## **HOUSE BILL No. 2614**

By Committee on Health and Human Services

AN ACT concerning the state board of pharmacy; powers, duties and sections; also repealing K.S.A. 2015 Supp. 65-1637b and 65-1651a. 1669, 65-1676, 65-2837a and 65-4202 and repealing the existing 65-1636, 65-1637, 65-1642, 65-1643, 65-1645, 65-1655, 65-1663, 65-1648, 65-1660 and 65-7007 and K.S.A. 2015 Supp. 65-1626, 65-1627, functions thereof; amending K.S.A. 65-669, 65-1633, 65-1635, 65-

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

follows: 65-1626. For the purposes of this act: Section 1. K.S.A. 2015 Supp. 65-1626 is hereby amended to read as

- or research subject by: injection, inhalation, ingestion or any other means, to the body of a patient (a) "Administer" means the direct application of a drug, whether by
- A practitioner or pursuant to the lawful direction of a practitioner;
- of the practitioner; or 3 the patient or research subject at the direction and in the presence
- thereto. a pharmacist as authorized in K.S.A. 65-1635a, and amendments

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- party logistics provider or dispenser but shall not include a common warehouseman's business. when acting in the usual and lawful course of the carrier's or carrier, public warehouseman or employee of the carrier or warehouseman the direction of a manufacturer, repackager, wholesale distributor, third (b) "Agent" means an authorized person who acts on behalf of or at
- service where the entity controls access to the application and maintains electronic prescription or pharmacy prescription applications as a hosted the software and records on its server. "Application service provider" means an entity that sells

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distributor—has a written agreement currently in effect with the eode, complies with any one of the following: (1) The wholesalewholesale distributor, as defined in section 1504 of the internal revenue when the wholesale distributor, including any affiliated group of the deemed to exist between such wholesale distributor and a manufacturer distribute the manufacturer's prescription drug. An engoing relationship is with whom a manufacturer has established an ongoing relationship to (d) "Authorized distributor of record" means a wholesale distributor

> Prepared by Renae Jefferies Technical Amendment February 12, 2016 Assistant Revisor

**HOUSE HEALTH & HUMAN SERVICES** 

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ATTACHMENT

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every prescription order filled at the pharmacy and a medication profile record system as provided under subsection (d). The book or file of prescription orders shall be kept for a period of not less than five years. The book or file of prescription orders shall at all times be open to inspection by members of the board, the secretary of health and environment, the duly authorized agents or employees of such board or secretary and other proper authorities.

- (c) (1) A medication profile record system shall be maintained in all pharmacies for persons for whom prescriptions are dispensed. The following information shall be recorded: (A) The name and address of the patient for whom the medication is intended; (B) the prescriber's name, the original date the prescription is dispensed and the number or designation identifying the prescription; (C) the name, strength and quantity of the drug dispensed and the name of the dispensing pharmacist; and (D) drug allergies and sensitivities.
- (2) Upon receipt of a prescription order, the pharmacist shall examine the patient's medication profile record before dispensing the medication to determine the possibility of a harmful drug interaction or reaction to medication. Upon recognizing a potential harmful drug interaction or reaction to the medication, the pharmacist shall take appropriate action to avoid or minimize the problem which shall, if necessary, include consultation with the prescriber with documentation of actions taken on the prescription record.
- (3) A medication profile record shall be maintained for a period of not less than five years from the date of the last entry in the record.

- (4) All prescription drug orders communicated by way of electronic transmission shall conform to federal and state laws and the provisions of the board's rules and regulations.
- (d) No registration shall be issued or continued for the conduct of a pharmacy until or unless the provisions of this section have been complied with.
- (e) Each pharmacy Hall comply with the requirements of the federal drug supply chain security act, 21 U.S.C. § 351 et seq.
- Sec. 9. K.S.A. 2015 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

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