STATE OF OKLAHOMA

2nd Session of the 56th Legislature (2018)

SENATE BILL 1175 By: Brecheen

AS INTRODUCED

An Act relating to the Uniform Controlled Dangerous Substances Act; amending 63 O.S. 2011, Section 2-309C, as last amended by Section 73, Chapter 15, O.S.L. 2013 (63 O.S. Supp. 2017, Section 2-309C), which relates to dispensers of Schedule II, III, IV or V controlled dangerous substances; adding medical marijuana as a substance which requires transmission to central repository; requiring certain facilities to be owned by person authorized to access and transmit to central repository; amending 63 O.S. 2011, Section 2-309D, as last amended by Section 35, Chapter 210, O.S.L. 2016 (63 O.S. Supp. 2017, Section 2-309D), which relates to central repository information; expanding central repository access to certain individuals; and providing a conditional effective date.

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

SECTION 1. AMENDATORY 63 O.S. 2011, Section 2-309C, as last amended by Section 73, Chapter 15, O.S.L. 2013 (63 O.S. Supp. 2017, Section 2-309C), is amended to read as follows:

Section 2-309C. A. A dispenser of a Schedule II, III, IV or V controlled dangerous substance dispensed pursuant to a valid prescription or of medical marijuana shall transmit to a central repository designated by the Oklahoma State Bureau of Narcotics and

Dangerous Drugs Control using the American Society for Automation in Pharmacy's (ASAP) Telecommunications Format for Controlled Substances version designated in rules by the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, the following information for each dispensation:

1. Recipient's and recipient's agent's name;

2. Recipient's and recipient's agent's address;

3. Recipient's and recipient's agent's date of birth;

4. Recipient's and recipient's agent's identification number;

5. National Drug Code number of the substance dispensed;

6. Date of the dispensation;

7. Quantity of the substance dispensed;

8. Prescriber's United States Drug Enforcement Agency registration number;

9. Dispenser's registration number; and

10. Other information as required by administrative rule.

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

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

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

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

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

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

F. Any facility which dispenses medical marijuana shall be owned by a person authorized to access and transmit to the central repository.

SECTION 2. AMENDATORY 63 O.S. 2011, Section 2-309D, as last amended by Section 35, Chapter 210, O.S.L. 2016 (63 O.S. Supp. 2017, Section 2-309D), is amended to read as follows:

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

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

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

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

a. Board of Podiatric Medical Examiners,

b. Board of Dentistry,

c. State Board of Pharmacy,

d. State Board of Medical Licensure and Supervision,

e. State Board of Osteopathic Examiners,

f. State Board of Veterinary Medical Examiners,

g. Oklahoma Health Care Authority,

h. Department of Mental Health and Substance Abuse Services,

i. Board of Examiners in Optometry,

j. Board of Nursing,

k. Office of the Chief Medical Examiner, and

1. State Board of Health;

4. A multicounty grand jury properly convened pursuant to the Multicounty Grand Jury Act;

5. Medical practitioners employed by the United States Department of Veterans Affairs, the United States Military, or other federal agencies treating patients in this state; ~~and~~

6. At the discretion of the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, medical practitioners and their staff, including those employed by the federal government in this state; and

7. Owners of facilities which dispense medical marijuana.

B. This section shall not prevent access, at the discretion of the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, to investigative information by peace officers and investigative agents of federal, state, county or municipal law enforcement agencies, district attorneys and the Attorney General in furtherance of criminal, civil or administrative investigations or prosecutions within their respective jurisdictions, designated legal, communications, and analytical employees of the Bureau, and to registrants in furtherance of efforts to guard against the diversion of controlled dangerous substances.

C. This section shall not prevent the disclosure, at the discretion of the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, of statistical information gathered

from the central repository to the general public which shall be limited to types and quantities of controlled substances dispensed and the county where dispensed.

D. This section shall not prevent the disclosure, at the discretion of the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, of prescription-monitoring-program information to prescription-monitoring programs of other states provided a reciprocal data-sharing agreement is in place.

E. The Department of Mental Health and Substance Abuse Services and the State Department of Health may utilize the information in the central repository for statistical, research, substance abuse prevention, or educational purposes, provided that consumer confidentiality is not compromised.

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

G. 1. Registrants shall have access to the central repository for the purposes of patient treatment and for determination in prescribing or screening new patients. The patient's history may be disclosed to the patient for the purposes of treatment of information at the discretion of the physician.

2. a. Prior to prescribing or authorizing for refill, if one hundred eighty (180) days have elapsed prior to the previous access and check, of opiates, synthetic opiates, semisynthetic opiates, benzodiazepine or carisoprodol to a patient of record, registrants or members of their medical or administrative staff shall be required until October 31, 2020, to access the information in the central repository to assess medical necessity and the possibility that the patient may be unlawfully obtaining prescription drugs in violation of the Uniform Controlled Dangerous Substances Act. The duty to access and check shall not alter or otherwise amend appropriate medical standards of care. The registrant or medical provider shall note in the patient file that the central repository has been checked and may maintain a copy of the information.

b. The requirements set forth in subparagraph a of this paragraph shall not apply:

(1) to medical practitioners who prescribe the controlled substances set forth in subparagraph a of this paragraph for hospice or end-of-life care, or

(2) for a prescription of a controlled substance set forth in subparagraph a of this paragraph that is issued by a practitioner for a patient residing in a nursing facility as defined by Section 1-1902 of this title, provided that the prescription is issued to a resident of such facility.

3. Registrants shall not be liable to any person for any claim of damages as a result of accessing or failing to access the information in the central repository and no lawsuit may be predicated thereon.

H. The State Board of Podiatric Examiners, the State Board of Dentistry, the State Board of Medical Licensure and Supervision, the State Board of Examiners in Optometry, the State Board of Nursing, the State Board of Osteopathic Examiners and the State Board of Veterinary Medical Examiners shall have the sole responsibility for enforcement of the provisions of subsection G of this section. Nothing in this section shall be construed so as to permit the Director of the State Bureau of Narcotics and Dangerous Drugs Control to assess administrative fines provided for in Section 2-304 of this title.

I. The Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, or a designee thereof, shall provide a monthly list to the Directors of the State Board of Podiatric

Examiners, the State Board of Dentistry, the State Board of Medical Licensure and Supervision, the State Board of Examiners in Optometry, the State Board of Nursing, the State Board of Osteopathic Examiners and the State Board of Veterinary Medical Examiners of the top twenty prescribers of controlled dangerous substances within their respective areas of jurisdiction. Upon discovering that a registrant is prescribing outside the limitations of his or her licensure or outside of drug registration rules or applicable state laws, the respective licensing board shall be notified by the Bureau in writing. Such notifications may be considered complaints for the purpose of investigations or other actions by the respective licensing board. Licensing boards shall have exclusive jurisdiction to take action against a licensee for a violation of subsection G of this section.

J. Information regarding fatal and nonfatal overdoses, other than statistical information as required by Section 2-106 of this title, shall be completely confidential. Access to this information shall be strictly limited to the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control or designee, the Chief Medical Examiner, state agencies and boards provided in subsection A of this section, and the registrant that enters the information. Registrants shall not be liable to any person for a claim of damages for information reported pursuant to the provisions of Section 2-105 of this title.

K. The Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control shall provide adequate means and procedures allowing access to central repository information for registrants lacking direct computer access.

L. Upon completion of an investigation in which it is determined that a death was caused by an overdose, either intentionally or unintentionally, of a controlled dangerous substance, the medical examiner shall be required to report the decedent's name and date of birth to the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control. The Oklahoma State Bureau of Narcotics and Dangerous Drugs Control shall be required to maintain a database containing the classification of medical practitioners who prescribed or authorized controlled dangerous substances pursuant to this subsection.

SECTION 3. This act shall become effective upon passage of State Question 788.

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