ENGROSSED SENATE AMENDMENTS
TO
ENGROSSED HOUSE
BILL NO. 1616 By: Derby of the House

and

Standridge of the Senate

An Act relating to Oklahoma Bureau of Narcotics and Dangerous Drugs Control; amending 63 O.S. 2011, Sections 2-103, as last amended by Section 70, Chapter 15, O.S.L. 2013, 2-105 and 2-110, as amended by Section 46, Chapter 259, O.S.L. 2012 (63 O.S. Supp. 2014, Sections 2-103 and 2-110), which relate to the Uniform Controlled Dangerous Substances Act; authorizing retired commissioned employees to purchase certain weapons; providing procedures for transferring ownership; modifying certain reporting requirement; authorizing use of state-owned vehicles by certain employees; amending 63 O.S. 2011, Sections 2-204, as last amended by Section 2, Chapter 154, O.S.L. 2014 and 2-208, as amended by Section 3, Chapter 80, O.S.L. 2012 (63 O.S. Supp. 2014, Sections 2-204 and 2-208), which relate to Schedule I and III substances; adding substances related to hallucinogenics and synthetic cannabinoids; deleting certain substance from Schedule III; amending 63 O.S. 2011, Section 2-315, which relates to the Anti-Drug Diversion Act; modifying submission requirement for destroying controlled dangerous substances; amending 63 O.S. 2011, Section 2-407, which relates to penalties for certain violations; and expanding scope of certain prohibited act.

AMENDMENT NO. 1. Page 4, lines 9-20, delete all proposed new language

AMENDMENT NO. 2. Page 6, line 16, delete Section 3 entirely and renumber subsequent sections

AMENDMENT NO. 3. Page 35, line 18 1/2, insert a new Section 5 to read as follows:

"SECTION 5. AMENDATORY 63 O.S. 2011, Section 2-210, as last amended by Section 4, Chapter 154, O.S.L. 2014 (63 O.S. Supp. 2014, Section 2-210), is amended to read as follows:

Section 2-210. Schedule IV.

A. Any material, compound, mixture, or preparation which contains any quantity of the following substances having a potential for abuse associated with a stimulant or depressant effect on the central nervous system:

1. Chloral betaine;

2. Chloral hydrate;

3. Ethchlorvynol;

4. Ethinamate;

5. Meprobamate;

6. Paraldehyde;

7. Petrichloral;

8. Diethylpropion;

9. Phentermine;

10. Pemoline;

11. Chlordiazepoxide;

12. Chlordiazepoxide and its salts, but not including chlordiazepoxide hydrochloride and clidinium bromide or chlordiazepoxide and water-soluble esterified estrogens;

13. Diazepam;

14. Oxazepam;

15. Clorazepate;

16. Flurazepam and its salts;

17. Clonazepam;

18. Barbital;

19. Mebutamate;

20. Methohexital;

21. Methylphenobarbital;

22. Phenobarbital;

23. Fenfluramine;

24. Pentazocine;

25. Propoxyphene;

26. Butorphanol;

27. Alprazolam;

28. Halazepam;

29. Lorazepam;

30. Prazepam;

31. Temazepam;

32. Triazolam;

33. Carisoprodol;

34. Dichloralphenazone;

35. Estazolam;

36. Eszopiclone;

37. Midazolam;

38. Modafinil;

39. Zaleplon;

40. Zolpidem;

41. Tramadol; ~~or~~

42. Bromazepam~~.~~ ; or

43. Suvorexant.

B. 1. The following nonnarcotic substances, which may, under the Federal Food, Drug, and Cosmetic Act (21 U.S.C., Section 301), be lawfully sold over the counter without a prescription, are excluded from all schedules of controlled substances under this title:

a. Breathe-Aid,

b. BronCare,

c. Bronchial Congestion,

d. Bronkaid Tablets,

e. Bronkaid Dual Action Caplets,

f. Bronkotabs,

g. Bronkolixir,

h. NeoRespin,

i. Pazo Hemorrhoid Ointment and Suppositories,

j. Primatene Tablets,

k. Primatene "Dual Action" Formula,

l. Quelidrine,

m. Resp, and

n. Vatronal Nose Drops.

2. At the request of any person, the Director may exempt any other drug product containing ephedrine from being included as a Schedule IV controlled substance if such product:

a. is labeled and marketed in a manner consistent with the pertinent OTC tentative final or final monograph issued by the FDA, and

b. is manufactured and distributed for legitimate medicinal use and in a manner that reduces or eliminates the likelihood of abuse.

3. In making a determination regarding a drug product, the Director, after notice and hearing, shall consider the following:

a. the history and current pattern of abuse,

b. the name and labeling of the product,

c. the intended manner of distribution, advertising and promotion of the product, and

d. other factors as may be relevant to and consistent with the public health and safety.

4. The hearing shall be held in accordance with the Administrative Procedures Act.

5. A list of current drug products meeting exemption requirements under this subsection may be obtained from the Bureau upon written request.

C. The Board of Pharmacy may except by rule any compound, mixture, or preparation containing any depressant substance listed

in subsection A of this section from the application of all or any part of the Uniform Controlled Dangerous Substances Act, Section 2-101 et seq. of this title, if the compound, mixture, or preparation contains one or more active medicinal ingredients not having a depressant effect on the central nervous system, and if the admixtures are included therein in combinations, quantity, proportion, or concentration that vitiate the potential for abuse of the substances which have a depressant effect on the central nervous system."

and renumber subsequent sections

AMENDMENT NO. 4. Page 1, restore the title to read:

"An Act relating to Oklahoma Bureau of Narcotics and Dangerous Drugs Control; amending 63 O.S. 2011, Section 2-103, as last amended by Section 70, Chapter 15, O.S.L. 2013, which relates to the Director of the Bureau; removing certain provisions relating to determination of salaries; amending 63 O.S. 2011, Section 2-105, which relates to certain reports; requiring State Medical Examiner to report deaths to certain persons; amending 63 O.S. 2011, Sections 2-204, as last amended by Section 2, Chapter 154, O.S.L. 2014, 2-208, as amended by Section 3, Chapter 80, O.S.L. 2012 (63 O.S. Supp. 2014, Sections 2-204 and 2-208), and 2-210, as last amended by Section 4, Chapter 154, O.S.L. 2014 (63 O.S. Supp. 2014, Section 2-210), which relate to Schedule I, III, and IV substances; adding substances related to hallucinogenics and synthetic cannabinoids; deleting certain substance from Schedule III; adding certain substance to Schedule IV; amending 63 O.S. 2011, Section 2-315, which relates to the Anti-Drug Diversion Act; modifying submission requirement for destroying controlled dangerous substances; amending 63 O.S. 2011, Section 2-407, which relates to penalties for certain violations; expanding scope of certain

prohibited act to include failure to disclose certain information."

Passed the Senate the 22nd day of April, 2015.

_____
Presiding Officer of the Senate

Passed the House of Representatives the ____ day of _____, 2015.

_____
Presiding Officer of the House of Representatives

ENGROSSED HOUSE
BILL NO. 1616 By: Derby of the House

and

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An Act relating to Oklahoma Bureau of Narcotics and Dangerous Drugs Control; amending 63 O.S. 2011, Sections 2-103, as last amended by Section 70, Chapter 15, O.S.L. 2013, 2-105 and 2-110, as amended by Section 46, Chapter 259, O.S.L. 2012 (63 O.S. Supp. 2014, Sections 2-103 and 2-110), which relate to the Uniform Controlled Dangerous Substances Act; authorizing retired commissioned employees to purchase certain weapons; providing procedures for transferring ownership; modifying certain reporting requirement; authorizing use of state-owned vehicles by certain employees; amending 63 O.S. 2011, Sections 2-204, as last amended by Section 2, Chapter 154, O.S.L. 2014 and 2-208, as amended by Section 3, Chapter 80, O.S.L. 2012 (63 O.S. Supp. 2014, Sections 2-204 and 2-208), which relate to Schedule I and III substances; adding substances related to hallucinogenics and synthetic cannabinoids; deleting certain substance from Schedule III; amending 63 O.S. 2011, Section 2-315, which relates to the Anti-Drug Diversion Act; modifying submission requirement for destroying controlled dangerous substances; amending 63 O.S. 2011, Section 2-407, which relates to penalties for certain violations; and expanding scope of certain prohibited act.

BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

SECTION 1. AMENDATORY 63 O.S. 2011, Section 2-103, as last amended by Section 70, Chapter 15, O.S.L. 2013 (63 O.S. Supp. 2014, Section 2-103), is amended to read as follows:

Section 2-103. A. The Director shall be appointed by the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control Commission. The Director of Narcotics and Dangerous Drugs Control on January 1, 1984, shall be initially appointed as Director. The succeeding Director shall, at the time of the appointment, have a Bachelor's Degree from an accredited college or university and at least five (5) years of experience in drug law enforcement. The Director may appoint necessary assistants, agents, and other personnel to perform the work of the office and may prescribe their titles and duties and fix their compensation~~, other than the salaries established in subsection A of Section 2-103a of this title,~~ pursuant to Merit System rules. The Director may appoint employees to the positions of Chief of Law Enforcement Information and Technology, Public Information/Education Officer, Training Officer, Program Administrators, Grants Administrator, Criminal Analysts, Legal Secretary, and Typist Clerk/Spanish Transcriptionists. The positions shall be unclassified and exempt from the rules and procedures of the Office of Management and Enterprise Services, except leave regulations. The office of the Director shall be located at a suitable place in Oklahoma City, Oklahoma.

B. 1. Agents appointed by the Director shall have the powers of peace officers generally; provided, the Director may appoint special agents and reserve special agents, who shall be unclassified

employees of the state, to meet specific investigatory needs. Special agents and reserve special agents shall not be required to meet the age and educational requirements as specified in this section.

2. Agents appointed on and after November 1, 1998, shall be at least twenty-one (21) years of age and shall have a Bachelor's Degree from an accredited college or university.

3. Each entering agent, with the exception of special agents, shall be required to serve one (1) year in a probationary status as a prerequisite to being placed on permanent status.

C. Agents appointed pursuant to the provisions of this section shall have the responsibility of investigating alleged violations and shall have the authority to arrest those suspected of having violated the provisions of the Uniform Controlled Dangerous Substances Act, as well as the crimes of money laundering and human trafficking, as otherwise set forth by laws of this state.

D. The Director may appoint reserve special agents who shall not be considered employees of the state and shall serve at the will of the Director. Reserve special agents shall complete a minimum of one hundred sixty (160) hours of training pursuant to Section 3311 of Title 70 of the Oklahoma Statutes and may not serve more than one hundred forty (140) hours per calendar month. Upon completion of training, reserve special agents appointed by the Director shall have general peace officer powers and the authority to arrest those

suspected of having violated the provisions of the Uniform Controlled Dangerous Substances Act. The agency may expend funds related to training and special reserve agents may receive travel expenses pursuant to the State Travel Reimbursement Act.

E. A commissioned employee of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control shall be entitled to receive upon retirement by reason of length of service, the continued custody and possession of the sidearm and badge carried by such employee immediately prior to retirement. In addition to the sidearm and badge, the commissioned employee may purchase the rifle, shotgun and additional service pistols issued to the commissioned employee immediately prior to retirement. The cost of purchasing the weapon shall be the cost of the weapon at the time of the initial purchase of the weapon. Upon payment of the replacement cost, the retired employee shall be entitled to ownership of the weapon. Any records regarding the ownership of each weapon transferred shall be modified to reflect the transfer to the retired commissioned employee. Proceeds from the purchase of the weapon shall be deposited in the Oklahoma Bureau of Narcotics Revolving Fund.

F. A commissioned employee of the Bureau may be entitled to receive, upon retirement by reason of disability, the continued custody and possession of the sidearm and badge carried by such

employee immediately prior to retirement upon written approval of the Director.

G. Custody and possession of the sidearm and badge of a commissioned employee killed in the line of duty may be awarded by the Director to the spouse or next of kin of the deceased employee.

H. Custody and possession of the sidearm and badge of a commissioned employee who dies while employed at the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control may be awarded by the Director to the spouse or next of kin of the deceased employee.

I. Any Director appointed on or after July 1, 2003, shall be eligible to participate in either the Oklahoma Public Employees Retirement System or in the Oklahoma Law Enforcement Retirement System and shall make an irrevocable election in writing to participate in one of the two retirement systems.

SECTION 2. AMENDATORY 63 O.S. 2011, Section 2-105, is amended to read as follows:

Section 2-105. A. It shall be the duty of all departments, officers, agencies, and employees of the state to cooperate with the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control in carrying out the functions of the office. The State Medical Examiner shall promptly report to the ~~office~~ offices of the Director of the Oklahoma Bureau of Narcotics and Dangerous Drugs Control, the Executive Director of the State Board of Medical Licensure and Supervision and the Executive Director of the State

Board of Osteopathic Examiners all deaths occurring within the state which were the result or probable result of abuse of a controlled dangerous substance.

B. The Bureau shall be required to compile a yearly report of all fatal and nonfatal drug overdoses for the State of Oklahoma. All registrants, as defined in the Anti-Drug Diversion Act, shall report any person appearing at a medical facility with a drug overdose to the central repository as provided in the Anti-Drug Diversion Act. The determination of a drug overdose shall be made solely at the discretion of the treating medical professional based on the education, experience and professional opinion of the medical professional. This information shall be considered part of the central repository pursuant to the Anti-Drug Diversion Act and shall be confidential and not open to the public pursuant to the provisions of Section 2-309D of this title.

SECTION 3. AMENDATORY 63 O.S. 2011, Section 2-110, as amended by Section 46, Chapter 259, O.S.L. 2012 (63 O.S. Supp. 2014, Section 2-110), is amended to read as follows:

Section 2-110. The Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control may employ attorneys, who shall be unclassified employees of the state, or contract with attorneys, as needed. These attorneys may advise the Director, the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control Commission and Bureau personnel on all legal matters and shall

appear for and represent the Director, the Commission and Bureau personnel in all administrative hearings and all litigation or other proceedings which may arise in the discharge of their duties. At the request of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control Commission, such attorney shall assist the district attorney in prosecuting charges of violators of the Uniform Controlled Dangerous Substances Act or any felony relating to or arising from a violation of the Uniform Controlled Dangerous Substances Act. Attorneys for the Bureau who have been certified by the Council on Law Enforcement Education and Training to carry a weapon or have been issued a handgun license pursuant to the provisions of the Oklahoma Self-Defense Act shall be allowed to carry weapons pursuant to paragraph 3 of subsection A of Section 1272 of Title 21 of the Oklahoma Statutes. Attorneys for the Bureau may use state-owned vehicles to provide transportation between the residence of the employee and the assigned place of employment and between the residence of the employee and any location other than the assigned place of employment to which the employee travels in the performance of the official duty of the employee. These attorneys, pursuant to this provision, shall not be considered eligible to participate in the Oklahoma Law Enforcement Retirement System. If a conflict of interest would be created by such attorney representing the Director, the Commission or Bureau personnel,

additional counsel may be hired upon approval of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control Commission.

SECTION 4. AMENDATORY 63 O.S. 2011, Section 2-204, as last amended by Section 2, Chapter 154, O.S.L. 2014 (63 O.S. Supp. 2014, Section 2-204), is amended to read as follows:

Section 2-204. The controlled substances listed in this section are included in Schedule I.

A. Any of the following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters, and ethers, unless specifically excepted, when the existence of these isomers, esters, ethers, and salts is possible within the specific chemical designation:

1. Acetylmethadol;

2. Allylprodine;

3. Alphacetylmethadol;

4. Alphameprodine;

5. Alphamethadol;

6. Benzethidine;

7. Betacetylmethadol;

8. Betameprodine;

9. Betamethadol;

10. Betaprodine;

11. Clonitazene;

12. Dextromoramide;

13. Dextrorphan (except its methyl ether);

14. Diampromide;

15. Diethylthiambutene;

16. Dimenoxadol;

17. Dimepheptanol;

18. Dimethylthiambutene;

19. Dioxaphetyl butyrate;

20. Dipipanone;

21. Ethylmethylthiambutene;

22. Etonitazene;

23. Etoxeridine;

24. Furethidine;

25. Hydroxypethidine;

26. Ketobemidone;

27. Levomoramide;

28. Levophenacylmorphan;

29. Morpheridine;

30. Noracymethadol;

31. Norlevorphanol;

32. Normethadone;

33. Norpipanone;

34. Phenadoxone;

35. Phenampromide;

36. Phenomorphan;

37. Phenoperidine;

38. Piritramide;

39. Proheptazine;

40. Properidine;

41. Racemoramide; or

42. Trimeperidine.

B. Any of the following opium derivatives, their salts, isomers, and salts of isomers, unless specifically excepted, when the existence of these salts, isomers, and salts of isomers is possible within the specific chemical designation:

1. Acetorphine;

2. Acetyldihydrocodeine;

3. Benzylmorphine;

4. Codeine methylbromide;

5. Codeine-N-Oxide;

6. Cyprenorphine;

7. Desomorphine;

8. Dihydromorphine;

9. Etorphine;

10. Heroin;

11. Hydromorphinol;

12. Methyldesorphine;

13. Methylhydromorphine;

14. Morphine methylbromide;

15. Morphine methylsulfonate;

16. Morphine-N-Oxide;

17. Myrophine;

18. Nicocodeine;

19. Nicomorphine;

20. Normorphine;

21. Phoclodine; or

22. Thebacon.

C. Any material, compound, mixture, or preparation which contains any quantity of the following hallucinogenic substances, their salts, isomers, and salts of isomers, unless specifically excepted, when the existence of these salts, isomers, and salts of isomers is possible within the specific chemical designation:

1. Methcathinone;

2. 3, 4-methylenedioxy amphetamine;

3. 3, 4-methylenedioxy methamphetamine;

4. 5-methoxy-3, 4-methylenedioxy amphetamine;

5. 3, 4, 5-trimethoxy amphetamine;

6. Bufotenine;

7. Diethyltryptamine;

8. Dimethyltryptamine;

9. 4-methyl-2, 5-dimethoxyamphetamine;

10. Ibogaine;

11. Lysergic acid diethylamide;

12. Marihuana;

13. Mescaline;

14. N-benzylpiperazine;

15. N-ethyl-3-piperidyl benzilate;

16. N-methyl-3-piperidyl benzilate;

17. Psilocybin;

18. Psilocyn;

19. 2, 5 dimethoxyamphetamine;

20. 4 Bromo-2, 5-dimethoxyamphetamine;

21. 4 methoxyamphetamine;

22. Cyclohexamine;

23. Salvia Divinorum;

24. Salvinorin A;

25. Thiophene Analog of Phencyclidine. Also known as: 1-(1-(2-thienyl) cyclohexyl) piperidine; 2-Thienyl Analog of Phencyclidine; TPCP, TCP;

26. Phencyclidine (PCP);

27. Pyrrolidine Analog for Phencyclidine. Also known as 1-(1-Phenylcyclohexyl) - Pyrrolidine, PCPy, PHP;

28. 1-(3-trifluoromethylphenyl) piperazine;

29. Flunitrazepam;

30. B-hydroxy-amphetamine;

31. B-ketoamphetamine;

32. 2,5-dimethoxy-4-nitroamphetamine;

33. 2,5-dimethoxy-4-bromophenethylamine;

34. 2,5-dimethoxy-4-chlorophenethylamine;

35. 2,5-dimethoxy-4-iodoamphetamine;

36. 2,5-dimethoxy-4-iodophenethylamine;

37. 2,5-dimethoxy-4-methylphenethylamine;

38. 2,5-dimethoxy-4-ethylphenethylamine;

39. 2,5-dimethoxy-4-fluorophenethylamine;

40. 2,5-dimethoxy-4-nitrophenethylamine;

41. 2,5-dimethoxy-4-ethylthio-phenethylamine;

42. 2,5-dimethoxy-4-isopropylthio-phenethylamine;

43. 2,5-dimethoxy-4-propylthio-phenethylamine;

44. 2,5-dimethoxy-4-cyclopropylmethylthio-phenethylamine;

45. 2,5-dimethoxy-4-tert-butylthio-phenethylamine;

46. 2,5-dimethoxy-4-(2-fluoroethylthio)-phenethylamine;

47. 5-methoxy-N, N-dimethyltryptamine;

48. N-methyltryptamine;

49. A-ethyltryptamine;

50. A-methyltryptamine;

51. N, N-diethyltryptamine;

52. N, N-diisopropyltryptamine;

53. N, N-dipropyltryptamine;

54. 5-methoxy-a-methyltryptamine;

55. 4-hydroxy-N, N-diethyltryptamine;

56. 4-hydroxy-N, N-diisopropyltryptamine;

57. 5-methoxy-N, N-diisopropyltryptamine;

58. 4-hydroxy-N-isopropyl-N-methyltryptamine;

59. 3,4-Methylenedioxymethcathinone (Methylone);

60. 3,4-Methylenedioxypyrovalerone (MDPV);

61. 4-Methylmethcathinone (Mephedrone);

62. 4-methoxymethcathinone;

63. 4-Fluoromethcathinone;

64. 3-Fluoromethcathinone;

65. 1-(8-bromobenzo 1,2-b;4,5-b' difuran-4-yl)-2-aminopropane;

66. 2,5-Dimethoxy-4-chloroamphetamine;

67. 4-Methylethcathinone;

68. Pyrovalerone;

69. N,N-diallyl-5-methoxytryptamine;

70. 3,4-Methylenedioxy-N-ethylcathinone (Ethylone);

71. B-keto-N-Methylbenzodioxolylbutanamine (Butylone);

72. B-keto-Methylbenzodioxolylpentanamine (Pentylone);

73. Alpha-Pyrrolidinopentiophenone;

74. 4-Fluoroamphetamine;

75. Pentredone;

76. 4'-Methyl-a-pyrrolidinohexaphenone;

77. 2,5-dimethoxy-4-(n)-propylphenethylamine;

78. 2,5-dimethoxyphenethylamine;

79. 1,4-Dibenzylpiperazine;

80. N,N-Dimethylamphetamine;

81. 4-Fluoromethamphetamine;

82. 4-Chloro-2,5-dimethoxy-N-(2-methoxybenzyl)phenethylamine (25C-NBOMe);

83. 4-Iodo-2,5-dimethoxy-N-(2-methoxybenzyl)phenethylamine (25I-NBOMe);

84. 4-Bromo-2,5-dimethoxy-N-(2-methoxybenzy)phenethylamine (25B-NBOMe); ~~or~~

85. 1-(4-Fluorophenyl)piperazine; or

86. Methoxetamine.

D. Unless specifically excepted or unless listed in a different schedule, any material, compound, mixture, or preparation which contains any quantity of the following substances having stimulant or depressant effect on the central nervous system:

1. Fenethylline;

2. Mecloqualone;

3. N-ethylamphetamine;

4. Methaqualone;

5. Gamma-Hydroxybutyric Acid, also known as GHB, gamma-hydroxybutyrate, 4-hydroxybutyrate, 4-hydroxybutanoic acid, sodium oxybate, and sodium oxybutyrate;

6. Gamma-Butyrolactone (GBL) as packaged, marketed, manufactured or promoted for human consumption, with the exception of legitimate food additive and manufacturing purposes;

7. Gamma Hydroxyvalerate (GHV) as packaged, marketed, or manufactured for human consumption, with the exception of legitimate food additive and manufacturing purposes;

8. Gamma Valerolactone (GVL) as packaged, marketed, or manufactured for human consumption, with the exception of legitimate food additive and manufacturing purposes; or

9. 1,4 Butanediol (1,4 BD or BDO) as packaged, marketed, manufactured, or promoted for human consumption with the exception of legitimate manufacturing purposes.

E. 1. The following industrial uses of Gamma-Butyrolactone, Gamma Hydroxyvalerate, Gamma Valerolactone, or 1,4 Butanediol are excluded from all schedules of controlled substances under this title:

a. pesticides,

b. photochemical etching,

c. electrolytes of small batteries or capacitors,

d. viscosity modifiers in polyurethane,

e. surface etching of metal coated plastics,

f. organic paint disbursements for water soluble inks,

g. pH regulators in the dyeing of wool and polyamide fibers,

h. foundry chemistry as a catalyst during curing,

i. curing agents in many coating systems based on urethanes and amides,

j. additives and flavoring agents in food, confectionary, and beverage products,

k. synthetic fiber and clothing production,

l. tetrahydrofuran production,

m. gamma butyrolactone production,

n. polybutylene terephthalate resin production,

o. polyester raw materials for polyurethane elastomers and foams,

p. coating resin raw material, and

q. as an intermediate in the manufacture of other chemicals and pharmaceuticals.

2. At the request of any person, the Director may exempt any other product containing Gamma-Butyrolactone, Gamma Hydroxyvalerate, Gamma Valerolactone, or 1,4 Butanediol from being included as a Schedule I controlled substance if such product is labeled, marketed, manufactured and distributed for legitimate industrial use in a manner that reduces or eliminates the likelihood of abuse.

3. In making a determination regarding an industrial product, the Director, after notice and hearing, shall consider the following:

a. the history and current pattern of abuse,

b. the name and labeling of the product,

c. the intended manner of distribution, advertising and promotion of the product, and

d. other factors as may be relevant to and consistent with the public health and safety.

4. The hearing shall be held in accordance with the procedures of the Administrative Procedures Act.

F. Any material, compound, mixture, or preparation, whether produced directly or indirectly from a substance of vegetable origin or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis, that contains any quantity of the following substances, or that contains any of their salts, isomers, and salts of isomers when the existence of these salts, isomers, and salts of isomers is possible within the specific chemical designation:

1. JWH-004;

2. JWH-007;

3. JWH-009;

4. JWH-015;

5. JWH-016;

6. JWH-018;

7. JWH-019;

8. JWH-020;

9. JWH-030;

10. JWH-046;

11. JWH-047;

12. JWH-048;

13. JWH-049;

14. JWH-050;

15. JWH-070;

16. JWH-071;

17. JWH-072;

18. JWH-073;

19. JWH-076;

20. JWH-079;

21. JWH-080;

22. JWH-081;

23. JWH-082;

24. JWH-094;

25. JWH-096;

26. JWH-098;

27. JWH-116;

28. JWH-120;

29. JWH-122;

30. JWH-145;

31. JWH-146;

32. JWH-147;

33. JWH-148;

34. JWH-149;

35. JWH-150;

36. JWH-156;

37. JWH-167;

38. JWH-175;

39. JWH-180;

40. JWH-181;

41. JWH-182;

42. JWH-184;

43. JWH-185;

44. JWH-189;

45. JWH-192;

46. JWH-193;

47. JWH-194;

48. JWH-195;

49. JWH-196;

50. JWH-197;

51. JWH-198;

52. JWH-199;

53. JWH-200;

54. JWH-201;

55. JWH-202;

56. JWH-203;

57. JWH-204;

58. JWH-205;

59. JWH-206;

60. JWH-207;

61. JWH-208;

62. JWH-209;

63. JWH-210;

64. JWH-211;

65. JWH-212;

66. JWH-213;

67. JWH-234;

68. JWH-235;

69. JWH-236;

70. JWH-237;

71. JWH-239;

72. JWH-240;

73. JWH-241;

74. JWH-242;

75. JWH-243;

76. JWH-244;

77. JWH-245;

78. JWH-246;

79. JWH-248;

80. JWH-249;

81. JWH-250;

82. JWH-251;

83. JWH-252;

84. JWH-253;

85. JWH-262;

86. JWH-292;

87. JWH-293;

88. JWH-302;

89. JWH-303;

90. JWH-304;

91. JWH-305;

92. JWH-306;

93. JWH-307;

94. JWH-308;

95. JWH-311;

96. JWH-312;

97. JWH-313;

98. JWH-314;

99. JWH-315;

100. JWH-316;

101. JWH-346;

102. JWH-348;

103. JWH-363;

104. JWH-364;

105. JWH-365;

106. JWH-367;

107. JWH-368;

108. JWH-369;

109. JWH-370;

110. JWH-371;

111. JWH-373;

112. JWH-386;

113. JWH-387;

114. JWH-392;

115. JWH-394;

116. JWH-395;

117. JWH-397;

118. JWH-398;

119. JWH-399;

120. JWH-400;

121. JWH-412;

122. JWH-413;

123. JWH-414;

124. JWH-415;

125. CP-55, 940;

126. CP-47, 497;

127. HU-210;

128. HU-211;

129. WIN-55, 212-2;

130. AM-2201;

131. AM-2233;

132. JWH-018 adamantyl-carboxamide;

133. AKB48;

134. JWH-122 N-(4-pentenyl)analog;

135. MAM2201;

136. URB597;

137. URB602;

138. URB754;

139. UR144;

140. XLR11;

141. A-796,260;

142. STS-135;

143. AB-FUBINACA;

144. AB-PINACA;

145. PB-22;

146. AKB48 N-5-Fluorpentyl;

147. AM1248;

148. FUB-PB-22;

149. ADB-FUBINACA;

150. BB-22;

151. 5-Fluoro PB-22; or

152. 5-Fluoro AKB-48.

G. In addition to those substances listed in subsection F of this section, unless specifically excepted or unless listed in another schedule, any material, compound, mixture, or preparation

which contains any quantity of a synthetic cannabinoid found to be in any of the following chemical groups:

1. Naphthoylindoles: any compound containing a 3-(1-naphthoyl)indole structure with or without substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3- morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the indole ring to any extent, and whether or not substituted on the naphthyl ring to any extent. Naphthoylindoles include, but are not limited to:

a. 1-[2-(4-morpholinyl)ethyl]-3-(1-naphthoyl)indole (JWH-200),

b. 1-(5-fluoropentyl)-3-(1-naphthoyl)indole (AM2201),

c. 1-pentyl-3-(1-naphthoyl)indole (JWH-018),

d. 1-butyl-3-(1-naphthoyl)indole (JWH-073),

e. 1-pentyl-3-(4-methoxy-1-naphthoyl)indole (JWH-081),

f. 1-propyl-2-methyl-3-(1-naphthoyl)indole (JWH-015),

g. 1-hexyl-3-(1-naphthoyl)indole (JWH-019),

h. 1-pentyl-3-(4-methyl-1-naphthoyl)indole (JWH-122),

i. 1-pentyl-3-(4-ethyl-1-naphthoyl)indole (JWH-210),

j. 1-pentyl-3-(4-chloro-1-naphthoyl)indole (JWH-398),

k. 1-pentyl-2-methyl-3-(1-naphthoyl)indole (JWH-007),

l. 1-pentyl-3-(7-methoxy-1-naphthoyl)indole (JWH-164),

m. 1-pentyl-2-methyl-3-(4-methoxy-1-naphthoyl)indole (JWH-098),

n. 1-pentyl-3-(4-fluoro-1-naphthoyl)indole (JWH-412),

o. 1-[1-(N-methyl-2-piperidinyl)methyl]-3-(1-naphthoyl)indole (AM-1220),

p. 1-(5-fluoropentyl)-3-(4-methyl-1-naphthoyl)indole (MAM-2201), or

q. 1-(4-cyanobutyl)-3-(1-naphthoyl)indole (AM-2232);

2. Naphthylmethylindoles: any compound containing a 1H-indol-3-yl-(1-naphthyl)methane structure with or without substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3- morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the indole ring to any extent, and whether or not substituted on the naphthyl ring to any extent. Naphthylmethylindoles include, but are not limited to, (1-pentylindol-3-yl)(1-naphthyl)methane (JWH-175);

3. Naphthoylpyrroles: any compound containing a 3-(1-naphthoyl)pyrrole structure with or without substitution at the nitrogen atom of the pyrrole ring by an alkyl, haloalkyl,

cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the pyrrole ring to any extent, and whether or not substituted on the naphthyl group to any extent. Naphthoylpyrroles include, but are not limited to:

a. 1-hexyl-2-phenyl-4-(1-naphthoyl)pyrrole (JWH-147),

b. 1-pentyl-5-(2-methylphenyl)-3-(1-naphthoyl)pyrrole (JWH-370),

c. 1-pentyl-3-(1-naphthoyl)pyrrole (JWH-030), or

d. 1-hexyl-5-phenyl-3-(1-naphthoyl)pyrrole (JWH-147);

4. Naphthylideneindenes: any compound containing a ~~naphthylideneindene~~ 1-(1-naphthylmethylene)indene structure with or without substitution at the 3-position of the indene ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the indene group to any extent, and whether or not substituted on the naphthyl group to any extent. Naphthylmethylindenes include, but are not

limited to, (1-[(3-pentyl)-1H-inden-1-ylidene)methyl]naphthalene (JWH-176);

5. Phenylacetylindoles: any compound containing a 3-phenylacetylindole structure with or without substitution at the nitrogen atom of the indole ring by alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3- morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the indole ring to any extent, and whether or not substituted on the phenyl ring to any extent. Phenylacetylindoles include, but are not limited to:

a. 1-pentyl-3-(2-methoxyphenylacetyl)indole (JWH-250),

b. 1-(2-cyclohexylethyl)-3-(2-methoxyphenylacetyl)indole (RCS-8),

c. 1-pentyl-3-(2-chlorophenylacetyl)indole (JWH-203),

d. 1-pentyl-3-(2-methylphenylacetyl)indole (JWH-251),

e. 1-pentyl-3-(4-methoxyphenylacetyl)indole (JWH-201), or

f. 1-pentyl-3-(3-methoxyphenylacetyl)indole (JWH-302);

6. Cyclohexylphenols: any compound containing a 2-(3-hydroxycyclohexyl)phenol structure with or without substitution at the 5-position of the phenolic ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl,

halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, and whether or not further substituted on the cyclohexyl ring to any extent. Cyclohexylphenols include, but are not limited to:

a. 5-(1,1-dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (CP-47,497),

b. 5-(1,1-dimethyloctyl)-2-[(1R,3S)-3-hydroxycyclohexyl]-phenol (cannabicyclohexanol; CP-47,497 C8 homologue), or

c. 5-(1,1-dimethylheptyl)-2-[(1R,2R)-5-hydroxy-2-(3-hydroxypropyl)cyclohexyl]-phenol (CP 55,~~490~~ 940);

7. Benzoylindoles: any compound containing a 3-(~~1-~~benzoyl)indole structure with or without substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3- morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted on the indole ring to any extent, and whether or not substituted on the phenyl group to any extent. Benzoylindoles include, but are not limited to:

a. 1-pentyl-3-(4-methoxybenzoyl)indole (RCS-4),

b. 1-[2-(4-morpholinyl)ethyl]-2-methyl-3-(4-methoxybenzoyl)indole (Pravadoline or WIN 48, 098),

c. 1-(5-fluoropentyl)-3-(2-iodobenzoyl)indole (AM-694),

d. 1-pentyl-3-(2-iodobenzoyl)indole (AM-679), or

e. 1-[1-(N-methyl-2-piperidinyl)methyl]-3-(2-iodobenzoyl)indole (AM-2233);

8. Cyclopropoylindoles: Any compound containing a 3-(cyclopropoyl)indole structure with substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3- morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted in the indole ring to any extent and whether or not substituted in the cyclopropoyl ring to any extent. Cyclopropoylindoles include, but are not limited to:

a. 1-pentyl-3-(2,2,3,3-tetramethylcyclopropoyl)indole (UR-144),

b. 1-(5-chloropentyl)-3-(2,2,3,3-tetramethylcyclopropoyl)indole (5Cl-UR-144), or

c. 1-(5-fluoropentyl)-3-(2,2,3,3-tetramethylcyclopropoyl)indole (XLR11);

9. Indole Amides: Any compound containing a 1H-Indole-3-carboxamide structure with or without substitution at ~~either~~ the nitrogen atom of the ~~indazole~~ indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, ~~or~~ (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not substituted at the carboxamide group by an adamantyl, ~~1-napthyl~~ naphthyl, ~~or phenol~~ phenyl, benzyl, quinolinyl, cycloalkyl, 1-amino-3-methyl-1-oxobutan-2-yl, 1-amino-3,3-dimethyl-1-oxobutan-2-yl, 1-methoxy-3-methyl-1-oxobutan-2-yl, 1-methoxy-3,3-dimethyl-1-oxobutan-2-yl or pyrrole group, and whether or not further substituted in the indole, adamantyl, naphthyl, ~~or~~ phenyl, pyrrole, quninolinyl, or cycloalkyl rings to any extent. Indole Amides include, but are not limited to:

a. N-(1-adamantyl)-1-pentyl-1H-indole-3-carboxamide (2NE1), ~~or~~

b. N-(1-adamantyl)-1-(5-fluoropentyl-1H-indole-3-carboxamide (STS-135),

c. N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-pentyl-1H-indole-3-carboxamide (ADBICA),

d. N-(1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-(5-fluoropentyl)-1H-indole-3-carboxamide (5F-ADBICA),

e. N-(naphthalen-1-yl)-1-pentyl-1H-indole-3-carboxamide (NNE1),

f. 1-(5-fluoropentyl)-N-(naphthalene-1-yl)-1H-indole-3-carboxamide (5F-NNE1),

g. N-benzyl-1-pentyl-1H-indole-3-carboxamide (SDB-006), or

h. N-benzyl-1-(5-fluoropentyl)-1H-indole-3-carboxamide (5F-SDB-006); ~~and~~

10. Indole Esters: Any compound containing a 1H-Indole-3-carboxylate structure with or without substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not substituted at the carboxylate group by an adamantyl, naphthyl, phenyl, benzyl, quinolinyl, cycloalkyl,1-amino-3-methyl-1-oxobutan-2-yl, 1-amino-3,3-dimethyl-1-oxobutan-2-yl, 1-methoxy-3-methyl-1-oxobutan-2-yl, 1-methoxy-3,3-dimethyl-1-oxobutan-2-yl or pyrrole group, and whether or not further substituted in the indole, adamantyl, naphthyl, phenyl, pyrrole, quinolinyl, or cycloalkyl rings to any extent. Indole Esters include, but are not limited to:

a. quinolin-8-yl 1-pentyl-1H-indole-3-carboxylate (PB-22),

b. quinolin-8-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate (5F-PB-22),

c. quinolin-8-yl 1-(cyclohexylmethyl)-1H-indole-3-carboxylate (BB-22),

d. naphthalen-1-yl 1-(4-fluorobenzyl)-1H-indole-3-carboxylate (FDU-PB-22), or

e. naphthalen-1-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate (NM2201);

11. Adamantanoylindoles: Any compound containing an adamantanyl-(1H-indol-3-yl)methanone structure with or without substitution at the nitrogen atom of the indole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, whether or not further substituted in the indole ring to any extent and whether or not substituted in the adamantyl ring to any extent. Adamantanoylindoles include, but are not limited to:

a. adamantan-1-yl[1-[(1-methyl-2-piperidinyl)methyl]-1H-indol-3-yl]methanone (AM1248), or

b. adamantan-1-yl-(1-pentyl-1H-indol-3-yl)methanone (AB-001);

12. Carbazole Ketone: Any compound containing (9H-carbazole-3-yl) methanone structure with or without substitution at the nitrogen atom of the carbazole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, with substitution at the carbon of the methanone group by an adamantyl, naphthyl, phenyl, benzyl, quinolinyl, cycloalkyl, 1-amino-3-methyl-1-oxobutan-2-yl, 1-amino-3,3-dimethyl-1-oxobutan-2-yl, 1-methoxy-3-methyl-1-oxobutan-2-yl, 1-methoxy-3,3-dimethyl-1-oxobutan-2-yl or pyrrole group, and whether or not further substituted at the carbazole, adamantyl, naphthyl, phenyl, pyrrole, quinolinyl, or cycloalkyl rings to any extent. Carbazole Ketones include, but are not limited to, naphthalen-1-yl(9-pentyl-9H-carbazol-3-yl)methanone (EG-018);

13. Benzimidazole Ketone: Any compound containing (benzimidazole-2-yl) methanone structure with or without substitution at either nitrogen atom of the benzimidazole ring by an alkyl, haloalkyl, cyanoalkyl, alkenyl, cycloalkylmethyl, cycloalkylethyl, benzyl, halobenzyl, 1-(N-methyl-2-piperidinyl)methyl, 2-(4-morpholinyl)ethyl, 1-(N-methyl-2-

pyrrolidinyl)methyl, 1-(N-methyl-3-morpholinyl)methyl, (tetrahydropyran-4-yl)methyl, 1-methylazepanyl, phenyl, or halophenyl group, with substitution at the carbon of the methanone group by an adamantyl, naphthyl, phenyl, benzyl, quinolinyl, cycloalkyl, 1-amino-3-methyl-1-oxobutan-2-yl, 1-amino-3,3-dimethyl-1-oxobutan-2-yl, 1-methoxy-3-methyl-1-oxobutan-2-yl, 1-methoxy-3,3-dimethyl-1-oxobutan-2-yl or pyrrole group, and whether or not further substituted in the benzimidazole, adamantyl, naphthyl, phenyl, pyrrole, quinolinyl, or cycloalkyl rings to any extent. Benzimidazole Ketones include, but are not limited to:

a. naphthalen-1-yl(1-pentyl-1H-benzo[d]imidazol-2-l)methanone (JWH-018 benzimidazole analog), or

b. (1-(5-fluoropentyl)-1H-benzo[d]imidazol-2-yl)(naphthalen-1-yl)methanone (FUBIMINA); and

14. Modified by Replacement: any compound defined in this subsection that is modified by replacement of a carbon with nitrogen in the indole, naphthyl, ~~or~~ indene, benzimidazole, or carbazole ring.

SECTION 5. AMENDATORY 63 O.S. 2011, Section 2-208, as amended by Section 3, Chapter 80, O.S.L. 2012 (63 O.S. Supp. 2014, Section 2-208), is amended to read as follows:

Section 2-208. The controlled substances listed in this section are included in Schedule III.

A. Unless listed in another schedule, any material, compound, mixture, or preparation, which contains any quantity of the following substances or any other substance having a potential for abuse associated with a stimulant or depressant effect on the central nervous system:

1. Any drug product containing gamma-hydroxybutyric acid, including its salts, isomers, and salts of isomers, for which an application has been approved under Section 505 of the Federal Food, Drug, and Cosmetic Act;

2. Any material, compound, mixture, or preparation which contains any quantity of the following hormonal substances or steroids, including their salts, isomers, esters and salts of isomers and esters, when the existence of these salts, isomers, esters, and salts of isomers and esters is possible within the specific chemical designation:

a. Boldenone,

b. Chlorotestosterone,

c. Clostebol,

d. Dehydrochlormethyltestosterone,

e. Dihydrotestosterone,

f. Drostanolone,

g. Ethylestrenol,

h. Fluoxymesterone,

i. Formebolone,

j. Mesterolone,

k. Methandienone,

l. Methandranone,

m. Methandriol,

n. Methandrostenolone,

o. Methenolone,

p. Methyltestosterone, except as provided in subsection E of this section,

q. Mibolerone,

r. Nandrolone,

s. Norethandrolone,

t. Oxandrolone,

u. Oxymesterone,

v. Oxymetholone,

w. Stanolone,

x. Stanozolol,

y. Testolactone,

z. Testosterone, except as provided in subsection E of this section, and

aa. Trenbolone;

3. Any substance which contains any quantity of a derivative of barbituric acid, or any salt of a derivative of barbituric acid;

4. Benzephetamine and its salts;

5. Buprenorphine;

6. Butalbital/acetaminophen/caffeine;

7. Chlorhexadol;

8. Chlorphentermine and its salts;

9. Clortermine;

10. Glutethimide;

11. ~~Hydrocodone with another active ingredient;~~

~~12.~~ Ketamine, its salts, isomers, and salts of isomers;

~~13.~~ 12. Lysergic acid;

~~14.~~ 13. Lysergic acid amide;

~~15.~~ 14. Mazindol;

~~16.~~ 15. Methyprylon;

~~17.~~ 16. Phendimetrazine;

~~18.~~ 17. Phenylacetone (P2P);

~~19.~~ 18. Sulfondiethylmethane;

~~20.~~ 19. Sulfonethylmethane;

~~21.~~ 20. Sulfonmethane;

~~22.~~ 21. Tetrahydrocannibinols;

~~23.~~ 22. 1-Phenycyclohexylamine; or

~~24.~~ 23. 1-Piperidinocychexanecarbo nitrile (PCC).

Livestock implants as regulated by the Federal Food and Drug Administration shall be exempt.

B. Nalorphine.

C. Unless listed in another schedule, any material, compound, mixture, or preparation containing limited quantities of any of the following narcotic drugs, or any salts thereof:

1. Not more than one and eight-tenths (1.8) grams of codeine or any of its salts, per one hundred (100) milliliters or not more than ninety (90) milligrams per dosage unit, with an equal or greater quantity of an isoquinoline alkaloid of opium;

2. Not more than one and eight-tenths (1.8) grams of codeine or any of its salts, per one hundred (100) milliliters or not more than ninety (90) milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

3. Not more than one and eight-tenths (1.8) grams of dihydrocodeine or any of its salts, per one hundred (100) milliliters or not more than ninety (90) milligrams per dosage unit, with one or more active, nonnarcotic ingredients in recognized therapeutic amounts;

4. Not more than three hundred (300) milligrams of ethylmorphine or any of its salts, per one hundred (100) milliliters or not more than fifteen (15) milligrams per dosage unit, with one or more ingredients in recognized therapeutic amounts;

5. Not more than five hundred (500) milligrams of opium per one hundred (100) milliliters or per one hundred (100) grams, or not more than twenty-five (25) milligrams per dosage unit, with one or

more active, nonnarcotic ingredients in recognized therapeutic amounts; or

6. Not more than fifty (50) milligrams of morphine or any of its salts, per one hundred (100) milliliters or per one hundred (100) grams with one or more active, nonnarcotic ingredients in recognized therapeutic amounts.

D. The Board of Pharmacy may except by rule any compound, mixture, or preparation containing any stimulant or depressant substance listed in subsections A and B of this section from the application of all or any part of the Uniform Controlled Dangerous Substances Act if the compound, mixture, or preparation contains one or more active medicinal ingredients not having a stimulant or depressant effect on the central nervous system, and if the admixtures are included therein in combinations, quantity, proportion, or concentration that vitiate the potential for abuse of the substances which have a stimulant or depressant effect on the central nervous system.

E. The following hormonal substances or steroids are exempt from classification as Schedule III controlled dangerous substances:

1. Estratest, containing 1.25 mg esterified estrogens and 2.5 mg methyltestosterone;

2. Estratest HS, containing 0.625 mg esterified estrogens and 1.25 mg methyltestosterone;

3. Premarin with Methyltestosterone, containing 1.25 mg conjugated estrogens and 10.0 mg methyltestosterone;

4. Premarin with Methyltestosterone, containing 0.625 mg conjugated estrogens and 5.0 mg methyltestosterone;

5. Testosterone Cypionate - Estrodiol Cypionate injection, containing 50 mg/ml Testosterone Cypionate; and

6. Testosterone Enanthate - Estradiol Valerate injection, containing 90 mg/ml Testosterone Enanthate and 4 mg/ml Estradiol Valerate.

SECTION 6. AMENDATORY 63 O.S. 2011, Section 2-315, is amended to read as follows:

Section 2-315. A. Except as otherwise provided by law, any person required to obtain an annual registration pursuant to Section 2-302 of this title, or any group home, or residential care home as defined by Section 1-820 of this title shall submit for destruction all controlled dangerous substances which are out of date, which are unwanted, unused or which are abandoned by their owner at their facility due to death or other circumstances.

B. All controlled dangerous substances described in subsection A of this section shall be submitted to the Oklahoma City laboratory of the Oklahoma State Bureau of Investigation, along with all required information on forms provided by the Oklahoma State Bureau of Investigation, to the federal Drug Enforcement Administration, to a duly registered reverse distributor, ~~or~~ to the original registered

supplier or their registered agent, to a duly registered retail pharmacy, or to a hospital or clinic with an on-site pharmacy pursuant to the rules set forth in Part 1317 of Title 21 of the Code of Federal Regulations. When any such substance is transported by private contract or common carrier or United States Postal Service for the purpose of destruction, the sender shall require a receipt from such private contract or common carrier or United States Postal Service, and such receipt shall be retained as a permanent record by the sender.

C. Controlled dangerous substances submitted to the Oklahoma State Bureau of Investigation pursuant to the provisions of this section shall be destroyed pursuant to the procedures provided in subsection A of Section 2-508 of this title.

Controlled dangerous substances submitted to any distributors, reverse distributors or their original registered suppliers pursuant to the provisions of this section shall be destroyed by incineration so as to make the substance absolutely unusable for human purposes. An official record listing the property destroyed, the location of destruction and disposal, and the name and title of the person supervising the destruction and disposal shall be submitted to the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control and the federal Drug Enforcement Administration office located nearest the destruction site.

D. The Office of the Chief Medical Examiner is hereby authorized to perform on-site incineration of all controlled dangerous substances which are obtained in the discharge of the official duties of the Chief Medical Examiner. Any record relating to destruction of a controlled dangerous substance shall be maintained as required by the state or federal government and shall be available for inspection by appropriate state or federal government regulatory agencies.

E. This section shall constitute a part of the Uniform Controlled Dangerous Substances Act.

SECTION 7. AMENDATORY 63 O.S. 2011, Section 2-407, is amended to read as follows:

Section 2-407. A. No person shall obtain or attempt to obtain any preparation excepted from the provisions of the Uniform Controlled Dangerous Substances Act pursuant to Section 2-313 of this title in a manner inconsistent with the provisions of paragraph 1 of subsection B of Section 2-313 of this title, or a controlled dangerous substance or procure or attempt to procure the administration of a controlled dangerous substance:

1. By fraud, deceit, misrepresentation, or subterfuge;

2. By the forgery of, alteration of, adding any information to or changing any information on a prescription or of any written order;

3. By the concealment of a material fact; ~~or~~

4. By the use of a false name or the giving of a false address; or

5. By knowingly failing to disclose the receipt of a controlled dangerous substance or a prescription for a controlled dangerous substance of the same or similar therapeutic use from another practitioner within the previous thirty (30) days.

B. Except as authorized by this act, a person shall not manufacture, create, deliver, or possess with intent to manufacture, create, or deliver or possess a prescription form, an original prescription form, or a counterfeit prescription form. This shall not apply to the legitimate manufacture or delivery of prescription forms, or a person acting as an authorized agent of the practitioner.

C. Information communicated to a physician in an effort unlawfully to procure a controlled dangerous substance, or unlawfully to procure the administration of any such drug, shall not be deemed a privileged communication.

D. Any person who violates this section is guilty of a felony punishable by imprisonment for not more than ten (10) years, by a fine of not more than Ten Thousand Dollars ($10,000.00), or by both such fine and imprisonment. A second or subsequent offense under this section is a felony punishable by imprisonment for not less than four (4) years nor more than twenty (20) years, by a fine of

not more than Twenty Thousand Dollars ($20,000.00), or by both such fine and imprisonment.

E. Convictions for second or subsequent violations of this section shall not be subject to statutory provisions for suspended sentences, deferred sentences, or probation.

F. Any person convicted of any offense described in this section shall, in addition to any fine imposed, pay a special assessment trauma-care fee of One Hundred Dollars ($100.00) to be deposited into the Trauma Care Assistance Revolving Fund created in Section ~~1-2522~~ 1-2530.9 of this title.

Passed the House of Representatives the 9th day of March, 2015.

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Presiding Officer of the House
of Representatives

Passed the Senate the ___ day of _____, 2015.

_____
Presiding Officer of the Senate