STATE OF OKLAHOMA

1st Session of the 50th Legislature (2005)

SENATE BILL 638 By: Adelson

AS INTRODUCED

An Act relating to professions and occupations and public health and safety; amending 59 O.S. 2001, Section 353.24, as last amended by Section 21, Chapter 523, O.S.L. 2004 (59 O.S. Supp. 2004, Section 353.24), which relates to unlawful acts within the Oklahoma Pharmacy Act; merging duplicate section; excluding certain persons from provisions related to possession of dangerous drugs without a valid prescription or license; amending 63 O.S. 2001, Section 2-302, as last amended by Section 59, Chapter 5, O.S.L. 2004 (63 O.S. Supp. 2004, Section 2-302), which relates to controlled dangerous substances registration requirements; excluding specified persons from registration requirements; repealing 59 O.S. 2001, Section 353.24, as last amended by Section 8, Chapter 374, O.S.L. 2004 (59 O.S. Supp. 2004, Section 353.24), which is a duplicate section relating to unlawful acts within the Oklahoma Pharmacy Act; and providing an effective date.

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

SECTION 1. AMENDATORY 59 O.S. 2001, Section 353.24, as last amended by Section 21, Chapter 523, O.S.L. 2004 (59 O.S. Supp. 2004, Section 353.24), is amended to read as follows:

Section 353.24 It shall be unlawful for any person, firm or corporation to:

1. Forge or increase the quantity of drug in any prescription, or to present a prescription bearing forged, fictitious or altered information or to possess any drug secured by such forged, fictitious or altered prescription;

2. Sell, offer for sale, barter or give away any unused quantity of drugs obtained by prescription, except through a program pursuant to the Utilization of Unused Prescription Medications Act or as otherwise provided by the Board of Pharmacy;

3. Sell, offer for sale, barter or give away any drugs damaged by fire, water, or other causes without first obtaining the written approval of the Board or the State Department of Health;

4. Enter into any arrangement whereby prescription orders are received, or prescriptions delivered at a place other than the pharmacy in which they are compounded and dispensed. However, nothing in this paragraph shall prevent a pharmacist or an employee of the pharmacy from personally receiving a prescription or delivering a legally filled prescription at a residence, office or place of employment of the patient for whom the prescription was written. Nothing in this paragraph shall prevent veterinary prescription drugs from being shipped directly from a wholesaler or distributor to a client; provided, such drugs may be dispensed only on prescription of a licensed veterinarian and only when an existing veterinary-client-patient relationship exists;

5. Sell, offer for sale or barter or buy any professional samples <u>except through a program pursuant to the Utilization of Unused Prescription Medications Act</u>. For purpose of this paragraph, "professional samples" means complimentary drugs packaged in accordance with federal and state statutes and regulations and provided to a licensed practitioner free of charge by manufacturers or distributors for the purpose of being distributed free of charge in such package by the licensed practitioner to a patient;

6. Refuse to permit or otherwise prevent members of the Board or such representatives thereof from entering and inspecting any and all places, including premises, equipment, contents, and records, where drugs, medicine, chemicals or poisons are stored, sold, vended, given away, compounded, dispensed or manufactured; or

7. Possess dangerous drugs without a valid prescription or a valid license to possess such drugs<u>; provided, however, this provision shall not apply to any Department of Mental Health and Substance Abuse Services employee or any person whose facility</u>

contracts with the Department of Mental Health and Substance Abuse Services, whose possession of such controlled dangerous substance is solely for the purpose of delivery to the home or residence of a mental health consumer.

SECTION 2. AMENDATORY 63 O.S. 2001, Section 2-302, as last amended by Section 59, Chapter 5, O.S.L. 2004 (63 O.S. Supp. 2004, Section 2-302), is amended to read as follows:

Section 2-302. A. Every person who manufactures, distributes, dispenses, prescribes, administers or uses for scientific purposes any controlled dangerous substance within 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 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 Narcotics and Dangerous Drugs Control, in accordance with rules

promulgated by the Director; provided that this provision shall not apply to wholesale distributors who ship controlled dangerous substances to pharmacies or other entities registered within this state in accordance with rules promulgated by the Director.

C. 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.

D. 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. 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; ~~and~~

7. Registered nurses and licensed practical nurses; and

8. Any Department of Mental Health and Substance Abuse Services employee or any person whose facility contracts with the Department of Mental Health and Substance Abuse Services, whose possession of such controlled dangerous substance is solely for the purpose of delivery to the home or residence of a mental health consumer.

F. 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. 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. The Director is authorized to inspect the establishment of a registrant or applicant for registration in accordance with rules promulgated by the Director.

I. 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.

J. 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. 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 of this section shall be subject to the registration requirements of Section 2-101 et seq. of this title.

L. The licensing board of any professional defined as a mid-level 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.

SECTION 3. REPEALER 59 O.S. 2001, Section 353.24, as last amended by Section 8, Chapter 374, O.S.L. 2004 (59 O.S. Supp. 2004, Section 353.24), is hereby repealed.

SECTION 4. This act shall become effective November 1, 2005.

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