STATE OF OKLAHOMA

2nd Session of the 58th Legislature (2022)

SENATE BILL 1151 By: Standridge

AS INTRODUCED

An Act relating to the Anti-Drug Diversion Act; amending 63 O.S. 2021, Sections 2-309B and 2-309D, which relate to definitions and central repository information; modifying definition; allowing certain disclosure; and providing an effective date.

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

SECTION 1. AMENDATORY 63 O.S. 2021, Section 2-309B, is amended to read as follows:

Section 2-309B. For the purposes of the Anti-Drug Diversion Act:

1. "Bureau" means the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control;

2. "Dispenser" means a person who distributes a Schedule II controlled dangerous substance, but does not include a licensed hospital pharmacy or a licensed nurse or medication aide who administers such a substance at the direction of a licensed physician;

3. "Dispenser's registration number" means the dispenser's Oklahoma State Bureau of Narcotics and Dangerous Drugs Control registration number or, in the case of a pharmacist, the National Association of Boards of Pharmacy number for the pharmacy where the dispensation is made;

4. "Exception report" means an output of data indicating Schedule II controlled dangerous substance dispensation which is outside expected norms for a prescriber practicing a particular specialty or field of health care, for a dispenser doing business in a particular location, or for a recipient;

5. "Recipient" means the person for whom a prescription is prescribed and who is the lawful intended ultimate user;

6. "Recipient's agent" means a person who is authorized by the ultimate user to pick up the recipient's medication and deliver it to the recipient or a person who claims a prescription other than the person to whom the medication is prescribed;

7. "Recipient's identification number" and "recipient's agent's identification number" means the unique number contained on ~~a valid passport, military identification card, driver license, or identification card issued to a recipient pursuant to Section 6-105 of Title 47 of the Oklahoma Statutes or similar statute of another state if the recipient is not a resident of the State of Oklahoma, or, if the recipient is less than eighteen (18) years old and has no such identification, the unique number contained on a valid~~

~~passport, military identification card, driver license, or identification card issued to the recipient's parent or guardian pursuant to Section 6-105 of Title 47 of the Oklahoma Statutes or similar statute of another state if the parent or guardian is not a resident of the State of Oklahoma, or, if the controlled dangerous substance is obtained for an animal, the unique number contained on the animal owner's valid driver license or identification card issued pursuant to Section 6-105 of Title 47 of the Oklahoma Statutes or similar statute of another state if the owner is not a resident of the State of Oklahoma. Nonresident drug outlets registered pursuant to the Oklahoma Pharmacy Act and resident drug outlets defined in Section 353.1 of Title 59 of the Oklahoma Statutes are exempt from the picture identification requirement if the nonresident and resident drug outlets have obtained the identification of the patient through the prescription benefit plan of the patient~~ forms of identification listed in 8 CFR 274a.2(b)(1)(v)(A) and (B);

8. "Registrant" means a person, persons, corporation or other entity who has been issued by the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control a registration pursuant to Section 2-302 of this title; and

9. "State" means any state, territory, or possession of the United States, the District of Columbia, or foreign nation.

SECTION 2. AMENDATORY 63 O.S. 2021, 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. Oklahoma State Board of Podiatric Medical Examiners,

b. Oklahoma 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. Oklahoma Board of Nursing,

k. Office of the Chief Medical Examiner, and

l. 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;

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. The members of the Opioid Overdose Fatality Review Board for the purpose of carrying out the duties prescribed by Section 2-1001 of this title.

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, tribal, 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~~ for statistical, research, substance abuse prevention, or educational purposes, provided that consumer confidentiality is not compromised.

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 to aid in the determination in prescribing or screening new patients. The physician or designee shall provide, upon request by the patient, the history of the patient or the query history of the patient.

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 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.

4. The failure of a registrant to access and check the central repository as required under state or federal law or regulation may, after investigation, be grounds for the licensing board of the registrant to take disciplinary action against the registrant.

H. The Oklahoma State Board of Podiatric Medical Examiners, the ~~State~~ Oklahoma Board of Dentistry, the State Board of Medical Licensure and Supervision, the ~~State~~ Board of Examiners in

Optometry, the ~~State~~ Oklahoma 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 Oklahoma State Board of Podiatric Medical Examiners, the ~~State~~ Oklahoma Board of Dentistry, the State Board of Medical Licensure and Supervision, the ~~State~~ Board of Examiners in Optometry, the ~~State~~ Oklahoma 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.

M. The Oklahoma State Bureau of Narcotics and Dangerous Drugs Control is authorized to provide unsolicited notification to the licensing board of a pharmacist or practitioner if a patient has received one or more prescriptions for controlled substances in quantities or with a frequency inconsistent with generally recognized standards of safe practice. An unsolicited notification to the licensing board of the practitioner pursuant to this section:

1. Is confidential;

2. May not disclose information that is confidential pursuant to this section; and

3. May be in a summary form sufficient to provide notice of the basis for the unsolicited notification.

SECTION 3. This act shall become effective November 1, 2022.

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