As Amended by House Committee

As Amended by Senate Committee

Session of 2007

SENATE BILL No. 270

By Senators V. Schmidt and D. Schmidt

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AN ACT concerning controlled substances; relating to ephedrine and pseudoephedrine; amending K.S.A. 2006 Supp. 65-1643, 65-4113, 65-4166 and 65-7006 and repealing the existing sections.

Be it enacted by the Legislature of the State of Kansas:

New Section 1. The attorney general shall adopt rules and regulations establishing criteria for self-certifications pursuant to K.S.A. 65-1643, and amendments thereto. The criteria shall provide that a separate certification is required for each place of business at which a licensed pharmacist, registered pharmacy technician or pharmacy intern or clerk supervised by a licensed pharmacist sells a controlled substance designated in subsection (e) or (f) of K.S.A. 65-4113, and amendments thereto. The attorney general shall establish a program regarding such certifications and training through an internet site of the office of the attorney general. Promptly after receiving a certification, the attorney general shall make available a copy of the certification to the Kansas bureau of investigation and local law enforcement officials.

— See. 2. Section 1. K.S.A. 2006 Supp. 65-1643 is hereby amended to read as follows: 65-1643. It shall be unlawful:

(a) For any person to operate, maintain, open or establish any pharmacy within this state without first having obtained a registration from the board. Each application for registration of a pharmacy shall indicate the person or persons desiring the registration, including the pharmacist in charge, as well as the location, including the street name and number, and such other information as may be required by the board to establish the identity and exact location of the pharmacy. The issuance of a registration for any pharmacy shall also have the effect of permitting such pharmacy to operate as a retail dealer without requiring such pharmacy to obtain a retail dealer's permit. On evidence satisfactory to the board: (1) That the pharmacy for which the registration is sought will be conducted in full compliance with the law and the rules and regulations of the board; (2) that the location and appointments of the pharmacy are

such that it can be operated and maintained without endangering the public health or safety; (3) that the pharmacy will be under the supervision of a pharmacist, a registration shall be issued to such persons as the board shall deem qualified to conduct such a pharmacy.

- (b) For any person to manufacture within this state any drugs except under the personal and immediate supervision of a pharmacist or such other person or persons as may be approved by the board after an investigation and a determination by the board that such person or persons is qualified by scientific or technical training or experience to perform such duties of supervision as may be necessary to protect the public health and safety; and no person shall manufacture any such drugs without first obtaining a registration so to do from the board. Such registration shall be subject to such rules and regulations with respect to requirements, sanitation and equipment, as the board may from time to time adopt for the protection of public health and safety.
- (c) For any person to distribute at wholesale any drugs without first obtaining a registration so to do from the board.
- (d) For any person to sell or offer for sale at public auction or private sale in a place where public auctions are conducted, any drugs without first having obtained a registration from the board so to do, and it shall be necessary to obtain the permission of the board in every instance where any of the products covered by this section are to be sold or offered for sale.
- (e) For any person to in any manner distribute or dispense samples of any drugs without first having obtained a permit from the board so to do, and it shall be necessary to obtain permission from the board in every instance where the samples are to be distributed or dispensed. Nothing in this subsection shall be held to regulate or in any manner interfere with the furnishing of samples of drugs to duly licensed practitioners, to mid-level practitioners, to pharmacists or to medical care facilities.
- (f) Except as otherwise provided in this subsection (f), for any person operating a store or place of business to sell, offer for sale or distribute any drugs to the public without first having obtained a registration or permit from the board authorizing such person so to do. No retail dealer who sells 12 or fewer different nonprescription drug products shall be required to obtain a retail dealer's permit under the pharmacy act of the state of Kansas or to pay a retail dealer new permit or permit renewal fee under such act. It shall be lawful for a retail dealer who is the holder of a valid retail dealer's permit issued by the board or for a retail dealer who sells 12 or fewer different nonprescription drug products to sell and distribute nonprescription drugs which are prepackaged, fully prepared by the manufacturer or distributor for use by the consumer and labeled in accordance with the requirements of the state and federal food, drug and

cosmetic acts. Such nonprescription drugs shall not include: (1) A controlled substance; (2) a prescription-only drug; or (3) a drug product intended for human use by hypodermic injection; but such a retail dealer shall not be authorized to display any of the words listed in subsection (u) of K.S.A. 65-1626 and amendments thereto, for the designation of a pharmacy or drugstore.

- (g) For any person to sell any drugs manufactured and sold only in the state of Kansas, unless the label and directions on such drugs shall first have been approved by the board.
- (h) For any person to operate an institutional drug room without first having obtained a registration to do so from the board. Such registration shall be subject to the provisions of K.S.A. 65-1637a and amendments thereto and any rules and regulations adopted pursuant thereto.
- (i) For any person to be a pharmacy student without first obtaining a registration to do so from the board, in accordance with rules and regulations adopted by the board, and paying a pharmacy student registration fee of \$25 to the board.
- (j) For any person to operate a veterinary medical teaching hospital pharmacy without first having obtained a registration to do so from the board. Such registration shall be subject to the provisions of K.S.A. 65-1662 and amendments thereto and any rules and regulations adopted pursuant thereto.
- (k) For any person to sell or distribute in a pharmacy a controlled substance designated in subsection (e) or (f) of K.S.A. 65-4113, and amendments thereto, unless:
- (1) (A) Such controlled substance is sold or distributed by a licensed pharmacist, a registered pharmacy technician or a pharmacy intern or clerk supervised by a licensed pharmacist; $\frac{1}{2}$
- (B) any person purchasing, receiving or otherwise acquiring any such controlled substance produces a photo identification showing the date of birth of the person and signs a log and enters in the log, or allows the seller to enter in the log, such person's address and the date and time of sale. The log or database required by the board shall be available for inspection during regular business hours to the board of pharmacy and any law enforcement officer; or
- (C) the seller determines that the name entered in the log corresponds to the name provided on such identification and that the date and time entered are correct; and
- (D) the seller enters in the log the name of the controlled substance and the quantity sold; \mathbf{or}
 - (2) there is a lawful prescription.; or.
- 42 (3) the pharmacy has submitted to the attorney general a self-certi-43 fication that any licensed pharmacist, registered pharmacy technician or

pharmacy intern or clerk supervised by a licensed pharmacist, employed by such pharmacy has undergone training provided by the seller to ensure that such individuals understand the requirements that apply under this subsection and in accordance with criteria established by the attorney general. The pharmacy shall maintain a copy of such certification and records demonstrating such individuals have undergone the training.

- (l) For any person to sell or distribute in a pharmacy four or more packages or containers of any controlled substance designated in subsection (e) or (f) of K.S.A. 65-4113, and amendments thereto, (1) to a specific customer within any seven-day period. in an amount which exceeds a daily amount of 3.6 grams, without regard to the number of transactions, or (2) unless such controlled substance is packaged in blister packs, each blister containing not more than two dosage units, or where the use of blister packs is technically infeasible, the product is packaged in unit dose packets or pouches.
- —(m) (l) For any pharmacy to allow customers to have direct access to any controlled substance designated in subsection (e) or (f) of K.S.A. 65-4113, and amendments there. Such controlled substance shall be placed behind the counter or stored in a locked cabinet that is located in an area of the pharmacy to which customers do not have direct access.
- (m) A seller who in good faith releases information in a log pursuant to subsection (k) to any law enforcement officer is immune from civil liability for such release unless the release constitutes gross negligence or intentional, wanton or willful misconduct.
- Sec. 3. 2. K.S.A. 2006 Supp. 65-4113 is hereby amended to read as follows: 65-4113. (a) The controlled substances or drugs, by whatever official name, common or usual name, chemical name or brand name designated, listed in this section are included in schedule V.
- (b) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture or preparation containing the following narcotic drug or its salts:

- (c) Any compound, mixture or preparation containing limited quantities of any of the following narcotic drugs which also contains one or more nonnarcotic active medicinal ingredients in sufficient proportion to confer upon the compound, mixture or preparation valuable medicinal qualities other than those possessed by the narcotic drug alone:
- (1) Not more than 200 milligrams of codeine or any of its salts per 100 milliliters or per 100 grams.
- (2) Not more than 100 milligrams of dihydrocodeine or any of its salts per 100 milliliters or per 100 grams.
- 42 (3) Not more than 100 milligrams of ethylmorphine or any of its salts 43 per 100 milliliters or per 100 grams.

- (4) Not more than 2.5 milligrams of diphenoxylate and not less than 25 micrograms of atropine sulfate per dosage unit.
- (5) Not more than 100 milligrams of opium per 100 milliliters or per 100 grams.
- (6) Not more than .5 milligram of difenoxin (9168) and not less than 25 micrograms of atropine sulfate per dosage unit.
- (d) Unless specifically excepted or unless listed in another schedule, any material, compound, mixture or preparation which contains any quantity of the following substances having a stimulant effect on the central nervous system, including its salts, isomers (whether optical, position or geometric) and salts of such isomers whenever the existence of such salts, isomers and salts of isomers is possible within the specific chemical designation:
- (1) Propylhexedrine (except when part of a compound used for nasal decongestion which is authorized to be sold lawfully over the counter without a prescription under the federal food, drug and cosmetic act, so long as it is used only for such purpose)......

- (e) Except as provided in subsection (g), Any compound, mixture or preparation containing any detectable quantity of ephedrine, its salts or optical isomers, or salts of optical isomers.
- (f) Except as provided in subsection (g), Any compound, mixture or preparation containing any detectable quantity of pseudoephedrine, its salts or optical isomers, or salts of optical isomers.
- (g) The scheduling of the substances in subsections (e) and (f) shall not apply to any compounds, mixtures or preparations of ephedrine or pseudoephedrine which are in liquid, liquid capsule or gel capsule form.
- Sec. 4. K.S.A. 2006 Supp. 65-4166 is hereby amended to read as follows: 65-4166. The provisions of this act K.S.A. 21-2501a, 65-1643, 65-4113, 65-4152, 65-4159, 65-7001 and 65-7006, and amendments thereto, and K.S.A. 2006 Supp. 75-722, and amendments thereto, and any rules and regulations promulgated thereunder shall be applicable and uniform throughout this state and in all cities and counties therein. No A city or county shall may enact or enforce any law, ordinance, rule, regulation or resolution in conflict with, in addition to, or supplemental to, the provisions of this act unless expressly authorized by law to do so. more stringent than such provisions. In such cases the more stringent local regulation shall control to the extent of any inconsistency between such regulation and such provisions.
- Sec. 5. 3. K.S.A. 2006 Supp. 65-7006 is hereby amended to read as follows: 65-7006. (a) It shall be unlawful for any person to possess ephedrine, pseudoephedrine, red phosphorus, lithium metal, sodium metal, io-

dine, anhydrous ammonia, pressurized ammonia or phenylpropanolamine, or their salts, isomers or salts of isomers with intent to use the product to manufacture a controlled substance.

- (b) It shall be unlawful for any person to market, sell, distribute, advertise, or label any drug product containing ephedrine, pseudoephedrine, red phosphorus, lithium metal, sodium metal, iodine, anhydrous ammonia, pressurized ammonia or phenylpropanolamine, or their salts, isomers or salts of isomers if the person knows or reasonably should know that the purchaser will use the product to manufacture a controlled substance.
- (c) It shall be unlawful for any person to market, sell, distribute, advertise or label any drug product containing ephedrine, pseudoephedrine, or phenylpropanolamine, or their salts, isomers or salts of isomers for indication of stimulation, mental alertness, weight loss, appetite control, energy or other indications not approved pursuant to the pertinent federal over-the-counter drug final monograph or tentative final monograph or approved new drug application.
- (d) It shall be unlawful for any person to purchase, receive or otherwise acquire at retail any compound, mixture or preparation containing more than 3.6 grams of pseudoephedrine base or ephedrine base in any single transaction or any compound, mixture or preparation containing more than nine grams of pseudoephedrine base or ephedrine base or ephedrine base or by means of shipping through any private or commercial carrier or the postal service 7.5 grams within any thirty-day period of any controlled substance designated in subsection (e) or (f) of K.S.A. 65-4113, and amendments thereto, within any thirty-day period.
- (e) For persons arrested and charged under this section, bail shall be at least \$50,000 cash or surety, unless the court determines on the record that the defendant is not likely to re-offend, the court imposes pretrial supervision or the defendant agrees to participate in a licensed or certified drug treatment program.
- (e) (f) A violation of this section subsection (a), (b) or (c) shall be a drug severity level 2 felony. A violation of subsection (d) shall be a class A nonperson misdemeanor
- Sec. 6. 4. K.S.A. 2006 Supp. 65-1643, 65-4113, 65-4166 and 65-7006 are hereby repealed.
- Sec. 7.5. This act shall take effect and be in force from and after its publication in the statute book.