

1 STATE OF OKLAHOMA

2 1st Session of the 55th Legislature (2015)

3 SENATE BILL 693

By: Griffin

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5
6 AS INTRODUCED

7 An Act relating to the Anti-Drug Diversion Act;
8 amending 63 O.S. 2011, Section 2-309D, as last
9 amended by Section 22, Chapter 293, O.S.L. 2014 (63
10 O.S. Supp. 2014, Section 2-309D), which relates to
11 central repository information; deleting obsolete
12 language; permitting certain persons to access
13 certain information; permitting disclosure of certain
14 information for certain purposes; requiring certain
15 persons to access central repository information for
16 certain purposes; and providing an effective date.

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

18 SECTION 1. AMENDATORY 63 O.S. 2011, Section 2-309D, as
19 last amended by Section 22, Chapter 293, O.S.L. 2014 (63 O.S. Supp.
20 2014, Section 2-309D), is amended to read as follows:

21 Section 2-309D. A. The information collected at the central
22 repository pursuant to the Anti-Drug Diversion Act shall be
23 confidential and shall not be open to the public. Access to the
24 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

1 of the Oklahoma State Bureau of Narcotics and Dangerous Drugs
2 Control;

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

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

7 a. Board of Podiatric Medical Examiners,

8 b. Board of Dentistry,

9 c. State Board of Pharmacy,

10 d. State Board of Medical Licensure and Supervision,

11 e. State Board of Osteopathic Examiners,

12 f. State Board of Veterinary Medical Examiners,

13 g. Oklahoma Health Care Authority,

14 h. Department of Mental Health and Substance Abuse
15 Services, and

16 i. State Board of Health;

17 provided, however, that the executive director or chief investigator
18 of each of these boards shall be limited to access to information
19 relevant to licensees of the employing board of such executive
20 director or chief investigator;

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

23 5. The Department of Mental Health and Substance Abuse Services
24 and the State Department of Health for statistical, research,

1 substance abuse prevention or educational purposes provided that the
2 consumer's confidentiality is not compromised; and

3 6. At the discretion of the Director of Narcotics and Dangerous
4 Drugs Control, medical practitioners and their staff, including
5 those employed by the federal government within this state.

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

15 C. This section shall not prevent the disclosure, at the
16 discretion of the Director ~~of the Oklahoma State Bureau~~ of Narcotics
17 and Dangerous Drugs Control, of statistical information gathered
18 from the central repository to the general public which shall be
19 limited to types and quantities of controlled substances dispensed
20 and the county where dispensed.

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

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

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

12 G. Information regarding nonfatal overdoses, other than
13 statistical information as required by Section 2-106 of this title,
14 shall be completely confidential. Access to this information shall
15 be strictly limited to the Director ~~of the Oklahoma State Bureau of~~
16 Narcotics and Dangerous Drugs Control or designee, the Chief Medical
17 Examiner, and the registrant that enters the information.
18 Registrants shall not be liable to any person for a claim of damages
19 for information reported pursuant to the provisions of Section 2-105
20 of this title.

21 H. Upon completion of an investigation in which it is
22 determined that a death was caused by an overdose, either
23 intentionally or unintentionally, of a controlled dangerous
24 substance, the medical examiner shall be required to report the

1 decedent's name and date of birth to the Oklahoma State Bureau of
2 Narcotics and Dangerous Drugs Control. The Oklahoma State Bureau of
3 Narcotics and Dangerous Drugs Control shall be required to maintain
4 a database containing the classification of medical practitioners
5 who prescribed or authorized controlled dangerous substances
6 pursuant to this subsection.

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

12 2. Prior to prescribing or authorizing for refill of all
13 hydrocodone products, all oxycodone products, all benzodiazepines,
14 diazepam, carisiprodal or ultram to the patient of record,
15 registrants or members of their medical or administrative staff
16 shall access and verify the information in the central repository to
17 assess medical necessity and the possibility that the patient may be
18 unlawfully obtaining prescription drugs in violation of the Uniform
19 Controlled Dangerous Substances Act.

20 SECTION 2. This act shall become effective November 1, 2015.

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