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STATE OF OKLAHOMA

1 1st Session of the 53rd Legislature (2011)

2 HOUSE BILL 1235 By: Sherrer

3 AS INTRODUCED

4 An Act relating to public health and safety; amending
5 63 O.S. 2001, Sections 2-208, as amended by Section
6 3, Chapter 283, O.S.L. 2005 and 2-212, as last
7 amended by Section 4, Chapter 458, O.S.L. 2010 (63
8 O.S. Supp. 2010, Sections 2-208 and 2-212), which
9 relate to the Uniform Controlled Dangerous Substances
10 Act; adding pseudoephedrine to Schedule III; deleting
11 pseudoephedrine from Schedule V; deleting procedures
12 for sale of certain products; deleting exemptions;
13 deleting defined terms; amending 63 O.S. 2001,
14 Sections 2-309C, as last amended by Section 5,
15 Chapter 458, O.S.L. 2010 and 2-309D, as last amended
16 by Section 3, Chapter 160, O.S.L. 2010 (63 O.S. Supp.
17 2010, Sections 2-309C and 2-309D), which relate to
18 the Anti-Drug Diversion Act; deleting reporting and
19 monitoring requirements for dispensers and
20 registrants who dispense certain product; and
21 providing an effective date.

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

23 SECTION 1. AMENDATORY 63 O.S. 2001, Section 2-208,
24 as amended by Section 3, Chapter 283, O.S.L. 2005 (63 O.S. Supp.

1 2010, Section 2-208), is amended to read as follows:

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

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

9 1. Any substance which contains any quantity of a
10 derivative of barbituric acid, or any salt of a derivative of
11 barbituric acid unless specifically excepted or unless listed in
12 another schedule;

13 2. Chlorhexadol;

14 3. Glutethimide;

15 4. Lysergic acid;

16 5. Lysergic acid amide;

17 6. Methyprylon;

18 7. Sulfondiethylmethane;

19 8. Sulfonethylmethane;

20 9. Sulfonmethane;

21 10. Benzphetamine and its salts;

22 11. Chlorphentermine and its salts;

23 12. Clortermine;

24 13. Mazindol;

25 14. Phendimetrazine;

26 15. Phenylacetone (P2P);

27 16. 1-Phenylcyclohexylamine;

- 1 17. 1-Piperidinocyclohexanecarbo nitrile (PCC);
- 2 18. Ketamine, its salts, isomers, and salts of isomers;
- 3 19. Any material, compound, mixture, or preparation which
- 4 contains any quantity of the following hormonal substances or
- 5 steroids, including their salts, isomers, esters and salts of
- 6 isomers and esters, when the existence of these salts, isomers,
- 7 esters, and salts of isomers and esters is possible within the
- 8 specific chemical designation:
- 9 a. Boldenone,
- 10 b. Chlorotestosterone,
- 11 c. Clostebol,
- 12 d. Dehydrochlormethyltestosterone,
- 13 e. Dihydrotestosterone,
- 14 f. Drostanolone,
- 15 g. Ethylestrenol,
- 16 h. Fluoxymesterone,
- 17 i. Formebolone,
- 18 j. Mesterolone,
- 19 k. Methandienone,
- 20 l. Methandranone,
- 21 m. Methandriol,
- 22 n. Methandrostenolone,
- 23 o. Methenolone,
- 24 p. Methyltestosterone, except as provided in subsection
- 25 E of this section,
- 26 q. Mibolerone,
- 27 r. Nandrolone,

- 1 s. Norethandrolone,
2 t. Oxandrolone,
3 u. Oxymesterone,
4 v. Oxymetholone,
5 w. Stanolone,
6 x. Stanozolol,
7 y. Testolactone,
8 z. Testosterone, except as provided in subsection E of

9 this section, and

10 aa. Trenbolone;

11 20. Tetrahydrocannabinols; ~~[or]~~

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

16 22. Any compound, mixture, or preparation containing any
17 detectable quantity of pseudoephedrine, its salts or optical
18 isomers, or salts of optical isomers.

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

21 B. Nalorphine.

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

25 1. Not more than one and eight-tenths (1.8) grams of
26 codeine or any of its salts, per one hundred (100) milliliters or

27 not more than ninety (90) milligrams per dosage unit, with an

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1 equal or greater quantity of an isoquinoline alkaloid of opium;

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

7 3. Not more than three hundred (300) milligrams of
8 dihydrocodeinone or any of its salts, per one hundred (100)
9 milliliters or not more than fifteen (15) milligrams per dosage
10 unit, with a fourfold or greater quantity of an isoquinoline
11 alkaloid of opium;

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

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

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

27

7. Not more than five hundred (500) milligrams of opium per

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1 one hundred (100) milliliters or per one hundred (100) grams, or
2 not more than twenty-five (25) milligrams per dosage unit, with
3 one or more active, nonnarcotic ingredients in recognized
4 therapeutic amounts; or

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

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

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

23 1. Estratest, containing 1.25 mg esterified estrogens and
24 2.5 mg methyltestosterone;

25 2. Estratest HS, containing 0.625 mg esterified estrogens

26 and 1.25 mg methyltestosterone;

27 3. Premarin with Methyltestosterone, containing 1.25 mg

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1 conjugated estrogens and 10.0 mg methyltestosterone;

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

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

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

9 SECTION 1. AMENDATORY 63 O.S. 2001, Section 2-212,
10 as last amended by Section 4, Chapter 458, O.S.L. 2010 (63 O.S.
11 Supp. 2010, Section 2-212), is amended to read as follows:

12 Section 2-212. ~~[A.]~~ The controlled substances listed in
13 this section are included in Schedule V.

14 ~~[1.]~~ Any compound, mixture, or preparation containing
15 limited quantities of any of the following narcotic drugs, which
16 also contains one or more nonnarcotic active medicinal ingredients
17 in sufficient proportion to confer upon the compound, mixture, or
18 preparation, valuable medicinal qualities other than those
19 possessed by the narcotic drug alone:

20 ~~[a.]~~ ~~[not]~~

21 1. Not more than two hundred (200) milligrams of codeine,
22 or any of its salts, per one hundred (100) milliliters or per one
23 hundred (100) grams~~[r]~~i;

24 ~~[b.]~~ ~~[not]~~

25 2. Not more than one hundred (100) milligrams of

26 dihydrocodeine, or any of its salts, per one hundred (100)
27 milliliters or per one hundred (100) grams~~[7]~~;

7

1 ~~[e.]~~ ~~[not]~~

2 3. Not more than one hundred (100) milligrams of
3 ethylmorphine, or any of its salts, per one hundred (100)
4 milliliters or per one hundred (100) grams~~[7]~~;

5 ~~[d.]~~ ~~[not]~~

6 4. Not more than two and five-tenths (2.5) milligrams of
7 diphenoxylate and not less than twenty-five (25) micrograms of
8 atropine sulfate per dosage unit~~[7]~~; or

9 ~~[e.]~~ ~~[not]~~

10 5. Not more than one hundred (100) milligrams of opium
11 per one hundred (100) milliliters or per one hundred (100) grams.

12 ~~[2. Any compound, mixture, or preparation containing any~~
13 ~~detectable quantity of pseudoephedrine, its salts or optical~~
14 ~~isomers, or salts of optical isomers. If any compound, mixture,~~
15 ~~or preparation as specified in this paragraph is dispensed, sold,~~
16 ~~or distributed in a pharmacy:]~~

17 ~~[a.] [it shall be dispensed, sold, or distributed only~~
18 ~~by, or under the supervision of, a licensed pharmacist or a~~
19 ~~registered pharmacy technician, and]~~

20 ~~[b.] [any person purchasing, receiving, or otherwise~~
21 ~~acquiring any compound, mixture, or preparation shall produce a~~
22 ~~driver license, passport, military identification, or other state-~~
23 ~~issued identification card and shall sign a written log, receipt,~~
24 ~~or other program or mechanism approved by the Oklahoma Bureau of~~

25 ~~Narcotics and Dangerous Drugs Control, showing:]~~

26 ~~[(1) [the date of the transaction,]~~

27 ~~[(2) [name of the purchaser,]~~

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1 ~~[(3) [driver license number, passport, military~~
2 ~~identification, or state issued identification number and state of~~
3 ~~residence of the purchaser,]~~

4 ~~[(4) [name and initials of the pharmacist or pharmacy~~
5 ~~technician conducting the transaction,]~~

6 ~~[(5) [the product being sold, and]~~

7 ~~[(6) [total quantity, in grams or milligrams, of~~
8 ~~pseudoephedrine purchased.]]~~

9 ~~[No person shall purchase, receive, or otherwise acquire more~~
10 ~~than nine (9) grams of any product, mixture, or preparation within~~
11 ~~any thirty day period. Provided, the requirements of this~~
12 ~~subsection shall not apply to any quantity of such product,~~
13 ~~mixture or preparation dispensed pursuant to a valid prescription.]~~

14 ~~[B. The Schedule, as specified in paragraph 2 of subsection~~
15 ~~A, shall not apply to any compounds, mixtures, or preparations~~
16 ~~which are in liquid, liquid capsule, or gel capsule form if~~
17 ~~pseudoephedrine is not the only active ingredient.]~~

18 ~~[C. The Director of the Oklahoma State Bureau of Narcotics~~
19 ~~and Dangerous Drugs Control, by rule, may exempt other products~~
20 ~~from this Schedule which the Director finds are not used in the~~
21 ~~illegal manufacture of methamphetamine or other controlled~~
22 ~~dangerous substances. A manufacturer of a drug product may apply~~
23 ~~for removal of the product from the Schedule if the product is~~
24 ~~determined by the Director to have been formulated in such a way~~

25 ~~as to effectively prevent the conversion of the active ingredient~~
26 ~~into methamphetamine.]~~

27 ~~[D. As used in this section:]~~

9

1 ~~[1. "Gel capsule" means any soft gelatin, liquid filled~~
2 ~~capsule that contains a liquid suspension, which, in the case of~~
3 ~~pseudoephedrine, is suspended in a matrix of glycerin,~~
4 ~~polyethylene glycol, and propylene glycol, along with other liquid~~
5 ~~substances. Regardless of product manufacturer labeling, a~~
6 ~~gelatin covered solid does not constitute a gel capsule under this~~
7 ~~definition; and]~~

8 ~~[2. "Active ingredient" shall include the matrix of~~
9 ~~glycerin, polyethylene glycol, and propylene glycol that is found~~
10 ~~in liquid capsules.]~~

11 SECTION 1. AMENDATORY 63 O.S. 2001, Section 2-309C,
12 as last amended by Section 5, Chapter 458, O.S.L. 2010 (63 O.S.
13 Supp. 2010, Section 2-309C), is amended to read as follows:

14 Section 2-309C. A. A dispenser of a Schedule II, III, IV
15 or V controlled dangerous substance including any compound mixture
16 or preparation containing any detectable quantity of
17 pseudoephedrine, its salts or optical isomers, or salts of optical
18 isomers when dispensed pursuant to a valid prescription shall
19 transmit to a central repository designated by the Oklahoma State
20 Bureau of Narcotics and Dangerous Drugs Control using the American
21 Society for Automation in Pharmacy's (ASAP) Telecommunications
22 Format for Controlled Substances version designated in rules by
23 the Oklahoma State Bureau of Narcotics and Dangerous Drugs

24 Control, the following information for each dispensation:

- 25 1. Recipient's name;
- 26 2. Recipient's address;
- 27 3. Recipient's date of birth;

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- 1 4. Recipient's identification number;
- 2 5. National Drug Code number of the substance dispensed;
- 3 6. Date of the dispensation;
- 4 7. Quantity of the substance dispensed;
- 5 8. Prescriber's United States Drug Enforcement Agency
- 6 registration number;
- 7 9. Dispenser's registration number; and
- 8 10. Other information as required by administrative rule.

9 B. The information required by this section shall be
10 transmitted:

11 1. In a format or other media designated acceptable by the
12 Oklahoma State Bureau of Narcotics and Dangerous Drugs Control;
13 and

14 2. Within twenty-four (24) hours of the time that the
15 substance is dispensed. Beginning January 1, 2012, all
16 information shall be submitted on a real-time log.

17 C. When a prescription is written or dispensed to a
18 resident of a nursing home or a person who is under the care of a
19 hospice program licensed pursuant to the provisions of the
20 Oklahoma Hospice Licensing Act who does not have an identification
21 card issued by the state or another form of a recipient
22 identification number pursuant to Section 2-309B of this title, a
23 Social Security number may be used for the purpose of complying

24 with the reporting requirements provided for in this section.

25 D. The provisions of subsection B of this section shall not
26 apply to a nonresident drug outlet registered pursuant to the
27 Oklahoma Pharmacy Act or to a resident drug outlet as defined in

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1 Section 353.1 of Title 59 of the Oklahoma Statutes if the
2 nonresident or resident drug outlet mails or delivers a controlled
3 substance to a patient or client. Nonresident and resident drug
4 outlets shall transmit the information required in this section
5 within seven (7) days of the date that the controlled substance is
6 dispensed.

7 E. Willful failure to transmit accurate information as
8 required by this section shall be a misdemeanor punishable, upon
9 conviction, by not more than one (1) year in the county jail, or
10 by a fine of not more than One Thousand Dollars (\$1,000.00), or by
11 both such imprisonment and fine, or administrative action may be
12 taken pursuant to Section 2-304 of this title.

13 F. The Director of the Bureau shall have the authority to
14 allow paper submissions on a form designated by the Oklahoma State
15 Bureau of Narcotics and Dangerous Drugs Control, if the dispenser
16 has an appropriate hardship.

17 ~~[G. The Oklahoma State Bureau of Narcotics and Dangerous~~
18 ~~Drugs Control is authorized, by any funds available to it, to~~
19 ~~implement a real time electronic logbook to monitor the sale of~~
20 ~~nonprescription Schedule V products containing any detectable~~
21 ~~quantity of pseudoephedrine, its salts or optical isomers, or~~
22 ~~salts of optical isomers. Dispensers of such pseudoephedrine~~

23 ~~products shall report all such sales electronically pursuant to~~
24 ~~rules promulgated by the Oklahoma State Bureau of Narcotics and~~
25 ~~Dangerous Drugs Control.]~~

26 ~~[H. The Oklahoma State Bureau of Narcotics and Dangerous~~
27 ~~Drugs Control shall have the authority to adopt rules for the~~

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1 ~~reporting of sales of Schedule V product containing any detectable~~
2 ~~quantity of pseudoephedrine, its salts or optical isomers, or~~
3 ~~salts of optical isomers.]~~

4 SECTION 1. AMENDATORY 63 O.S. 2001, Section 2-309D,
5 as last amended by Section 3, Chapter 160, O.S.L. 2010 (63 O.S.
6 Supp. 2010, Section 2-309D), is amended to read as follows:

7 Section 2-309D. A. The information collected at the
8 central repository pursuant to the Anti-Drug Diversion Act shall
9 be confidential and shall not be open to the public. Access to
10 the information shall be limited to:

11 1. Peace officers certified pursuant to Section 3311 of
12 Title 70 of the Oklahoma Statutes who are employed as
13 investigative agents of the Oklahoma State Bureau of Narcotics and
14 Dangerous Drugs Control;

15 2. The United States Drug Enforcement Administration
16 Diversion Group Supervisor;

17 3. The executive director or chief investigator, as
18 designated by each board, of the following state boards:

19 a. Board of Podiatric Medical Examiners,

20 b. Board of Dentistry,

21 c. State Board of Pharmacy,

22 d. State Board of Medical Licensure and Supervision,

23 e. State Board of Osteopathic Examiners,
24 f. State Board of Veterinary Medical Examiners, and
25 g. Oklahoma Health Care Authority;
26 provided, however, that the executive director or chief
27 investigator of each of these boards shall be limited to access to

1 information relevant to licensees of the employing board of such
2 executive director or chief investigator; and

3 4. A multicounty grand jury properly convened pursuant to
4 the Multicounty Grand Jury Act, Sections 350 through 363 of Title
5 22 of the Oklahoma Statutes.

6 B. This section shall not prevent the disclosure, at the
7 discretion of the Director of the Oklahoma Bureau of Narcotics and
8 Dangerous Drugs Control, of investigative information to peace
9 officers and investigative agents of federal, state, county or
10 municipal law enforcement agencies, district attorneys and the
11 Attorney General in furtherance of criminal investigations or
12 prosecutions within their respective jurisdictions, and to
13 registrants in furtherance of efforts to guard against the
14 diversion of controlled dangerous substances.

15 C. Any unauthorized disclosure of any information collected
16 at the central repository provided by the Anti-Drug Diversion Act
17 shall be a misdemeanor. Violation of the provisions of this
18 section shall be deemed willful neglect of duty and shall be
19 grounds for removal from office.

20 D. Notwithstanding the provisions of subsection B,
21 registrants shall have no requirement or obligation to access or

22 check the information in the central repository prior to
23 dispensing or administering medications or as part of their
24 professional practices. Registrants shall not be liable to any
25 person for any claim of damages as a result of accessing or
26 failing to access the information in the central repository and no
27 lawsuit may be predicated thereon. ~~[Nothing herein shall be~~

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1 ~~construed to relieve a registrant from any duty to monitor and~~
2 ~~report the sales of certain products pursuant to subsection E of~~
3 ~~Section 2-309C of this title.]~~

4 E. Information regarding nonfatal overdoses, other than
5 statistical information as required by Section 2-106 of this
6 title, shall be completely confidential. Access to this
7 information shall be strictly limited to the Director of the
8 Oklahoma State Bureau of Narcotics and Dangerous Drugs Control or
9 designee, the Chief Medical Examiner, and the registrant that
10 enters the information. Registrants shall not be liable to any
11 person for a claim of damages for information reported pursuant to
12 the provisions of Section 2-105 of this title.

13 SECTION 1. This act shall become effective November 1,
14 2011.

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