Session of 2008

Substitute for SENATE BILL No. 549

By Committee on Public Health and Welfare

2-22

10AN ACT relating to the board of pharmacy; concerning continuous qual-11 ity improvement programs and nonresident pharmacy; amending 12 K.S.A. 65-1657 and repealing the existing section. 13 14Be it enacted by the Legislature of the State of Kansas: 15New Section 1. (a) No later than July 1, 2009, each pharmacy shall establish a continuous quality improvement (CQI) program. The purpose 1617of the CQI program shall be to assess errors that occur in the pharmacy in dispensing or furnishing prescription medications so that the pharmacy 1819may take appropriate action to prevent a recurrence. 20(b) Reports and records Reports, memoranda, proceedings, find-21ings and other records generated as part of a pharmacy's CQI program 22 shall be considered confidential and privileged peer review documents 23 and not subject to discovery, subpoena or other means of legal compulsion 24 for their release to any person or entity and shall not be admissible in any 25civil or administrative action other than an administrative proceeding in-26 itiated by the board of pharmacy. Nothing in this section shall be con-27 strued to prohibit a patient from accessing such patient's own prescription 28records. Nothing in this section shall affect the discoverability of any rec-29 ord not solely generated for or maintained as a part of a pharmacy's CQI 30 program. 31(c) No person in attendance at any meeting being conducted as part 32 of a CQI program shall be compelled to testify in any civil, criminal or 33 administrative action, other than an administrative proceeding initiated 34 by the board of pharmacy as to any discussions or decisions which oc-35 curred as part of the CQI program. 36 (d) All reports and records generated as part of a pharmacy's CQI 37 program shall be available for inspection by the board of pharmacy within 38 a time period established by the board in rules and regulations. 39 In conducting a disciplinary proceeding in which admission of any (e) 40 matters that are confidential and privileged under subsection (b) are pro-41posed, the board of pharmacy shall hold the hearing in closed session 42when any report, record or testimony is disclosed. Unless otherwise pro-43 vided by law, the board of pharmacy in conducting a disciplinary proceeding may close only that portion of the hearing in which disclosure of
 such privileged matters are proposed. In closing a portion of a hearing as
 provided in this subsection, the presiding officer may exclude any person
 from the hearing except **members of the board**, the licensee, the li censee's attorney, the agency's attorney, the witness, the court reporter
 and appropriate staff support for either counsel.

7 The board of pharmacy shall make the portions of the administrative 8 record in which such privileged matters are disclosed subject to a pro-9 tective order prohibiting further disclosure. Such privileged matters shall 10 not be subject to discovery, subpoena or other means of legal compulsion 11 for their release to any person or entity. No person in attendance at a 12closed portion of a disciplinary proceeding shall be required to testify at 13 a subsequent civil, criminal or administrative hearing regarding the priv-14ileged matters, nor shall such testimony be admitted into evidence in any 15subsequent civil, criminal or administrative hearing.

16The board of pharmacy may review peer review committee records, 17any matters that are confidential and privileged under subsection 18(b) testimony or reports in conducting a disciplinary proceeding but must 19prove its findings with independently obtained testimony or records 20which shall be presented as part of the disciplinary proceeding in an open 21meeting of the board of pharmacy. Offering such testimony or records in 22 an open public hearing shall not be deemed a waiver of the peer review 23 privilege relating to any peer review committee testimony, record or 24 report.

(f) The board may establish by rules and regulations requirements
regarding the functions and record keeping of a pharmacy CQI program.
(g) This section shall be part of and supplemental to the pharmacy
act of the state of Kansas.

29 Sec. 2. K.S.A. 65-1657 is hereby amended to read as follows: 65-30 1657. (a) No nonresident pharmacy shall ship, mail or deliver, in any 31 manner, prescription drugs to a patient in this state unless registered 32 under this section as a nonresident pharmacy. Applications for a nonres-33 ident pharmacy registration under this section shall be made on a form 34 furnished by the board. A nonresident pharmacy registration shall be 35 granted for a period of one year upon compliance by the nonresident 36 pharmacy with the provisions of this section and rules and regulations adopted pursuant to this section and upon payment of the registration 37 38 fee established under K.S.A. 65-1645, and amendments thereto, for a 39 pharmacy registration. A nonresident pharmacy registration shall be re-40 newed annually on forms provided by the board, upon compliance by the 41nonresident pharmacy with the provisions of this section and rules and 42regulations adopted pursuant to this section and upon payment of the

43 renewal fee established under K.S.A. 65-1645, and amendments thereto,

1 for the renewal of a pharmacy registration.

2 (b) As conditions for the granting of a registration and for the renewal
3 of a registration for a nonresident pharmacy, the nonresident pharmacy
4 shall comply with the following:

5 (1) Provide information to the board to indicate the person or persons 6 applying for the registration, the location of the pharmacy from which 7 the prescription drugs will be dispensed, the names and titles of all prin-8 cipal owners and corporate officers, if any, and the names of all phar-9 macists dispensing prescription drugs to residents of Kansas;

10 (2) be registered and in good standing in the state in which such 11 pharmacy is located;

(3) maintain, in readily retrievable form, records of prescription drugsdispensed to Kansas patients;

(4) supply upon request, all information needed by the board to carry
out the board's responsibilities under this section and rules and regulations adopted pursuant to this section;

(5) maintain pharmacy hours that permit the timely dispensing of
drugs to Kansas patients and provide reasonable access for the patients
to consult with a licensed pharmacist about such patients' medications;

20 (6) provide toll-free telephone communication consultation between 21 a Kansas patient and a pharmacist at the pharmacy who has access to the 22 patient's records, and ensure that the telephone number(s) will be placed 23 upon the label affixed to each prescription drug container dispensed in 24 Kansas; and

(7) provide to the board such other information as the board mayreasonably request to administer the provisions of this section.

(c) When any nonresident pharmacy fails to supply requested information to the board or fails to respond to proper inquiry of the board,
after receiving notice by certified mail, the board may assess a civil fine
in accordance with the provisions in K.S.A. 65-1658, and amendments
thereto.

32 (e) (d) Each nonresident pharmacy shall comply with the following 33 unless compliance would be in conflict with specific laws or rules and 34 regulations of the state in which the pharmacy is located:

(1) All statutory and regulatory requirements of Kansas for controlled
 substances, including those that are different from federal law;

(2) labeling of all prescriptions dispensed, to include but not be lim-ited to identification of the product and quantity dispensed;

(3) all the statutory and regulatory requirements of Kansas for dis pensing prescriptions in accordance with the quantities indicated by the
 prescriber; and

42 (4) the Kansas law regarding the maintenance and use of the patient43 medication profile record system.

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1 (d) (e) In addition to subsection (c) requirements, each nonresident 2 pharmacy shall comply with all the statutory and regulatory requirements 3 of Kansas regarding drug product selection laws whether or not such 4 compliance would be in conflict with specific laws or rules and regulations $\mathbf{5}$ of the state in which the pharmacy is located, except that compliance 6 which constitutes only a minor conflict with specific laws or rules and 7 regulations of the state in which the pharmacy is located would not be 8 required under this subsection.

9 (c) (f) Each nonresident pharmacy shall develop and provide the 10 board with a policy and procedure manual that sets forth:

11 (1) Normal delivery protocols and times;

(2) the procedure to be followed if the patient's medication is not
available at the nonresident pharmacy, or if delivery will be delayed beyond the normal delivery time;

(3) the procedure to be followed upon receipt of a prescription for
an acute illness, which policy shall include a procedure for delivery of the
medication to the patient from the nonresident pharmacy at the earliest
possible time, or an alternative that assures the patient the opportunity
to obtain the medication at the earliest possible time; and

(4) the procedure to be followed when the nonresident pharmacy is advised that the patient's medication has not been received within the normal delivery time and that the patient is out of medication and requires interim dosage until mailed prescription drugs become available.

24 (f) (g) Except in emergencies that constitute an immediate threat to 25the public health and require prompt action by the board, the board may 26file a complaint against any nonresident pharmacy that violates any pro-27 vision of this section. This complaint shall be filed with the regulatory or 28licensing agency of the state in which the nonresident pharmacy is lo-29 cated. If the regulatory or licensing agency of the state in which the non-30 resident pharmacy is located fails to resolve the violation complained of 31 within a reasonable time, not less than 180 days from the date that the 32 complaint is filed, disciplinary proceedings may be initiated by the board. 33 The board also may initiate disciplinary actions against a nonresident 34 pharmacy if the regulatory or licensing agency of the state in which the 35 nonresident pharmacy is located lacks or fails to exercise jurisdiction.

36 $(\underline{g})(h)$ The board shall adopt rules and regulations that make excep-37 tions to the requirement of registration by a nonresident pharmacy when 38 the out-of-state pharmacy supplies lawful refills to a patient from a pre-39 scription that was originally filled and delivered to a patient within the 40 state in which the nonresident pharmacy is located, or when the prescrip-41 tions being mailed into the state of Kansas by a nonresident pharmacy 42 occurs only in isolated transactions. In determining whether the prescrip-

43 tions being mailed into the state of Kansas by a nonresident pharmacy

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1 are isolated transactions, the board shall consider whether the pharmacy

has promoted its services in this state and whether the pharmacy has a
contract with any employer or organization to provide pharmacy services
to employees or other beneficiaries in this state.

5 (h)(i) It is unlawful for any nonresident pharmacy which is not reg-6 istered under this act to advertise its services in this state, or for any 7 person who is a resident of this state to advertise the pharmacy services 8 of a nonresident pharmacy which has not registered with the board, with 9 the knowledge that the advertisement will or is likely to induce members 10 of the public in this state to use the pharmacy to fill prescriptions. A 11 violation of this section is a class C misdemeanor.

12 (i) (j) Upon request of the board, the attorney general may bring an 13 action in a court of competent jurisdiction for injunctive relief to restrain 14 a violation of the provisions of this section or any rules and regulations 15 adopted by the board under authority of this section. The remedy pro-16 vided under this subsection shall be in addition to any other remedy 17 provided under this section or under the pharmacy act of the state of 18 Kansas.

19 (i) (k) The board may adopt rules and regulations as necessary and 20 as are consistent with this section to carry out the provisions of this 21 section.

22 (k) (l) The executive secretary of the board shall remit all moneys 23 received from fees under this section to the state treasurer in accordance 24 with the provisions of K.S.A. 75-4215, and amendments thereto. Upon 25 receipt of each such remittance, the state treasurer shall deposit the entire 26 amount in the manner specified under K.S.A. 74-1609, and amendments 27 thereto.

28 (1) (m) This section shall be part of and supplemental to the pharmacy 29 act of the state of Kansas.

30 Sec. 3. K.S.A. 65-1657 is hereby repealed.

31 Sec. 4. This act shall take effect and be in force from and after its 32 publication in the statute book.