Approved: 3-10-98

MINUTES OF THE SENATE COMMITTEE ON PUBLIC HEALTH AND WELFARE.

The meeting was called to order by Chair Sandy Praeger at 10:00 a.m. on February 24, 1998 in Room 526-S of the Capitol.

All members were present except:

Committee staff present: Emalene Correll, Legislative Research Department

Robin Kempf, Legislative Research Department

Norman Furse, Revisor of Statutes

Jo Ann Bunten, Committee Secretary

Conferees appearing before the committee:

Others attending: See attached list

Appointment of Subcommittee on <u>SB 622</u> - Providing for licensure of physician assistants and <u>SB 655</u> - Board of Medical Professions

The Chair announced appointment of a subcommittee on <u>SB 622</u> and <u>SB 655</u> with Senator Praeger, Chair, and Senator Hardenburger and Senator Jones as members. Meetings of the subcommittee will be held after the "turnaround" in March.

Action on <u>SB 284</u> - Communications accommodations under the Developmental Disabilities Reform Act

Staff briefed the Committee on the term, "communications accommodations", which is used in SB 284 as a reference to supporting services such as specific types of sign language, Braille, large print and tape recordings of written materials. During Committee discussion it was noted that the definition, "deaf-blindness" in the original bill was dropped because opponents of the bill felt that it would broaden language under the DD Reform Act. Senator Hardenburger felt that Rules and Regulations may resolve concerns as expressed by the entities involved without changing the statutes. It was suggested that the bill not be passed until such Rules and Regs can be reviewed. Senator Hardenburger made a motion that the Committee recommend the bill be reported adversely, seconded by Senator Bleeker. It was pointed out during Committee discussion that the Deaf-Blind Task Force and Envision could submit testimony to the Joint Committee on Administrative Rules and Regulations, and through that process would receive input. It was also noted that if the Joint Committee on Administrative Rules and Regs did not address their specific concerns, then a new bill could be considered next year. The motion carried.

Action on SB 485 - Schedule IV controlled substances.

During Committee discussion on <u>SB 485 and SB 595</u>, a copy of the statute was distributed to the Committee showing what the Board of Pharmacy considers when scheduling, rescheduling or deleting a controlled substance. (Attachment 1) It was pointed out that <u>SB 595</u> would not be considered until the drugs referred to in the bill have been through the review process by the Board of Pharmacy.

Senator Hardenburger made a motion that the Committee recommend SB 485 favorably for passage, seconded by Senator Jones. The motion carried.

Action on <u>SB 507</u> - Pharmacy practice act amendments, civil fines, license suspension or revocation

Staff briefed the Committee on a balloon of <u>SB 507</u> showing proposed amendments relating to application fees and examination fee payments. (Attachment 2)

During Committee discussion concern was expressed relating to raising biennial renewal fees of pharmacists from not more than \$100 to \$200 as shown in the balloon of the bill on page 7, line 17. Senator Hardenburger made a motion that the Committee amend SB 507 as shown in the balloon of the bill except on

CONTINUATION SHEET

MINUTES OF THE SENATE COMMITTEE ON PUBLIC HEALTH AND WELFARE, Room 526-S Statehouse, at 10:00 a.m. on February 24, 1998.

page 7, line 17, by reinserting, "\$100" and striking "\$200", seconded by Senator Steineger. The motion carried.

Senator Hardenburger made a motion that the Committee recommend SB 507 as amended favorably for passage, seconded by Senator Steineger. The motion carried.

Action on SB 533 - Sale of nonprescription medicines and drugs through vending machines

A balloon of <u>SB 533</u> was distributed to the Committee showing a proposed amendment that would require vending machines to be in a climate controlled area, and that the drugs offered for sale or sold in such vending machines should not be older than the manufacturer's expiration date. (Attachment 3)

Senator Steineger made a motion that the Committee adopt the balloon amendments to SB 533, and that the Committee recommend SB 533 as amended favorably for passage, seconded by Senator Hardenburger. The motion carried.

Adjournment

The meeting was adjourned at 11:00 a.m.

The next meeting is scheduled for March 10, 1998.

SENATE PUBLIC HEALTH AND WELFARE COMMITTEE GUEST LIST

DATE: 2-24-98

NAME REPRESENTING	
KETTH R LANDIS	CHRISTIAN SCIENCE COMM.
alison Schmidt	Bothel College, north newton
Shana Budge	Bethel College, North Newton
Katie Ethenne for	Bethel College
Anni Empacher	Bethel Colley
Dibbi Stone	Bethel College
Morica Dask	()
XX	507
Jerho Dam Or	Bohel ouse
Markeliger	Bethel College
Angela Dreier	Bethel College
Krystal Stucky	Bethel College,
Susan anderson	Hein + Weir
Preson Barton	1× CDD
BIU Grosz	SLSM AS
Meg Draper	KM S
Brod Smoot	HLR
LARRY FROELICK	BOARD of PHARMACY
A DOLD LEHM	MADE

SENATE PUBLIC HEALTH AND WELFARE COMMITTEE GUEST LIST /

DATE: 2-24-98

NAME	REPRESENTING
Michael Byington	Envision
Don Cot	Emision.
Rich novis	Emraien
Dick Ourry	Envision
box Acreson	Ks. PHARMACISTS ASSOC.
Bob Willems	Ks. Dipermenists Associ
Miko & AINERS LAMONYON	Modi-XwiX
Dan nelson	Envisian
Cin Bogles	Emvision
Dare Bun	Emision
Rich Cotthice	Health Midwost
Warren Hock	Enflisar
Jeneen Hock	Engelson
Bag Swith	KAPA
Bethy Cranford	Rethel
Gen Red	KSNA
Mold Herzon	PARS
Crang Kalist	KCDD
Wicki Delmidt	Ko. Board of Pharmacy

SENATE PUBLIC HEALTH AND WELFARE COMMITTEE GUEST LIST

NAME	REPRESENTING
John Peterson	Stlde Stanne Missin Helt
	w.
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- (x) "Ultimate user" means a person who lawfully possesses a controlled substance for such person's own use or for the use of a member of such person's household or for administering to an animal owned by such person or by a member of such person's household.
- (y) "Isomer" means all enantiomers and diastereomers.
- (z) "Medical care facility" shall have the meaning ascribed to that term in K.S.A. 65-425 and amendments thereto.

(aa) "Cultivate" means the planting or promotion of growth of five or more plants which contain or can produce controlled substances.

(bb) (1) "Controlled substance analog" means a substance the chemical structure of which is substantially similar to the chemical structure of a controlled substance listed in or added to the schedules designated in K.S.A. 65-4105 or 65-4107 and amendments thereto; and:

(A) Which has a stimulant, depressant or hallucinogenic effect on the central nervous system substantially similar to the stimulant, depressant or hallucinogenic effect on the central nervous system of a controlled substance included in the schedules designated in K.S.A. 65-4105 or 65-4107 and amendments thereto; or

- (B) with respect to a particular individual, which the individual represents or intends to have a stimulant, depressant or hallucinogenic effect on the central nervous system substantially similar to the stimulant, depressant or hallucinogenic effect on the central nervous system of a controlled substance included in the schedules designated in K.S.A. 65-4105 or 65-4107 and amendments thereto.
- (2) "Controlled substance analog" does not include:
 - (A) A controlled substance;

(B) a substance for which there is an approved new drug application;

(C) a substance with respect to which an exemption is in effect for investigational use by a particular person under section 505 of the federal food, drug, and cosmetic act (21 U.S.C. 355) to the extent conduct with respect to the substance

is permitted by the exemption; or
(D) any substance to the extent not intended for human consumption before an exemption takes effect with respect to the substance.

History: L. 1972, ch. 234, § 1; L. 1974, ch. 258, § 1; L. 1975, ch. 332, § 1; L. 1980, ch. 195, § 1; L. 1985, ch. 214, § 2; L. 1989, ch. 192, § 4;

L. 1990, ch. 100, § 7; L. 1994, ch. 160, § 1; July

CASE ANNOTATIONS

24. Release of respondent attorney from court-ordered probation noted for compliance with order regarding use or possession of controlled substances. In re McKenna, 251 K. 615, 840 P.2d 412 (1992).

25. Circumstantial and direct evidence sufficient to prove defendant guilty of cultivation of marijuana. State v. Heiskell, 21 K.A.2d 105, 111, 896 P.2d 1106 (1995).

65-4102. Board of pharmacy to administer act; authority to control; report to speaker of house and president of senate on substances proposed for scheduling, rescheduling or deletion; scheduling of the controlled substance analog. (a) The board shall administer this act and may adopt rules and regulations relating to the registration and control of the manufacture, distribution and dispensing of controlled substances within this state. All rules and regulations of the board shall be adopted in conformance with article 4 of chapter 77 of the Kansas Statutes Annotated and the procedures prescribed by this act.

(b) Annually, the board shall submit to the speaker of the house of representatives and the president of the senate a report on substances proposed by the board for scheduling, rescheduling or deletion by the legislature with respect to any one of the schedules as set forth in this act, and reasons for the proposal shall be submitted by the board therewith. In making a determination regarding the proposal to schedule, reschedule or delete a substance, the board shall consider the following:

The actual or relative potential for abuse;

(2) the scientific evidence of its pharmacological effect, if known;

(3) the state of current scientific knowledge regarding the substance;

(4) the history and current pattern of abuse;

(5) the scope, duration and significance of abuse;

(6) the risk to the public health;

(7) the potential of the substance to produce psychological or physiological dependence liability; and

(8) whether the substance is an immediate precursor of a substance already controlled under this article.

(c) The board shall not include any nonnarcotic substance within a schedule if such substance may be lawfully sold over the counter without a prescription and cosmetic act.

(d) Authority does not extend to erages or tobacco

(e) Upon rece Supp. 65-4105a, 1 ing of the controllgency basis purscheduling of a si expires one year a uling rule. With r minent hazard to: consider whether uled on a tempo: factors set forth in and may also con manufacture or d formation concerin subsection (b). der this subsection lemaking proceed respect to the sui this subsection lar rulemaking proces (a) with respect to

History: L. 1 258, § 2; L. 1982, § 2; July 1.

65-4105. St. ule I. (a) The consection are include set forth opposite DEA controlled stassigned to it.

(b) Any of the their isomers, estermers, esters and cepted, whenever esters, ethers and cific chemical designation.

- (1) Acetyl-alpha-me 2-phenethyl)-4phenylacetamid
- (2) Acetylmethadol
- (3) Allylprodine (4) Alphacetylmeth: (except levo-alp levo-alpha-acety tate or LAAM)
- (5) Alphameproding
- (6) Alphamethadol

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out a prescription under the federal food, drug and cosmetic act.

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iarubith(d) Authority to control under this section does not extend to distilled spirits, wine, malt beverages or tobacco.

(e) Upon receipt of notice under K.S.A. 1997 Supp. 65-4105a, the board shall initiate scheduling of the controlled substance analog on an emergency basis pursuant to this subsection. The scheduling of a substance under this subsection expires one year after the adoption of the scheduling rule. With respect to the finding of an imminent hazard to the public safety, the board shall consider whether the substance has been scheduled on a temporary basis under federal law or factors set forth in subsections (b)(4), (5) and (6), and may also consider clandestine importation, manufacture or distribution, and if available, information concerning the other factors set forth in subsection (b). A rule may not be adopted under this subsection until the board initiates a rulemaking proceeding under subsection (a) with respect to the substance. A rule adopted under this subsection lapses upon the conclusion of the rulemaking proceeding initiated under subsection (a) with respect to the substance.

History: L. 1972, ch. 234, § 2; L. 1974, ch. 258, § 2; L. 1982, ch. 269, § 1; L. 1994, ch. 160, § 2; July 1.

65-4105. Substances included in schedule I. (a) The controlled substances listed in this section are included in schedule I and the number set forth opposite each drug or substance is the DEA controlled substances code which has been assigned to it.

(b) Any of the following opiates, including their isomers, esters, ethers, salts, and salts of isomers, esters and ethers, unless specifically excepted, whenever the existence of these isomers, esters, ethers and salts is possible within the specific chemical designation:

(1)	Acetyl-alpha-methylfentanyl (N-[1-(1-methyl-2-phenethyl)-4-piperidinyl] -N-	
	phenylacetamide	9815
(2)	Acetylmethadol	9601
	Allylprodine	9602
(4)	Alphacetylmethadol	9603
	tate or LAAM)	
(5)	Alphameprodine	9604
(6)	Alphamethadol	9605

(7)	Alpha-methylfentanyl (N-[1-(alpha-methyl-	
	beta-phenyl)ethyl-4-piperidyl] propionanilide;	
	1-(1-methyl-2-phenylethyl)-4-(N-propanilido)	0014
(8)	piperidine) Alpha-methylthiofentanyl (N-[1-methyl-2-(2-	9814
(0)	thienyl) ethyl-4-piperidinyl] -N-	
	phenylpropanamide)	9832
(9)	Benzethidine	9606
(10)	Betacetylmethadol	9607
(11)	Beta-hydroxyfentanyl (N-[1-(2-hydroxy-2-phe-	0000
(12)	nethyl) 4-piperidinyl] -N-phenylpropanamide Beta-hydroxy-3-methylfentanyl (other name:	9830
(12)	N-[1-(2-hydroxy-2-phenethyl)-3-methyl -4-pi-	
	peridinyl]-N-phenylpropanamide	9831
(13)	Betameprodine	9608
(14)	Betamethadol	9609
(15)	Betaprodine	9611
(16) (17)	Clonitazene Dextromoramide	9612
(18)	Diampromide	9613 9615
(19)	Diethylthiambutene	9616
(20)	Difenoxin	9168
(21)	Dimenoxadol	9617
(22)	Dimepheptanol	9618
(23)	Dimethylthiambutene	9619
(24) (25)	Dioxaphetyl butyrate Dipipanone	9621 9622
(26)	Ethylmethylthiambutene	9623
(27)	Etonitazene	9624
(28)	Etoxeridine	9625
(29)	Furethidine	9626
(30)	Hydroxypethidine	9627
(31) (32)	Ketobemidone	9628 9629
(33)	Levomoramide	9631
(34)	3-Methylfentanyl (N-[3-methyl-1-(2-phenyle-	0001
88 88	thyl)-4-piperidyl] -N-phenylpropanamide)	9813
(35)	3-Methylthiofentanyl (N-[(3-methyl-1-(2-	
	thienyl)ethyl-4-piperidinyl]-	
(36)	N-phenylpropanamide)	9833
(37)	▲	9632
(01)	propionoxypiperidine)	9661
(38)	Noracymethadol	9633
(39)	Norlevorphanol	9634
(40)		9635
(41) (42)		9636
(42)	(2-phenethyl) -4-piperidinyl] propanamide	9812
(43)		0012
	acetoxypiperidine)	9663
(44)		9637
(45)		9638
(46)		9647
(47) (48)	.	9641
(49)		9642 9643
(50)		9644
(51)	Propiram	9649
(52)	Racemoramide	9645
(53)	Thiofentanyl (N-phenyl-N-[1-(2-thienyl)ethyl-	
15 11	4- piperidinyl]-propanamide	9835
(54)		9750 9646

SENATE BILL No. 507

By Committee on Public Health and Welfare

1-27

AN ACT concerning the state board of pharmacy; relating to grounds for disciplinary actions; concerning civil fines; licensure; amending K.S.A. 65-1627f, 65-1632 and 65-1645 and K.S.A. 1997 Supp. 65-1627 and 65-1658 and repealing the existing sections.

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

Section 1. K.S.A. K.S.A. 1997 Supp. 65-1627 is hereby amended to read as follows: 65-1627. (a) The board may revoke, suspend, place in a probationary status or deny a renewal of any license of any pharmacist upon a finding that:

- (1) The license was obtained by fraudulent means;
- (2) the licensee has been convicted of a felony and the licensee fails to show that the licensee has been sufficiently rehabilitated to warrant the public trust;
- (3) the licensee is found by the board to be guilty of unprofessional conduct or professional incompetency;
- (4) the licensee is addicted to the liquor or drug habit to such a degree as to render the licensee unfit to practice the profession of pharmacy;
- (5) the licensee has violated a provision of the federal or state food, drug and cosmetic act, the uniform controlled substances act of the state of Kansas, or any rule and regulation adopted under any such act;
- (6) the licensee is found by the board to have filled a prescription not in strict accordance with the directions of the practitioner;
- (7) the licensee is found to be mentally or physically incapacitated to such a degree as to render the licensee unfit to practice the profession of pharmacy;
- (8) the licensee has violated any of the provisions of the pharmacy act of the state of Kansas or any rule and regulation adopted by the board pursuant to the provisions of such pharmacy act;
- (9) the licensee has failed to comply with the requirements of the board relating to the continuing education of pharmacists;
- (10) the licensee as a pharmacist in charge or consultant pharmacist der the provisions of subsection (c) or (d) of K.S.A. 65-1648 and endments thereto has failed to comply with the requirements of subsection (c) or (d) of K.S.A. 65-1648 and amendments thereto;

Proposed Amendments Mark-up Application fee? Examination fee payment?

Senate Public Health and Welfare Date: ユニスダーア8 Attachment No. カ

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(11) the licensee has knowingly submitted a misleading, deceptive, untrue or fraudulent misrepresentation on a claim form, bill or statement;

(12) the licensee has had a license to practice pharmacy revoked, suspended or limited, has been censured or has had other disciplinary action taken, or voluntarily surrendered the license after formal proceedings have been commenced, or has had an application for license denied, by the proper licensing authority of another state, territory, District of Columbia or other country, a certified copy of the record of the action of the other jurisdiction being conclusive evidence thereof; or

(13) the licensee has self-administered any controlled substance with-

out a practitioner's prescription order.

(b) In determining whether or not the licensee has violated subsection (a)(3), (a)(4), (a)(7) or (a)(13), the board upon reasonable suspicion of such violation has authority to compel a licensee to submit to mental or physical examination or drug screen, or any combination thereof, by such persons as the board may designate. To determine whether reasonable suspicion of such violation exists, the investigative information shall be presented to the board as a whole. Information submitted to the board as a whole and all reports, findings and other records shall be confidential and not subject to discovery by or release to any person or entity. The licensee shall submit to the board a release of information authorizing the board to obtain a report of such examination or drug screen, or both. A person affected by this subsection shall be offered, at reasonable intervals, an opportunity to demonstrate that such person can resume the competent practice of pharmacy with reasonable skill and safety to patients. For the purpose of this subsection, every person licensed to practice pharmacy and who shall accept the privilege to practice pharmacy in this state by so practicing or by the making and filing of an annual biennial renewal to practice pharmacy in this state shall be deemed to have consented to submit to a mental or physical examination or a drug screen, or any combination thereof, when directed in writing by the board and further to have waived all objections to the admissibility of the testimony, drug screen or examination report of the person conducting such examination or drug screen, or both, at any proceeding or hearing before the board on the ground that such testimony or examination or drug screen report constitutes a privileged communication. In any proceeding by the board pursuant to the provisions of this subsection, the record of such board proceedings involving the mental and physical examination or drug screen, or any combination thereof, shall not be used in any other administrative or judicial proceeding.

(c) The board may suspend, revoke, place in a probationary status or hy a renewal of any retail dealer's permit issued by the board when information in possession of the board discloses that such operations for

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application

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which the permit was issued are not being conducted according to law or the rules and regulations of the board.

- (d) The board may revoke, suspend, place in a probationary status or deny a renewal of the registration of a pharmacy upon a finding that: (1) Such pharmacy has been operated in such manner that violations of the provisions of the pharmacy act of the state of Kansas or of the rules and regulations of the board have occurred in connection therewith, (2) the owner or any pharmacist employed at such pharmacy is convicted, subsequent to such owner's acquisition of or such employee's employment at such pharmacy, of a violation of the pharmacy act or uniform controlled substances act of the state of Kansas, or the federal or state food, drug and cosmetic act; of (3) the owner or any pharmacist employed by such pharmacy has fraudulently claimed money for pharmaceutical services; or (4) the registrant has had a registration revoked, suspended or limited, has been censured or has had other disciplinary action taken, or an application for registration denied, by the proper registering authority of another state, territory, District of Columbia or other country, a certified copy of the record of the action of the other jurisdiction being conclusive evidence thereof.
- (e) A registration to manufacture or to distribute at wholesale a drug or a registration for the place of business where any such operation is conducted may be suspended, revoked, placed in a probationary status or the renewal of such registration may be denied by the board upon a finding that the registrant or the registrant's agent: (1) Has materially falsified any application filed pursuant to or required by the pharmacy act of the state of Kansas; (2) has been convicted of a felony under any federal or state law relating to the manufacture or distribution of drugs; (3) has had any federal registration for the manufacture or distribution of drugs suspended or revoked; (4) has refused to permit the board or its duly authorized agents to inspect the registrant's establishment in accordance with the provisions of K.S.A. 65-1629 and amendments thereto; (5) has failed to keep, or has failed to file with the board or has falsified records required to be kept or filed by the provisions of the pharmacy act of the state of Kansas or by the board's rules and regulations; or (6) has violated the pharmacy act of the state of Kansas or rules and regulations adopted by the state board of pharmacy under the pharmacy act of the state of Kansas or has violated the uniform controlled substances act or rules and regulations adopted by the state board of pharmacy under the uniform controlled substances act.
- (f) Orders under this section, and proceedings thereon, shall be subject to the provisions of the Kansas administrative procedure act.
- Sec. 2. K.S.A 65-1627f is hereby amended to read as follows: 65-1627f. (a) Depositions may be used by either party. Upon the completion



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hearing held hereunder, the board shall have the power to enter an order of revocation, suspension, probation or denial of the renewal of a license, registration or permit. The license, registrant or permit holder shall not engage in the activity authorized by such license, registration or permit after a license, registration or permit is revoked or the renewal thereof denied or during the time for which it is suspended. If a license, registration or permit is suspended or placed on probation, the suspension or probation shall be for a definite period of time to be fixed by the board, and the license, registration or permit shall be reinstated and any limitations or conditions thereon removed upon the expiration of such period if all annual terminal renewal fees have been paid. If such license, registration or permit is revoked, such revocation shall be for all time, except that at any time after the expiration of one year, application may be made for reinstatement of any license, registrant or permit holder whose license, registration or permit shall have been revoked, and such application shall be addressed to the executive secretary of the board. Such application shall be processed in accordance with the provisions of the Kansas administrative procedure act.

(b) All final orders entered in any proceeding shall be the action of

the board with a quorum present at such meeting.

Sec. 3. K.S.A. 65-1632 is hereby amended to read as follows: 65-1632. (a) Each license of to practice as a pharmacist issued by the board shall expire for the period ending on June 30 following the date of issuance, 1998 shall be renewed in accordance with this section. Each application for renewal of a license as a pharmacist shall be made on a form prescribed and furnished by the board. Except as otherwise provided in this subsection, the application, when accompanied by the renewal fee and received by the executive secretary of the board on or before the date of expiration of the license, shall have the effect of temporarily renewing the applicant's license until actual issuance or denial of the renewal. If at the time of filing a proceeding is pending before the board which may result in the suspension, probation, revocation or denial of the applicant's license, the board may by emergency order declare that the application for renewal shall not have the effect of temporarily renewing such applicant's license. Every licensed pharmacist shall pay to the secretary of the board annually biennially a renewal fee fixed by the board as provided in K.S.A. 65-1645 and amendments thereto. The renewal fee fixed by the board under this section immediately prior to the effective date of the act of which this section is amendatory shall continue in effect uni different renewal fee is fixed by the board by rules and regulations ded in K.S.A. 65-1645 and amendments thereto.

Gommencing on June 30, 1998, each holder of a pharmacist license, which is numbered with an even number, who is qualified under shall expire on June 30 of the year specified by the board for the expiration of the license and

on a biennial basis

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KS.A. 65-1631, and amendments thereto, shall have such license to practice issued or renewed for a period which shall expire on June 30, 2000. Commencing on June 30, 2000, in order to issue or renew such holder's license to practice on a biennial basis, each such even-pumbered license shall be issued or renewed for a period ending on the June 30 which occurs in the next even-numbered year.

(c) Commencing on June 30, 1998, each holder of a license to practice as a pharmacist, which is numbered with an odd number, who is qualified under 65-1631 and amendments thereto, shall have such license issued or renewed for a period which shall expire on June 30, 1999. Commencing on June 30, 1999 in order to issue or renew licenses to practice for such holders on a biennial basis, each such odd-numbered license shall be issued of renewed for a period ending on the June 30 which occurs in the next odd-numbered yeak?

(e) (d) The board may deny renewal of any license of a pharmacist on any ground which would authorize the board to deny an initial application for licensure or on any ground which would authorize the board to suspend, revoke or place on probation a license previously granted. Orders under this section, and proceedings thereon, shall be subject to

the provisions of the Kansas administrative procedure act.

[(e)] The payment of the renewal fee by a person who is a holder of a [(d) license as a pharmacist shall entitle the person to renewal of license if no grounds exist for denying the renewal of the license and if the person has furnished satisfactory evidence to the board that the person has successfully complied with the rules and regulations of the board relating to continuing professional education. These educational requirements shall be fixed by the board at not less than 40 20 clock hours nor more than 20 40 clock hours unnually biennially of a program of continuing education approved by the board. The maximum number of continuing education hours required by the board to meet the requirements for cancellation of inactive status licensure and renewal of license under subsection tal (f) or reinstatement of license because of nonpayment of fees under subsection (e) (g) shall not exceed 30 60.

(d) (f) The payment of the renewal fee by the person who is a holder (e) of a license as a pharmacist but who has not complied with the continuing education requirements fixed by the board, if no grounds exist for denying the renewal of the license other than that the person has not complied with the continuing education requirements fixed by the board, shall entitle the person to inactive status licensure by the board. No person holding an inactive status license from the board shall engage in the practice f pharmacy in this state. Upon furnishing satisfactory evidence to the ard of compliance with the continuing education requirements fixed by the board and upon the payment to the board of all applicable fees, a

Commencing with the renewal of licenses which expire on June 30, 1998, each license shall be renewed on a biennial basis. To provide for a system of biennial renewal of licenses, the board may provide rules by that licenses issued or renewed regulations may expire less than two years from the date of issuance or renewal.

Continuing education hours may be prorated for licensure periods which are less than biennial in accordance with rules and regulations of the board.

Policy: CE hours for cancellation of inactive status licensure and of renewal license.

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person holding an inactive status license from the board shall be entitled to cancellation of the inactive status license and to renewal of licensure as a pharmacist.

(f)

(e) (g) If the renewal fee for any pharmacist's license has not been paid by August 1 of any the renewal year, the license is hereby declared void, and no license shall be reinstated except upon payment of any unpaid renewal fee plus a penalty fee fixed by the board as provided in K.S.A. 65-1645 and amendments thereto and proof satisfactory to the board of compliance with the continuing education requirements fixed by the board. The penalty fee established by this section immediately prior to the effective date of the act shall continue in effect until a different penalty fee is fixed by the board by rules and regulations as provided in K.S.A. 65-1645 and amendments thereto. Payment of any unpaid renewal fee plus a penalty fee and the submission of proof satisfactory to the board of compliance with the continuing education requirements fixed by the board shall entitle the license to be reinstated. The nonpayment of renewal fees by a previously licensed pharmacist for a period exceeding three years shall not deprive the previously licensed pharmacist of the right to reinstate the license upon the payment of any unpaid fees and penalties and upon compliance with the continuing education requirements fixed by the board, except that the board may require such previously licensed pharmacist to take and pass an examination approved by the board for reinstatement as a pharmacist and to pay any applicable examination fee.

Sec. 4. K.S.A. 65-1645 is hereby amended to read as follows: 65-1645. (a) Application for licenses registrations or permits under K.S.A. 65-1643 and amendments thereto shall be made on a form prescribed and furnished by the board. Applications for registration to distribute at wholesale any drugs shall contain such information as may be required by the board in accordance with the provisions of K.S.A. 65-1655 and amendments thereto. The application shall be accompanied by the fee prescribed by the board under the provisions of this section. When such application and fees are received by the executive secretary of the board on or before the due date, such application shall have the effect of temporarily renewing the applicant's registration or permit until actual issuance or denial of the renewal. However, if at the time of filing a proceeding is pending before the board which may result in the suspension, probation, revocation or denial of the applicant's registration or permit, the board may declare, by emergency order, that such application for renewal shall not have the effect of temporarily renewing such applicant's registration or permit. Separate applications shall be made and separate registrations or permits issued for each separate place at which is carried on any of the operations for which a registration or permit is required by

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Renewal applications for pharmacist, $65\text{--}1632 \rho$

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K.S.A. 65-1643 and amendments thereto except that the board may provide for a single registration for a business entity registered to manufacture any drugs or registered to distribute at wholesale any drugs and operating more than one facility within the state, or for a parent entity with divisions, subsidiaries or affiliate companies, or any combination thereof, within the state when operations are conducted at more than one location and there exists joint ownership and control among all the entities.

- (b) The fees required for the issuing of the licenses, registrations or permits required by K.S.A. 65 1643 and amendments thereto shall be fixed by the board as herein provided, subject to the following:
- (1) Pharmacy, new registration not more than \$150, renewal not more than \$125;
 - (2) pharmacist, examination fee not more than \$350;
- (3) pharmacist, examination fee for previously licensed pharmacist not more than \$250;
 - (4) pharmacist renewal fee not more than \$100 \$200;
- (5) pharmacist, evaluation fee not more than \$250;
- (6) pharmacist, reciprocal licensure fee not more than \$250;
- (7) pharmacist, penalty fee, not more than \$250 \$500;
- (8) manufacturer, new registration not more than \$500, renewal not more than \$400;
- (9) wholesaler, new registration not more than \$500, renewal not more than \$400, except that a wholesaler dealing exclusively in nonprescription drugs, the manufacturing, distributing or dispensing of which does not require registration under the uniform controlled substances act, shall be assessed a fee for registration and reregistration not to exceed \$50;
 - (10) special auction not more than \$50;
 - (11) samples distribution not more than \$50;
- (12) institutional drug room, new registration not more than \$40, renewal not more than \$35;
- (13) retail dealer selling more than 12 different nonprescription drug products, new permit not more than \$12, renewal not more than \$12; or
- (14) certification of grades for each applicant for examination and registration not more than \$25.
- (c) For the purpose of fixing fees, the board may establish classes of retail dealers' permits for retail dealers selling more than 12 different nonprescription drug products, and the board may fix a different fee for each such class of permit.
- (d) The board shall determine annually the amount necessary to carry but and enforce the provisions of this act for the next ensuing fiscal year and shall fix by rules and regulations the fees authorized for such year at

under the pharmacy act of the state of Kansas

Pharmacist license application fee?

biennial

Stay

Policy: Penalty fee charge. Current penalty . fee fixed at \$150. 68-11-1.

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the sum deemed necessary for such purposes. The fees fixed by the board under this section immediately prior to the effective date of this act shall continue in effect until different fees are fixed by the board by rules and regulations as provided under this section.

(e) The board may deny renewal of any registration or permit required by K.S.A. 65-1643 and amendments thereto on any ground which would authorize the board to suspend, revoke or place on probation a registration or permit previously granted pursuant to the provisions of K.S.A. 65-1643 and amendments thereto. Registrations and permits issued under the provisions of K.S.A. 65-1643 and 65-1644 and amendments thereto shall be conspicuously displayed in the place for which the registration or permit was granted. Such registrations or permits shall not be transferable. All such registrations and permits except retail dealer permits shall expire on June 30 following date of issuance. Retail dealers' permits shall expire on the last day of February. All registrations and permits shall be renewed annually. Application blanks for renewal of registrations and permits shall be mailed by the board to each registrant or permittee at least 30 days prior to expiration of the registration or permit. If application for renewal is not made before 30 days after such expiration, the existing registration or permit shall lapse and become null and void on the date of its expiration, and no new registration or permit shall be granted except upon payment of the required renewal fee plus a penalty equal to the renewal fee. Failure of any registrant or permittee to receive such application blank shall not relieve the registrant or permittee from the penalty hereby imposed if the renewal is not made as prescribed.

Sec. 5. K.S.A. 1997 Supp. 65-1658 is hereby amended to read as follows: 65-1658. The state board of pharmacy, in addition to any other penalty prescribed under the pharmacy act of the state of Kansas, may assess a civil fine, after notice and an opportunity to be heard in accordance with the Kansas administrative procedure act, against any licensee or registrant under subsections (a), (c), (d) and (e) of K S.A. 65-1627 and amendments thereto for violation of the pharmacy act of the state of Kansas or rules and regulations of the state board of pharmacy adopted under the pharmacy act of the state of Kansas or for violation of the uniform controlled substances act or rules and regulations of the state board of pharmacy adopted under the uniform controlled substances act, in an amount not to exceed \$500 \$5,000 for each violation. All fines assessed and collected under this section shall be remitted to the state treasurer. Upon receipt thereof, the state treasurer shall deposit the entire amount in the state treasury and credit such amount to the state general fund.

Sec. 6. K.S.A. 65-1627f, 65-1632 and 65-1645 and K.S.A. 1997 Supp. 65-1627 and 65-1658 are hereby repealed.

(f) In each case in which a license of a pharmacist is issued or renewed for a period of time less than two years, the board shall prorate to the nearest whole month the license or renewal fee established pursuant to K.S.A. 65-1645 and amendments thereto.

Policy: Increase in maximum civil fine amount.

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Sec. 7. This act shall take effect and be in force from and after its publication in the statute book.

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SENATE BILL No. 533

By Committee on Public Health and Welfare

1-28

AN ACT concerning the sale of medicines and drugs through vending machines; amending K.S.A. 65-650 and repealing the existing section.

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

Section 1. K.S.A. 65-650 is hereby amended to read as follows: 65-650. Any person, firm or corporation who shall offer for sale or sell or distribute any prescription medicine, prescription-only drug or poison through or by means of any vending machine or other mechanical device, or who shall use any vending machine in or for the sale or distribution of any prescription medicine, prescription-only drug or poison, shall be deemed guilty of a class C nonperson misdemeanor and upon conviction shall be fined not less than twenty five dollars (\$25) \$25 nor more than five hundred dollars (\$500) \$500. No nonprescription drugs shall be sold through a vending machine in anything other than the manufacturer's original, tamper evident and expiration-dated packet.

Sec. 2. K.S.A. 65-650 is hereby repealed.

Sec. 3. This act shall take effect and be in force from and after its publication in the statute book.

(a)

vending Any machine in which nonprescription drugs are offered for sale or sold shall be located climate controlled area, and the drugs offered for sale or sold in such vending not be older machine shall than the manufacturer's expiration date. violation of this subsection (b) is class C nonperson misdemeanor and upon conviction the violator shall be fined not less than \$25 nor more than \$500.