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AN ACT concerning professional regulation.

2 Be it enacted by the People of the State of Illinois, 3 represented in the General Assembly:

4 Section 5. The Pharmacy Practice Act is amended by changing 5 Sections 3, 9, 9.5, 16a, 25.15, 30, and 35.16 as follows:

6 (225 ILCS 85/3) (from Ch. 111, par. 4123)

(Section scheduled to be repealed on January 1, 2018)

8 Sec. 3. Definitions. For the purpose of this Act, except 9 where otherwise limited therein:

"Pharmacy" or "drugstore" means and includes every 10 (a) shop, pharmacy department, or other place where 11 store, pharmacist care is provided by a pharmacist (1) where drugs, 12 medicines, or poisons are dispensed, sold or offered for sale 13 14 at retail, or displayed for sale at retail; or (2) where prescriptions of physicians, dentists, advanced practice 15 16 nurses, physician assistants, veterinarians, podiatrists, or 17 optometrists, within the limits of their licenses, are compounded, filled, or dispensed; or (3) which has upon it or 18 displayed within it, or affixed to or used in connection with 19 20 it, a sign bearing the word or words "Pharmacist", "Druggist", 21 "Pharmacy", "Pharmaceutical Care", "Apothecary", "Drugstore", "Medicine Store", "Prescriptions", "Drugs", "Dispensary", 22 "Medicines", or any word or words of similar or like import, 23

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either in the English language or any other language; or (4) where the characteristic prescription sign (Rx) or similar design is exhibited; or (5) any store, or shop, or other place with respect to which any of the above words, objects, signs or designs are used in any advertisement.

6 (b) "Drugs" means and includes (1) articles recognized in 7 the official United States Pharmacopoeia/National Formulary 8 (USP/NF), or any supplement thereto and being intended for and 9 having for their main use the diagnosis, cure, mitigation, 10 treatment or prevention of disease in man or other animals, as 11 approved by the United States Food and Drug Administration, but 12 does not include devices or their components, parts, or 13 accessories; and (2) all other articles intended for and having 14 for their main use the diagnosis, cure, mitigation, treatment 15 or prevention of disease in man or other animals, as approved 16 by the United States Food and Drug Administration, but does not 17 include devices or their components, parts, or accessories; and (3) articles (other than food) having for their main use and 18 19 intended to affect the structure or any function of the body of 20 man or other animals; and (4) articles having for their main 21 use and intended for use as a component or any articles 22 specified in clause (1), (2) or (3); but does not include 23 devices or their components, parts or accessories.

(c) "Medicines" means and includes all drugs intended for
 human or veterinary use approved by the United States Food and
 Drug Administration.

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(d) "Practice of pharmacy" means (1) the interpretation and 1 2 the provision of assistance in the monitoring, evaluation, and implementation of prescription drug orders; (2) the dispensing 3 of prescription drug orders; (3) participation in drug and 4 5 device selection; (4) drug administration limited to the administration of oral, topical, injectable, and inhalation as 6 7 follows: in the context of patient education on the proper use or delivery of medications; vaccination of patients 14 years of 8 9 age and older pursuant to a valid prescription or standing 10 order, by a physician licensed to practice medicine in all its 11 branches, upon completion of appropriate training, including 12 how to address contraindications and adverse reactions set forth by rule, with notification to the patient's physician and 13 14 appropriate record retention, or pursuant to hospital pharmacy 15 and therapeutics committee policies and procedures; (5) drug 16 regimen review; (6) drug or drug-related research; (7) the 17 provision of patient counseling; (8) the practice of telepharmacy; (9) the provision of those acts or services 18 19 necessary to provide pharmacist care; (10) medication therapy management; and (11) the responsibility for compounding and 20 21 labeling of drugs and devices (except labeling by а 22 manufacturer, repackager, or distributor of non-prescription 23 drugs and commercially packaged legend drugs and devices), proper and safe storage of drugs and devices, and maintenance 24 25 of required records. A pharmacist who performs any of the acts defined as the practice of pharmacy in this State must be 26

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1 actively licensed as a pharmacist under this Act.

2 (e) "Prescription" means and includes any written, oral, 3 facsimile, or electronically transmitted order for drugs or medical devices, issued by a physician licensed to practice 4 5 medicine in all its branches, dentist, veterinarian, or podiatrist, or optometrist, within the limits of their 6 7 licenses, by a physician assistant in accordance with 8 subsection (f) of Section 4, or by an advanced practice nurse 9 in accordance with subsection (q) of Section 4, containing the 10 following: (1) name of the patient; (2) date when prescription 11 was issued; (3) name and strength of drug or description of the 12 medical device prescribed; and (4) quantity, (5) directions for 13 use, (6) prescriber's name, address and signature, and (7) DEA number where required, for controlled substances. DEA numbers 14 15 shall not be required on inpatient drug orders.

16 (f) "Person" means and includes a natural person, 17 copartnership, association, corporation, government entity, or 18 any other legal entity.

(g) "Department" means the Department of Financial andProfessional Regulation.

(h) "Board of Pharmacy" or "Board" means the State Board of Pharmacy of the Department of Financial and Professional Regulation.

24 (i) "Secretary" means the Secretary of Financial and25 Professional Regulation.

26 (j) "Drug product selection" means the interchange for a

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1 prescribed pharmaceutical product in accordance with Section 2 25 of this Act and Section 3.14 of the Illinois Food, Drug and 3 Cosmetic Act.

(k) "Inpatient drug order" means an order issued by an 4 5 authorized prescriber for a resident or patient of a facility 6 licensed under the Nursing Home Care Act or the Hospital 7 Licensing Act, or "An Act in relation to the founding and operation of the University of Illinois Hospital and the 8 9 conduct of University of Illinois health care programs", 10 approved July 3, 1931, as amended, or a facility which is operated by the Department of Human Services (as successor to 11 12 Department of Mental Health the and Developmental 13 Disabilities) or the Department of Corrections.

14 (k-5) "Pharmacist" means an individual health care 15 professional and provider currently licensed by this State to 16 engage in the practice of pharmacy.

(1) "Pharmacist in charge" means the licensed pharmacist whose name appears on a pharmacy license and who is responsible for all aspects of the operation related to the practice of pharmacy.

21 (m) "Dispense" or "dispensing" means the interpretation, 22 evaluation, and implementation of a prescription drug order, 23 including the preparation and delivery of a drug or device to a 24 patient or patient's agent in a suitable container 25 appropriately labeled for subsequent administration to or use 26 by a patient in accordance with applicable State and federal HB1293 Enrolled - 6 - LRB096 09536 ASK 19695 b

laws and regulations. "Dispense" or "dispensing" does not mean 1 2 physical delivery to a patient or the а patient's representative in a home or institution by a designee of a 3 pharmacist or by common carrier. "Dispense" or "dispensing" 4 5 also does not mean the physical delivery of a drug or medical 6 device to a patient or patient's representative by a 7 pharmacist's designee within a pharmacy or drugstore while the 8 pharmacist is on duty and the pharmacy is open.

9 (n) "Nonresident pharmacy" means a pharmacy that is located 10 in a state, commonwealth, or territory of the United States, 11 other than Illinois, that delivers, dispenses, or distributes, 12 through the United States Postal Service, commercially 13 acceptable parcel delivery service, or other common carrier, to 14 Illinois residents, any substance which requires а 15 prescription.

16 (o) "Compounding" means the preparation and mixing of 17 components, excluding flavorings, (1) as the result of a prescriber's prescription drug order or initiative based on the 18 prescriber-patient-pharmacist relationship in the course of 19 20 professional practice or (2) for the purpose of, or incident to, research, teaching, or chemical analysis and not for sale 21 22 or dispensing. "Compounding" includes the preparation of drugs 23 or devices in anticipation of receiving prescription drug on routine, regularly observed dispensing 24 orders based patterns. Commercially available products may be compounded 25 for dispensing to individual patients only if all of the 26

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1 following conditions are met: (i) the commercial product is not 2 reasonably available from normal distribution channels in a 3 timely manner to meet the patient's needs and (ii) the 4 prescribing practitioner has requested that the drug be 5 compounded.

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(q) (Blank).

(p) (Blank).

(r) "Patient counseling" means the communication between a 8 9 pharmacist or a student pharmacist pharmacy intern under the 10 supervision of a pharmacist and a patient or the patient's 11 representative about the patient's medication or device for the 12 purpose of optimizing proper use of prescription medications or 13 devices. "Patient counseling" may include without limitation (1) obtaining a medication history; (2) acquiring a patient's 14 allergies and health conditions; (3) facilitation of the 15 patient's understanding of the intended use of the medication; 16 17 (4) proper directions for use; (5) significant potential adverse events; (6) potential food-drug interactions; and (7) 18 19 the need to be compliant with the medication therapy. A 20 pharmacy technician may only participate in the following aspects of patient counseling under the supervision of a 21 22 pharmacist: (1) obtaining medication history; (2) providing 23 the offer for counseling by a pharmacist or student pharmacist intern; and (3) acquiring a patient's allergies and health 24 conditions. 25

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(s) "Patient profiles" or "patient drug therapy record"

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means the obtaining, recording, and maintenance of patient prescription information, including prescriptions for controlled substances, and personal information.

4 (t) (Blank).

5 (11) "Medical device" means an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent, or 6 7 other similar or related article, including any component part 8 or accessory, required under federal law to bear the label 9 "Caution: Federal law requires dispensing by or on the order of 10 a physician". A seller of goods and services who, only for the 11 purpose of retail sales, compounds, sells, rents, or leases 12 medical devices shall not, by reasons thereof, be required to 13 be a licensed pharmacy.

14 (v) "Unique identifier" means an electronic signature, 15 handwritten signature or initials, thumb print, or other 16 acceptable biometric or electronic identification process as 17 approved by the Department.

18 (w) "Current usual and customary retail price" means the 19 price that a pharmacy charges to a non-third-party payor.

20 (x) "Automated pharmacy system" means a mechanical system
21 located within the confines of the pharmacy or remote location
22 that performs operations or activities, other than compounding
23 or administration, relative to storage, packaging, dispensing,
24 or distribution of medication, and which collects, controls,
25 and maintains all transaction information.

26 (y) "Drug regimen review" means and includes the evaluation

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of prescription drug orders and patient records for (1) known 1 2 allergies; (2) drug or potential therapy contraindications; duration 3 reasonable dose, of use, and route (3) of administration, taking into consideration factors such as age, 4 5 gender, and contraindications; (4) reasonable directions for 6 use; (5) potential or actual adverse drug reactions; (6) drug-drug interactions; (7) drug-food interactions; 7 (8) 8 drug-disease contraindications; (9) therapeutic duplication; 9 (10) patient laboratory values when authorized and available; 10 (11) proper utilization (including over or under utilization) 11 and optimum therapeutic outcomes; and (12) abuse and misuse.

12 "Electronic transmission prescription" (Z) means any 13 prescription order for which a facsimile or electronic image of the order is electronically transmitted from a licensed 14 15 prescriber to а pharmacy. "Electronic transmission 16 prescription" includes both data and image prescriptions.

17 "Medication therapy management services" means a (aa) distinct service or group of services offered by licensed 18 pharmacists, physicians licensed to practice medicine in all 19 20 its branches, advanced practice nurses authorized in a written 21 agreement with a physician licensed to practice medicine in all 22 its branches, or physician assistants authorized in guidelines 23 by a supervising physician that optimize therapeutic outcomes for individual patients through improved medication use. In a 24 25 retail or other non-hospital pharmacy, medication therapy shall consist of the evaluation of 26 management services

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prescription drug orders and patient medication records to 1 2 resolve conflicts with the following: 3 (1) known allergies; (2) drug or potential therapy contraindications; 4 5 (3) reasonable dose, duration of use, and route of administration, taking into consideration factors such as 6 7 age, gender, and contraindications; (4) reasonable directions for use; 8 9 (5) potential or actual adverse drug reactions; 10 (6) drug-drug interactions; 11 (7) drug-food interactions; 12 (8) drug-disease contraindications; 13 (9) identification of therapeutic duplication; 14 (10) patient laboratory values when authorized and 15 available; 16 (11) proper utilization (including over or under 17 utilization) and optimum therapeutic outcomes; and (12) drug abuse and misuse. 18 19 "Medication therapy management services" includes the 20 following: delivered 21 (1)documenting the services and 22 communicating the information provided to patients' 23 prescribers within an appropriate time frame, not to exceed 48 hours: 24 25 (2) providing patient counseling designed to enhance a 26 patient's understanding and the appropriate use of his or

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her medications; and

2 (3) providing information, support services, and 3 resources designed to enhance a patient's adherence with 4 his or her prescribed therapeutic regimens.

5 "Medication therapy management services" may also include 6 patient care functions authorized by a physician licensed to 7 practice medicine in all its branches for his or her identified 8 patient or groups of patients under specified conditions or 9 limitations in a standing order from the physician.

10 "Medication therapy management services" in a licensed 11 hospital may also include the following:

12 (1) reviewing assessments of the patient's health13 status; and

14 (2) following protocols of a hospital pharmacy and 15 therapeutics committee with respect to the fulfillment of 16 medication orders.

(bb) "Pharmacist care" means the provision by a pharmacist of medication therapy management services, with or without the dispensing of drugs or devices, intended to achieve outcomes that improve patient health, quality of life, and comfort and enhance patient safety.

(cc) "Protected health information" means individually identifiable health information that, except as otherwise provided, is:

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(1) transmitted by electronic media;

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(2) maintained in any medium set forth in the

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definition of "electronic media" in the federal Health
 Insurance Portability and Accountability Act; or

3 (3) transmitted or maintained in any other form or 4 medium.

5 "Protected health information" does not include individually6 identifiable health information found in:

7 (1) education records covered by the federal Family
8 Educational Right and Privacy Act; or

9 (2) employment records held by a licensee in its role 10 as an employer.

(dd) "Standing order" means a specific order for a patient or group of patients issued by a physician licensed to practice medicine in all its branches in Illinois.

14 (ee) "Address of record" means the address recorded by the 15 Department in the applicant's or licensee's application file or 16 license file, as maintained by the Department's licensure 17 maintenance unit.

18 (ff) "Home pharmacy" means the location of a pharmacy's 19 primary operations.

20 (Source: P.A. 94-459, eff. 1-1-06; 95-689, eff. 10-29-07.)

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(225 ILCS 85/9) (from Ch. 111, par. 4129)

22 (Section scheduled to be repealed on January 1, 2018)

23 Sec. 9. Registration as pharmacy technician. Any person 24 shall be entitled to registration as a registered pharmacy 25 technician who is of the age of 16 or over, has not engaged in

conduct or behavior determined to be grounds for discipline 1 2 under this Act, is attending or has graduated from an accredited high school or comparable school or educational 3 institution or received a GED, and has filed a written 4 5 application for registration on a form to be prescribed and 6 furnished by the Department for that purpose. The Department 7 shall issue a certificate of registration as a registered 8 pharmacy technician to any applicant who has qualified as 9 aforesaid, and such registration shall be the sole authority 10 required to assist licensed pharmacists in the practice of 11 pharmacy, under the supervision of a licensed pharmacist. A 12 registered pharmacy technician may, under the supervision of a 13 pharmacist, assist in the practice of pharmacy and perform such 14 functions as assisting in the dispensing process, offering 15 counseling, receiving new verbal prescription orders, and 16 having prescriber contact concerning prescription drug order 17 clarification. A registered pharmacy technician may not engage in patient counseling, drug regimen review, or clinical 18 conflict resolution. 19

Beginning on January 1, 2010, within 2 years after <u>initial</u> <u>registration</u> being employed as a registered technician, a pharmacy technician must become certified by successfully passing the Pharmacy Technician Certification Board (PTCB) examination or another Board-approved pharmacy technician examination <u>and register as a certified pharmacy technician</u> <u>with the Department</u> in order to continue to perform pharmacy

technician's duties. This requirement does not apply to 1 2 pharmacy technicians registered hired prior to January 1, 2008. 3 Any person registered as a pharmacy technician who is also 4 enrolled in a first professional degree program in pharmacy in 5 a school or college of pharmacy or a department of pharmacy of 6 a university approved by the Department or has graduated from 7 such a program within the last 18 months, shall be considered a 8 "student pharmacist pharmacy intern" and entitled to use the 9 "<u>student pharmacist"</u> pharmacy intern". title А student 10 pharmacist pharmacy intern must meet all of the requirements 11 for registration as a pharmacy technician set forth in this 12 Section excluding the requirement of certification prior to the 13 second registration renewal and pay the required pharmacy 14 technician registration fees. A student pharmacist may, under the supervision of a pharmacist, assist in the practice of 15 16 pharmacy and perform any and all functions delegated to him or 17 her by the pharmacist.

Any person seeking licensure as a pharmacist who has 18 19 graduated from a pharmacy program outside the United States 20 must register as a pharmacy technician and shall be considered a "student pharmacist" and be entitled to use the title 21 22 "student pharmacist" while completing the 1,200 clinical hours 23 of training approved by the Board of Pharmacy described and for 24 no more than 18 months after completion of these hours. These 25 individuals are not required to become certified pharmacy technicians while completing their Board approved clinical 26

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1 training, but must become licensed as a pharmacist or become a
2 certified pharmacy technician before the second pharmacy
3 technician registration renewal following completion of the
4 Board approved clinical training.

5 The Department shall not renew the pharmacy technician license of any person who has been registered as a "student 6 pharmacist" and has dropped out of or been expelled from an 7 ACPE accredited college of pharmacy, who has failed to complete 8 9 his or her 1,200 hours of Board approved clinical training 10 within 24 months or who has failed the pharmacist licensure 11 examination 3 times and shall require these individuals to meet 12 the requirements of and become registered a certified pharmacy 13 technician.

14 The Department, upon the recommendation of the Board, may 15 take any action set forth in Section 30 of this Act with regard 16 to <u>registrations</u> certificates pursuant to this Section.

17 Any person who is enrolled in a non-traditional Pharm.D. program at an ACPE accredited college of pharmacy and is a 18 licensed pharmacist under the laws of another United States 19 20 jurisdiction shall be permitted to engage in the program of practice experience required in the academic program by virtue 21 22 of such license. Such person shall be exempt from the 23 of registration as registered requirement а pharmacv technician while engaged in the program of practice experience 24 25 required in the academic program.

26 An applicant for registration as a pharmacy technician may

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assist a pharmacist in the practice of pharmacy for a period of 1 2 up to 60 days prior to the issuance of a certificate of 3 registration if the applicant has submitted the required fee and an application for registration to the Department. The 4 5 applicant shall keep a copy of the submitted application on the premises where the applicant is assisting in the practice of 6 7 pharmacy. The Department shall forward confirmation of receipt 8 of the application with start and expiration dates of practice 9 pending registration. 10 (Source: P.A. 95-689, eff. 10-29-07.) 11 (225 ILCS 85/9.5) 12 (Section scheduled to be repealed on January 1, 2018) 13 Sec. 9.5. Certified pharmacy technician. 14 (a) An individual registered as a pharmacy technician under 15 this Act may be registered receive certification as a certified 16 pharmacy technician, if he or she meets all of the following 17 requirements: (1) He or she has submitted a written application in 18 19 the form and manner prescribed by the Department Board. (2) He or she has attained the age of 18. 20 21 (3) He or she is of good moral character, as determined 22 by the Department. 23 (4) He or she has (i) graduated from pharmacy

24 technician training meeting the requirements set forth in 25 subsection (a) of Section 17.1 of this Act or (ii) obtained HB1293 Enrolled - 17 - LRB096 09536 ASK 19695 b

1 documentation from the pharmacist-in-charge of the 2 pharmacy where the applicant is employed verifying that he or she has successfully completed a training program and 3 successfully completed an objective assessment 4 has 5 mechanism prepared in accordance with rules established by 6 the Department Board.

7 (5) He or she has successfully passed an examination
8 accredited by the National Organization of Certifying
9 Agencies, as approved and required by the Board.

10 (6) He or she has paid the required certification fees.
11 (b) No pharmacist whose license has been denied, revoked,
12 suspended, or restricted for disciplinary purposes may be
13 eligible to be registered as a certified pharmacy technician.

14 (c) The <u>Department</u> Board may, by rule, establish any
 15 additional requirements for certification under this Section.

16 <u>(d) A person who is not a registered pharmacy technician</u> 17 <u>and meets the requirements of this Section may register as a</u> 18 <u>certified pharmacy technician without first registering as a</u> 19 pharmacy technician.

20 (Source: P.A. 95-689, eff. 10-29-07.)

(225 ILCS 85/16a) (from Ch. 111, par. 4136a)
(Section scheduled to be repealed on January 1, 2018)
Sec. 16a. (a) The Department shall establish rules and
regulations, consistent with the provisions of this Act,
governing nonresident pharmacies, including pharmacies

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providing services via the Internet, which sell, or offer for sale, drugs, medicines, or other pharmaceutical services in this State.

(b) The Department Board shall require and provide for an 4 5 annual nonresident special pharmacy registration for all located outside of this State that 6 pharmacies dispense 7 medications for Illinois residents and mail, ship, or deliver 8 prescription medications into this State. Nonresident special 9 pharmacy registration shall be granted by the Department Board 10 upon the disclosure and certification by a pharmacy:

(1) that it is licensed in the state in which the dispensing facility is located and from which the drugs are dispensed;

14 (2) of the location, names, and titles of all principal
 15 corporate officers and all pharmacists who are dispensing
 16 drugs to residents of this State;

(3) that it complies with all lawful directions and requests for information from the board of pharmacy of each state in which it is licensed or registered, except that it shall respond directly to all communications from the Board <u>or Department</u> concerning <u>any</u> <u>emergency</u> circumstances arising from the dispensing of drugs to residents of this State;

(4) that it maintains its records of drugs dispensed to
residents of this State so that the records are readily
retrievable from the records of other drugs dispensed;

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1 (5) that it cooperates with the Board <u>or Department</u> in 2 providing information to the board of pharmacy of the state 3 in which it is licensed concerning matters related to the 4 dispensing of drugs to residents of this State; and

5 (6) that during its regular hours of operation, but not less than 6 days per week, for a minimum of 40 hours per 6 7 a toll-free telephone service is provided to week, 8 facilitate communication between patients in this State 9 and a pharmacist at the pharmacy who has access to the patients' records. The toll-free number must be disclosed 10 11 on the label affixed to each container of drugs dispensed 12 to residents of this State.

13 (Source: P.A. 95-689, eff. 10-29-07.)

14 (225 ILCS 85/25.15)

15 (Section scheduled to be repealed on January 1, 2018)

16 Sec. 25.15. Telepharmacy.

(a) In this Section, "telepharmacy" means the provision of pharmacist care by a pharmacist that is accomplished through the use of telecommunications or other technologies to patients or their agents who are at a distance and are located within the United States, and which follows all federal and State laws, rules, and regulations with regard to privacy and security.

(b) Any pharmacy engaged in the practice of telepharmacymust meet all of the following conditions:

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(1) All events involving the contents of an automated
 pharmacy system must be stored in a secure location and may
 be recorded electronically.

4 (2) An automated pharmacy or prescription dispensing 5 machine system may be used in conjunction with the 6 pharmacy's practice of telepharmacy after inspection and 7 approval by the Department.

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(3) The pharmacist in charge shall:

9 (A) be responsible for the practice of 10 telepharmacy performed at a remote pharmacy, including 11 the supervision of any prescription dispensing machine 12 or automated medication system;

(B) ensure that the home pharmacy has sufficient
pharmacists on duty for the safe operation and
supervision of all remote pharmacies;

16 (C) ensure, through the use of a video and auditory 17 communication system, that a certified pharmacy 18 technician at the remote pharmacy has accurately and 19 correctly prepared any prescription for dispensing 20 according to the prescription;

21 (D) be responsible for the supervision and 22 training of certified pharmacy technicians at remote 23 pharmacies who shall be subject to all rules and 24 regulations; and

25 (E) ensure that patient counseling at the remote 26 pharmacy is performed by a pharmacist or <u>student</u> HB1293 Enrolled - 21 - LRB096 09536 ASK 19695 b

1	pharmacist pharmacist intern.
2	(Source: P.A. 95-689, eff. 10-29-07.)
3	(225 ILCS 85/30) (from Ch. 111, par. 4150)
4	(Section scheduled to be repealed on January 1, 2018)
5	Sec. 30. <u>Refusal, revocation, or suspension.</u>
6	(a) The Department may refuse to issue or renew, or may
7	revoke a license or registration, or may suspend, place on
8	probation, fine, or take any disciplinary or non-disciplinary
9	action as the Department may deem proper, including fines not
10	to exceed \$10,000 for each violation, with regard to any
11	<u>licensee or registrant</u> In accordance with Section 11 of this
12	Act, the Department may refuse to issue, restore, or renew, or
13	may revoke, suspend, place on probation, or reprimand as the
14	Department may deem proper with regard to any license or
15	certificate of registration or may impose a fine upon a
16	licensee or registrant not to exceed \$10,000 per violation for
17	any one or combination of the following causes:
18	1. Material misstatement in furnishing information to
19	the Department.
20	2. Violations of this Act, or the rules promulgated
21	hereunder.
22	3. Making any misrepresentation for the purpose of
23	obtaining licenses.
24	4. A pattern of conduct which demonstrates
25	incompetence or unfitness to practice.

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5. Aiding or assisting another person in violating any
 provision of this Act or rules.

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6. Failing, within 60 days, to respond to a written request made by the Department for information.

7. Engaging in <u>unprofessional</u>, dishonorable, or unethical conduct of a character likely to deceive, defraud or harm the public.

8 8. Discipline by another U.S. jurisdiction or foreign 9 nation, if at least one of the grounds for the discipline 10 is the same or substantially equivalent to those set forth 11 herein.

9. Directly or indirectly giving to or receiving from any person, firm, corporation, partnership or association any fee, commission, rebate or other form of compensation for any professional services not actually or personally rendered.

10. A finding by the Department that the licensee,
after having his license placed on probationary status has
violated the terms of probation.

20 11. Selling or engaging in the sale of drug samples21 provided at no cost by drug manufacturers.

12. Physical illness, including but not limited to, deterioration through the aging process, or loss of motor skill which results in the inability to practice the profession with reasonable judgment, skill or safety.

13. A finding that licensure or registration has been

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applied for or obtained by fraudulent means.

14. The applicant or licensee has been convicted in state or federal court of or entered a plea of guilty, nolo contendere, or the equivalent in a state or federal court to any crime which is a felony or any misdemeanor related to the practice of pharmacy <u>or</u>, of which an essential element is dishonesty.

8 15. Habitual or excessive use or addiction to alcohol, 9 narcotics, stimulants or any other chemical agent or drug 10 which results in the inability to practice with reasonable 11 judgment, skill or safety.

12 16. Willfully making or filing false records or reports 13 in the practice of pharmacy, including, but not limited to 14 false records to support claims against the medical 15 assistance program of the Department of Healthcare and 16 Family Services (formerly Department of Public Aid) under 17 the Public Aid Code.

17. Gross and willful overcharging for professional 18 19 services including filing false statements for collection 20 of fees for which services are not rendered, including, but 21 not limited to, filing false statements for collection of 22 monies for services not rendered from the medical 23 assistance program of the Department of Healthcare and 24 Family Services (formerly Department of Public Aid) under 25 the Public Aid Code.

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18. <u>Dispensing</u> Repetitiously dispensing prescription

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drugs without receiving a written or oral prescription <u>in</u>
 violation of law.

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19. Upon a finding of a substantial discrepancy in a Department audit of a prescription drug, including controlled substances, as that term is defined in this Act or in the Illinois Controlled Substances Act.

7 20. Physical or mental illness or any other impairment 8 or disability, including without limitation deterioration 9 through the aging process or loss of motor skills that 10 results in the inability to practice with reasonable 11 judgment, skill or safety, or mental incompetence, as 12 declared by a court of competent jurisdiction.

13 21. Violation of the Health Care Worker Self-Referral14 Act.

15 22. Failing to sell or dispense any drug, medicine, or
16 poison in good faith. "Good faith", for the purposes of
17 this Section, has the meaning ascribed to it in subsection
18 (u) of Section 102 of the Illinois Controlled Substances
19 Act. "Good faith", as used in this item (22), shall not be
20 limited to the sale or dispensing of controlled substances,
21 but shall apply to all prescription drugs.

22 23. Interfering with the professional judgment of a
23 pharmacist by any registrant under this Act, or his or her
24 agents or employees.

25 24. Failing to report within 60 days to the Department26 any adverse final action taken against a pharmacist,

pharmacist technician, or certified pharmacist technician by another licensing jurisdiction in any other state or any territory of the United States or any foreign jurisdiction, any governmental agency, any law enforcement agency, or any court for acts or conduct similar to acts or conduct that would constitute grounds for discipline as defined in this Section.

8 25. Failing to comply with a subpoena issued in 9 accordance with Section 35.5 of this Act.

10 <u>26. Disclosing protected health information in</u>
 11 violation of any State or federal law.

12 (b) The Department may refuse to issue or may suspend the license or registration of any person who fails to file a 13 14 return, or to pay the tax, penalty or interest shown in a filed 15 return, or to pay any final assessment of tax, penalty or 16 interest, as required by any tax Act administered by the 17 Illinois Department of Revenue, until such time as the requirements of any such tax Act are satisfied. 18

19 (c) The Department shall revoke the license or certificate 20 of registration issued under the provisions of this Act or any prior Act of this State of any person who has been convicted a 21 22 second time of committing any felony under the Illinois 23 Controlled Substances Act, or who has been convicted a second time of committing a Class 1 felony under Sections 8A-3 and 24 25 8A-6 of the Illinois Public Aid Code. A person whose license or 26 certificate of registration issued under the provisions of this

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Act or any prior Act of this State is revoked under this
 subsection (c) shall be prohibited from engaging in the
 practice of pharmacy in this State.

(d) The Department may adopt rules for the imposition of 4 5 fines in disciplinary cases, not to exceed \$10,000 for each 6 violation of this Act. Fines may be imposed in conjunction with 7 other forms of disciplinary action, but shall not be the 8 exclusive disposition of any disciplinary action arising out of 9 conduct resulting in death or injury to a patient. Fines shall 10 be paid within 60 days or as otherwise agreed to by the 11 Department. Any funds collected from such fines shall be 12 deposited in the Illinois State Pharmacy Disciplinary Fund.

13 (e) The entry of an order or judgment by any circuit court 14 establishing that any person holding a license or certificate 15 under this Act is a person in need of mental treatment operates 16 as a suspension of that license. A licensee may resume his or 17 her practice only upon the entry of an order of the Department based upon a finding by the Board that he or she has been 18 determined to be recovered from mental illness by the court and 19 20 upon the Board's recommendation that the licensee be permitted to resume his or her practice. 21

(f) The Department shall issue quarterly to the Board a status of all complaints related to the profession received by the Department.

(g) In enforcing this Section, the Board or the Department,upon a showing of a possible violation, may compel any licensee

or applicant for licensure under this Act to submit to a mental 1 2 or physical examination or both, as required by and at the 3 expense of the Department. The examining physician, or multidisciplinary team involved in providing physical and 4 mental examinations led by a physician consisting of one or a 5 combination of licensed physicians, licensed clinical 6 7 psychologists, licensed clinical social workers, licensed 8 clinical professional counselors, and other professional and 9 administrative staff, shall be those specifically designated 10 by the Department. The Board or the Department may order the 11 examining physician or any member of the multidisciplinary team 12 to present testimony concerning this mental or physical examination of the licensee or applicant. No information, 13 14 report, or other documents in any way related to the 15 examination shall be excluded by reason of any common law or 16 statutory privilege relating to communication between the 17 licensee or applicant and the examining physician or any member of the multidisciplinary team. The individual to be examined 18 19 may have, at his or her own expense, another physician of his 20 or her choice present during all aspects of the examination. Failure of any individual to submit to a mental or physical 21 22 examination when directed shall be grounds for suspension of 23 his or her license until such time as the individual submits to the examination if the Board finds, after notice and hearing, 24 25 that the refusal to submit to the examination was without 26 reasonable cause. If the Board finds a pharmacist, certified HB1293 Enrolled - 28 - LRB096 09536 ASK 19695 b

1 pharmacy technician, or pharmacy technician unable to practice 2 because of the reasons set forth in this Section, the Board shall require such pharmacist, certified pharmacy technician, 3 or pharmacy technician to submit to care, counseling, or 4 5 treatment by physicians or other appropriate health care 6 providers approved or designated by the Board as a condition 7 for continued, reinstated, or renewed licensure to practice. Any pharmacist, certified pharmacy technician, or pharmacy 8 9 technician whose license was granted, continued, reinstated, 10 renewed, disciplined, or supervised, subject to such terms, 11 conditions, or restrictions, and who fails to comply with such 12 terms, conditions, or restrictions or to complete a required program of care, counseling, or treatment, as determined by the 13 14 chief pharmacy coordinator or a deputy pharmacy coordinator, 15 shall be referred to the Secretary for a determination as to 16 whether the licensee shall have his or her license suspended 17 immediately, pending a hearing by the Board. In instances in which the Secretary immediately suspends a license under this 18 19 subsection (g), a hearing upon such person's license must be 20 convened by the Board within 15 days after such suspension and completed without appreciable delay. The Board shall have the 21 22 authority to review the subject pharmacist's, certified 23 pharmacy technician's, or pharmacy technician's record of 24 treatment and counseling regarding the impairment. 25 (Source: P.A. 95-331, eff. 8-21-07; 95-689, eff. 10-29-07.)

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(225 ILCS 85/35.16) (from Ch. 111, par. 4155.16) 1 2 (Section scheduled to be repealed on January 1, 2018) 3 Sec. 35.16. The Secretary Director may temporarily suspend the license of a pharmacist or pharmacy, or the registration of 4 <u>a</u>, pharmacy technician or <u>certified</u> pharmacy technician 5 registration as a distributor, without 6 а hearing, 7 simultaneously with the institution of proceedings for a hearing provided for in Section 35.2 of this Act, if the 8 9 Secretary **Director** finds that evidence in his possession 10 indicates that a continuation in practice would constitute an 11 imminent danger to the public. In the event that the Secretary 12 Director suspends, temporarily, this license or registration certificate without a hearing, a hearing by the Department must 13 be held within 15 days after such suspension has occurred, and 14 15 be concluded without appreciable delay.

16 (Source: P.A. 95-689, eff. 10-29-07.)