excepted or unless listed in another schedule, any material, compound, mixture or preparation containing any of the following narcotic drugs or their salts, calculated as the free anhydrous base or alkaloid, in quantities is hereby enumerated on schedule III:

(a) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit, with an equal or greater quantity of an isoquinoline alkaloid of opium;

(b) Not more than 1.8 grams of codeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(c) Not more than 300 milligrams of dihydrocodeinone (hydrocodone) per 100 milliliters or not more than 15 milligrams per dosage unit, with a fourfold or greater quantity of an isoquinoline alkaloid of opium;

(d) Not more than 300 milligrams of dihydrocodeinone (hydrocodone) per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(e) Not more than 1.8 grams of dihydrocodeine per 100 milliliters or not more than 90 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(f) Not more than 300 milligrams of ethylmorphine per 100 milliliters or not more than 15 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

(g) Not more than 500 milligrams of opium per 100 milliliters or per 100 grams, or not more than 25 milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts; or

(h) Not more than 50 milligrams of morphine per 100 milliliters or per 100 grams, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.

6. Except as otherwise provided in subsections 7 and 8, or unless listed in another schedule, any material, compound, mixture or preparation which contains any quantity of:

(a) Ephedrine, pseudoephedrine or phenylpropanolamine, their optical isomers, salts and salts of optical isomers, other than an over-the-counter ephedrine, pseudoephedrine or phenylpropanolamine drug product;

(b) N-methylephedrine, its optical isomers, salts and salts of optical isomers;

(c) Hydriodic acid; or

(d) Hydrogen iodide gas,

➥ are, as immediate precursors, controlled, the control of which is necessary to prevent, curtail or limit the manufacture of the controlled substances methamphetamine and N, N-dimethylamphetamine.

7. Ephedrine sulfate injection, as a solution, in either single-dose or multiple-dose ampules or vials in the possession of a practitioner or other person licensed by the Board to possess drugs is not a controlled substance. ***Dangerous drugs that contain ephedrine, pseudoephedrine or phenylpropanolamine are not controlled substances.***

8. Mahuang or other botanical products of genus ***Ephedra*** used in their natural state as a preparation for human consumption are not controlled substances for the purposes of this section.

9. Except as otherwise provided in subsections 10 and 11, or specifically excepted or listed in another schedule, any material, compound, mixture or preparation containing any quantity of

anabolic steroids, including their salts, isomers, esters and salts of isomers, whenever the existence of such salts of isomers is possible within the specific chemical designation, is hereby enumerated on schedule III:

(a) Androisoxazole;
(b) Androstenediol;
(c) Bolandiol;
(d) Bolasterone;
(e) Boldenone;
(f) Chlormethandienone;
(g) Clostebol;
(h) Chorionic gonadotropin (HGC);
(i) Dihydrochlormethyltestosterone;
(j) Dihydromesterone;
(k) Drostanolone;
(l) Ethylestrenol;
(m) Fluoxymesterone;
(n) Formebolone;
(o) Formyldienolone;
(p) 4-Hydroxy-19-nortestosterone;
(q) Mesterolone;
(r) Methandrenone;
(s) Methandriol;
(t) Methandrostenolone;
(u) Methenolone;
(v) 17-Methyltestosterone;
(w) Methyltrienolone;
(x) Mibolerone;
(y) Nandrolone;
(z) Norbolethone;
(aa) Norethandrolone;
(bb) Normethandrolone;
(cc) Oxandrolone;
(dd) Oxymesterone;
(ee) Oxymetholone;
(ff) Quinbolone;
(gg) Stanolone;
(hh) Stanozolol;
(ii) Stenbolone;
(jj) Testolactone;
(kk) Testosterone; or
(ll) Trenbolone.

10. Any anabolic steroid described in subsection 9 which is used solely for implantation in cattle or any other nonhuman species and is approved by the Food and Drug Administration for that use is not a controlled substance.

11. The following classifications are not controlled substances for the purposes of this section:

(a) Oral combinations containing therapeutic doses of estrogen and androgen;

(b) Parenteral preparations containing therapeutic doses of estrogen and androgen;

(c) Topical preparations containing androgens or combinations of androgen and estrogen; and

(d) Vaginal preparations.

12. Ketamine HCL is hereby enumerated on schedule III.

13. Synthetic Dronabinol in sesame oil encapsulated in a soft gelatin capsule in a drug product approved by the Food and Drug Administration (some trade or other names: (6aR-trans)-6a,7,8,10a-tetrahydro-6; 6,9-trimethyl-3-pentyl-6H-dibenzo [b,d]pyran- 1-ol; (-)-delta-9-(trans)-tetrahydrocannabinol; Marinol) is hereby enumerated on schedule III.

14. Gamma-hydroxybutyrate prepared by a registered pharmaceutical manufacturer of the Food and Drug Administration which is properly labeled, including lot numbers, and is available for medicinal purposes through a distribution system approved by the Food and Drug Administration is hereby enumerated on schedule III.

15. Human growth hormone (HGH) is hereby enumerated on schedule III.

16. As used in this section, "over-the-counter ephedrine, pseudoephedrine or phenylpropanolamine drug product" means a drug product that is packaged and sold in compliance with 21 U.S.C. §§ 801 et seq.