1	HOUSE OF REPRESENTATIVES - FLOOR VERSION
2	STATE OF OKLAHOMA
3	2nd Session of the 56th Legislature (2018)
4	COMMITTEE SUBSTITUTE
5	FOR HOUSE BILL NO. 2796 By: Downing, McCall, Sanders,
6	West (Tammy), Blancett, Bush and Frix of the House
7	and
8	Griffin of the Senate
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11	COMMITTEEL CLID CHITTIEL
12	COMMITTEE SUBSTITUTE
13	An Act relating to public health and safety; amending 63 O.S. 2011, Section 2-302, which relates to the
14	Uniform Controlled Dangerous Substances Act; requiring manufacturers and distributers to make certain data available for review to the Director of
15	the Oklahoma State Bureau of Narcotics and Dangerous
16	Drugs Control; requiring manufacturers and distributers to make data available on a monthly basis; making information maintained and provided to
17	the Bureau confidential; allowing certain persons access to information at the discretion of the
18	Director; and providing an effective date.
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21	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:
22	SECTION 1. AMENDATORY 63 O.S. 2011, Section 2-302, is
23	amended to read as follows:
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Section 2-302. A. Every person who manufactures, distributes, dispenses, prescribes, administers or uses for scientific purposes any controlled dangerous substance within or into this state, or who proposes to engage in the manufacture, distribution, dispensing, prescribing, administering or use for scientific purposes of any controlled dangerous substance within or into this state shall obtain a registration issued by the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control, in accordance with rules promulgated by the Director. Persons registered by the Director under Section 2-101 et seq. of this title to manufacture, distribute, dispense, or conduct research with controlled dangerous substances may possess, manufacture, distribute, dispense, or conduct research with those substances to the extent authorized by their registration and in conformity with the other provisions of this article. Every wholesaler, manufacturer or distributor of any drug product containing pseudoephedrine or phenylpropanolamine, or their salts, isomers, or salts of isomers shall obtain a registration issued by the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control in accordance with rules promulgated by the Director and as provided for in Section 2-332 of this title.

B. Out-of-state pharmaceutical suppliers who provide controlled dangerous substances to individuals within this state shall obtain a registration issued by the Director of the Oklahoma State Bureau of

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Narcotics and Dangerous Drugs Control, in accordance with rules

promulgated by the Director; provided that this. This provision

shall not also apply to wholesale distributors who ship distribute

controlled dangerous substances to pharmacies or other entities

registered within this state in accordance with rules promulgated by

the Director.

- C. Beginning November 1, 2018, every manufacturer and distributor required to register under the provisions of this section shall, at such time or times and in such form as the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control may require, make available for review all data required pursuant to federal law, federal rules and regulations and Section 827 of Title 21 of the United States Code. Beginning November 1, 2019, every manufacturer and distributor required to register under the provisions of this section shall provide all data required pursuant to federal law, federal rules and regulations and Section 827 of Title 21 of the United States Code on a monthly basis to the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control in accordance with rules promulgated by the Director.
- D. The information maintained and provided pursuant to subsection C of this section shall be confidential and not open to the public. Access to the information shall, at the discretion of the Director, be limited to:

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- 1 1. Peace officers certified pursuant to the provisions of
 2 Section 3311 of Title 70 of the Oklahoma Statutes who are employed
 3 as investigative agents of the Oklahoma State Bureau of Narcotics
 4 and Dangerous Drugs Control or the Office of the Attorney General;
 - 2. The United States Drug Enforcement Administration Diversion Group Supervisor; and
 - 3. A multicounty grand jury properly convened pursuant to the provisions of the Multicounty Grand Jury Act.
 - E. Manufacturers, distributors, home care agencies, hospices, home care services, and scientific researchers shall obtain a registration annually. Other practitioners shall obtain a registration for a period to be determined by the Director that will be for a period not less than one (1) year nor more than three (3) years.
 - $\frac{P}{F}$. Every trainer or handler of a canine controlled dangerous substances detector who, in the ordinary course of such trainer's or handler's profession, desires to possess any controlled dangerous substance, annually, shall obtain a registration issued by the Director for a fee of Seventy Dollars (\$70.00). Such persons shall be subject to all applicable provisions of Section 2-101 et seq. of this title and such applicable rules promulgated by the Director for those individuals identified in subparagraph a of paragraph 32 of Section 2-101 of this title. Persons registered by the Director pursuant to this subsection may possess controlled dangerous

substances to the extent authorized by their registration and in conformity with the other provisions of this article.

- E. G. The following persons shall not be required to register and may lawfully possess controlled dangerous substances under the provisions of Section 2-101 et seq. of this title:
- 1. An agent, or an employee thereof, of any registered manufacturer, distributor, dispenser or user for scientific purposes of any controlled dangerous substance, if such agent is acting in the usual course of such agent's or employee's business or employment;
- 2. Any person lawfully acting under the direction of a person authorized to administer controlled dangerous substances under Section 2-312 of this title;
- 3. A common or contract carrier or warehouser, or an employee thereof, whose possession of any controlled dangerous substance is in the usual course of such carrier's or warehouser's business or employment;
- 4. An ultimate user or a person in possession of any controlled dangerous substance pursuant to a lawful order of a practitioner;
- 5. An individual pharmacist acting in the usual course of such pharmacist's employment with a pharmacy registered pursuant to the provisions of Section 2-101 et seq. of this title;
 - 6. A nursing home licensed by this state;

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- 8. Registered nurses and licensed practical nurses.
- F. H. The Director may, by rule, waive the requirement for registration or fee for registration of certain manufacturers, distributors, dispensers, prescribers, administrators, or users for scientific purposes if the Director finds it consistent with the public health and safety.
- G. I. A separate registration shall be required at each principal place of business or professional practice where the applicant manufactures, distributes, dispenses, prescribes, administers, or uses for scientific purposes controlled dangerous substances.
- $H.\ J.$ The Director is authorized to inspect the establishment of a registrant or applicant for registration in accordance with rules promulgated by the Director.
- $\overline{\text{H.}}$ No person engaged in a profession or occupation for which a license to engage in such activity is provided by law shall be registered under this act unless such person holds a valid license of such person's profession or occupation.

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J. L. Registrations shall be issued on the first day of
November of each year. Registrations may be issued at other times,
however, upon certification of the professional licensing board.

K. M. The licensing boards of all professions and occupations to which the use of controlled dangerous substances is incidental shall furnish a current list to the Director, not later than the first day of October of each year, of the persons holding valid licenses. All such persons except persons exempt from registration requirements under subsection E G of this section shall be subject to the registration requirements of Section 2-101 et seq. of this title.

L. N. The licensing board of any professional defined as a midlevel practitioner shall notify and furnish to the Director, not
later than the first day of October of each year that such
professional holds a valid license, a current listing of individuals
licensed and registered with their respective boards to prescribe,
order, select, obtain and administer controlled dangerous
substances. The licensing board shall immediately notify the
Director of any action subsequently taken against any such
individual.

M. O. Beginning November 1, 2010, each registrant that prescribes, administers or dispenses methadone shall be required to check the prescription profile of the patient on the central

1	repository of the Oklahoma State Bureau of Narcotics and Dangerous
2	Drugs Control.
3	SECTION 2. This act shall become effective November 1, 2018.
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5	COMMITTEE REPORT BY: COMMITTEE ON JUDICIARY, dated 03/01/2018 - DO
6	PASS, As Amended and Coauthored.
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HB2796 HFLR BOLD FACE denotes Committee Amendments.