HB1179

LRB094 06611 RAS 36703 b

94TH GENERAL ASSEMBLY

State of Illinois

2005 and 2006

HB1179

Introduced 02/08/05, by Rep. Angelo Saviano - Frank J. Mautino - Arthur L. Turner - William B. Black

SYNOPSIS AS INTRODUCED:

225 ILCS 80/3	from Ch. 111, par. 3903
225 ILCS 80/15.1	
720 ILCS 570/102	from Ch. 56 1/2, par. 1102
720 ILCS 570/103	from Ch. 56 1/2, par. 1103

Amends the Illinois Optometric Practice Act of 1987. Makes a technical change in a Section concerning the practice of optometry. Changes the definition of "therapeutic ocular pharmaceutical agents". Provides that medication administered by injection may be used only for the treatment of anaphylaxis. Amends the Illinois Controlled Substances Act. Adds references to optometrists in the definitions of "practitioner", "prescriber", and "prescription". Provides that nothing in the Illinois Controlled Substances Act limits the lawful authority granted by the Optometric Practice Act of 1987.

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A BILL FOR

AN ACT concerning regulation.

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

Section 5. The Illinois Optometric Practice Act of 1987 is amended by changing Sections 3 and 15.1 as follows:

(225 ILCS 80/3) (from Ch. 111, par. 3903)

(Section scheduled to be repealed on January 1, 2007)

Sec. 3. Practice of optometry defined; referrals; manufacture of lenses and prisms.

(a) The practice of optometry is defined as the employment of any and all means for the examination, diagnosis, and treatment of the human visual system, the human eye, and its appendages without the use of surgery, including but not limited to: the appropriate use of diagnostic ocular pharmaceutical agents and therapeutic ocular pharmaceutical agents; refraction and other determinants of visual function; prescribing corrective lenses or prisms; prescribing, dispensing, or management of contact lenses; vision therapy; visual rehabilitation; or any other procedures taught in schools and colleges of optometry approved by the Department, and not specifically restricted in this Act, subject to demonstrated competency and training as required by the Board, and pursuant to rule or regulation approved by the Board and adopted by the Department.

A person shall be deemed to be practicing optometry within the meaning of this Act who:

(1) In any way presents himself or herself to be qualified to practice optometry.

(2) Performs refractions or employs any other determinants of visual function.

(3) Employs any means for the adaptation of lenses or prisms.

(4) Prescribes corrective lenses, prisms, vision therapy, visual rehabilitation, or ocular pharmaceutical agents.

(5) Prescribes or manages contact lenses for refractive, cosmetic, or therapeutic purposes.

(6) Evaluates the need for, or prescribes, low vision aids to partially sighted persons.

(7) Diagnoses or treats any ocular abnormality, disease, or visual or muscular anomaly of the human eye or visual system.

(8) Practices, or offers or attempts to practice, optometry as defined in this Act either on his or her own behalf or as an employee of a person, firm, or corporation, whether under the supervision of his or her employer or not.

Nothing in this Section shall be interpreted (i) to prevent a person from functioning as an assistant under the direct supervision of a person licensed by the State of Illinois to practice optometry or medicine in all of its branches or (ii) to prohibit visual screening programs that are conducted without a fee (other than voluntary donations), by charitable organizations acting in the public welfare under the supervision of a committee composed of persons licensed by the State of Illinois to practice optometry or persons licensed by the State of Illinois to practice medicine in all of its branches.

(b) When, in the course of providing optometric services to any person, an optometrist licensed under this Act finds an indication of a disease or condition of the eye which in his or her professional judgment requires professional service outside the scope of practice as defined in this Act, he or she shall refer such person to a physician licensed to practice medicine in all of its branches, or other appropriate health care practitioner. Nothing in this Act shall preclude an optometrist who is therapeutically certified from rendering appropriate non-surgical ~~nonsurgical ophthalmic~~ emergency

care.

(c) Nothing contained in this Section shall prohibit a person from manufacturing ophthalmic lenses and prisms or the fabrication of contact lenses according to the specifications prescribed by an optometrist or a physician licensed to practice medicine in all of its branches, but shall specifically prohibit the sale or delivery of ophthalmic lenses, prisms, and contact lenses without a prescription signed by an optometrist or a physician licensed to practice medicine in all of its branches.

(d) Nothing in this Act shall restrict the filling of a prescription by a pharmacist licensed under the Pharmacy Practice Act of 1987.

(Source: P.A. 90-655, eff. 7-30-99; 91-141, eff. 7-16-99.)

(225 ILCS 80/15.1)

(Section scheduled to be repealed on January 1, 2007)

Sec. 15.1. Diagnostic and therapeutic certification.

(a) Any licensed optometrist may apply to the Department, in the form the Department may prescribe, for a certificate to use diagnostic topical ocular pharmaceutical agents and the Department shall certify the applicant if:

(1) the applicant has received appropriate training and certification from a properly accredited institution of higher learning for the certificate; and

(2) the applicant has demonstrated training and competence to use diagnostic topical ocular pharmaceutical agents as required by the Board pursuant to rule or regulation approved by the Board and adopted by the Department.

A certificate to use topical ocular pharmaceutical agents for diagnostic purposes previously issued by the Department that is current and valid on the effective date of this amendatory Act of 1995 is valid until its expiration date and entitles the holder of the certificate to use diagnostic topical ocular pharmaceutical agents as provided in this Act.

(b) Any licensed optometrist may apply to the Department, in the form the Department may prescribe, for a certificate to use therapeutic ocular pharmaceutical agents and the Department shall certify the applicant if:

(1) the applicant has received a certificate to use diagnostic topical ocular pharmaceutical agents under subsection (a);

(2) the applicant has received appropriate training and certification from a properly accredited institution of higher learning for the certificate; and

(3) the applicant has demonstrated training and competence to use therapeutic ocular pharmaceutical agents as required by the Board pursuant to rule or regulation approved by the Board and adopted by the Department.

All applicants for license renewal after January 1, 2006 must apply for and maintain certification to use therapeutic ocular pharmaceutical agents.

(c) For purposes of the Act, "diagnostic topical ocular pharmaceutical agents" means anesthetics, mydriatics, cycloplegics, and miotics used for diagnostic purposes as defined by the Board pursuant to rule approved by the Board and adopted by the Department.

(d) For the purposes of the Act, "therapeutic ocular pharmaceutical agents" means <u>all drugs and pharmaceutical agents appropriate for the diagnosis, treatment, and management of diseases and conditions of the eye and adnexa. Medication administered by injection may be used only for the treatment of anaphylaxis.</u> ~~the following when used for diagnostic or therapeutic purposes: topical anti infective agents, topical anti allergy agents, topical anti glaucoma agents, topical anti-inflammatory agents, topical anesthetic agents, over the counter agents, non-narcotic oral analgesic agents, and mydriatic reversing agents.~~

(e) A licensed optometrist who is therapeutically certified may remove superficial foreign bodies from the human eye and adnexa.

(e-5) A licensed optometrist who is therapeutically certified may give orders for patient care related to the use of therapeutic ocular pharmaceutical agents to a nurse licensed to practice under Illinois law.

(f) An optometrist's certificate to use diagnostic topical ocular pharmaceutical agents shall be revoked or suspended by the Department upon recommendation of the Board based on the misuse of any diagnostic topical ocular pharmaceutical agent.

(g) An optometrist's certificate to use therapeutic ocular pharmaceutical agents shall be revoked or suspended by the Department upon recommendation of the Board based on the misuse of any therapeutic ocular pharmaceutical agent.

(h) An optometrist's license shall be revoked or suspended by the Department upon recommendation of the Board based upon either of the following causes:

(1) grave or repeated misuse of any diagnostic or therapeutic ocular pharmaceutical agent; and

(2) the use of any agent or procedure in the course of optometric practice by an optometrist not properly certified under this Section.

(i) The provisions of Sections 26.2, 26.3, 26.5, 26.10, 26.11, 26.14, and 26.15 of this Act shall apply to all disciplinary proceedings brought under this Section.

(j) The Director may temporarily suspend a certificate to use diagnostic topical ocular pharmaceuticals or a certificate to use therapeutic ocular pharmaceuticals or a license to practice optometry, without a hearing, simultaneously with the institution of proceedings for a hearing based upon a violation of subsection (f), (g), or (h) of this Section, if the Director finds that evidence in his or her possession indicates that the continued use of diagnostic topical ocular pharmaceuticals, or therapeutic ocular pharmaceuticals, or continued practice of optometry would constitute an immediate danger to the public. In the event that the Director temporarily suspends a certificate to use diagnostic topical ocular pharmaceuticals, therapeutic ocular pharmaceuticals, or a license to practice

optometry without a hearing, a hearing by the Board shall be commenced within 15 days after suspension has occurred, and concluded without appreciable delay.

(k) The Director of the Department of Professional Regulation shall notify the Director of the Department of Public Health as to the categories of ocular pharmaceutical agents permitted for use by an optometrist. The Director of the Department of Public Health shall in turn notify every licensed pharmacist in the State of the categories of ocular pharmaceutical agents that can be utilized and prescribed by an optometrist.

(l) Nothing in this Act prohibits the use of diagnostic topical ocular pharmaceutical agents or therapeutic ocular pharmaceutical agents in the practice of optometry by optometrists certified for such use under this Section.

(Source: P.A. 90-73, eff. 7-8-97; 91-141, eff. 7-16-99.)

Section 10. The Illinois Controlled Substances Act is amended by changing Sections 102 and 103 as follows:

(720 ILCS 570/102) (from Ch. 56 1/2, par. 1102)

Sec. 102. Definitions. As used in this Act, unless the context otherwise requires:

(a) "Addict" means any person who habitually uses any drug, chemical, substance or dangerous drug other than alcohol so as to endanger the public morals, health, safety or welfare or who is so far addicted to the use of a dangerous drug or controlled substance other than alcohol as to have lost the power of self control with reference to his addiction.

(b) "Administer" means the direct application of a controlled substance, whether by injection, inhalation, ingestion, or any other means, to the body of a patient, research subject, or animal (as defined by the Humane Euthanasia in Animal Shelters Act) by:

(1) a practitioner (or, in his presence, by his authorized agent),

(2) the patient or research subject at the lawful direction of the practitioner, or

(3) a euthanasia technician as defined by the Humane Euthanasia in Animal Shelters Act.

(c) "Agent" means an authorized person who acts on behalf of or at the direction of a manufacturer, distributor, or dispenser. It does not include a common or contract carrier, public warehouseman or employee of the carrier or warehouseman.

(c-1) "Anabolic Steroids" means any drug or hormonal substance, chemically and pharmacologically related to testosterone (other than estrogens, progestins, and corticosteroids) that promotes muscle growth, and includes:

(i) boldenone,

(ii) chlorotestosterone,

(iii) chostebol,

(iv) dehydrochlormethyltestosterone,

(v) dihydrotestosterone,

(vi) drostanolone,

(vii) ethylestrenol,

(viii) fluoxymesterone,

(ix) formebulone,

(x) mesterolone,

(xi) methandienone,

(xii) methandranone,

(xiii) methandriol,

(xiv) methandrostenolone,

(xv) methenolone,

(xvi) methyltestosterone,

(xvii) mibolerone,

(xviii) nandrolone,

(xix) norethandrolone,

(xx) oxandrolone,

(xxi) oxymesterone,

(xxii) oxymetholone,

(xxiii) stanolone,

(xxiv) stanozolol,

(xxv) testolactone,

(xxvi) testosterone,

(xxvii) trenbolone, and

(xxviii) any salt, ester, or isomer of a drug or substance described or listed in this paragraph, if that salt, ester, or isomer promotes muscle growth.

Any person who is otherwise lawfully in possession of an anabolic steroid, or who otherwise lawfully manufactures, distributes, dispenses, delivers, or possesses with intent to deliver an anabolic steroid, which anabolic steroid is expressly intended for and lawfully allowed to be administered through implants to livestock or other nonhuman species, and which is approved by the Secretary of Health and Human Services for such administration, and which the person intends to administer or have administered through such implants, shall not be considered to be in unauthorized possession or to unlawfully manufacture, distribute, dispense, deliver, or possess with intent to deliver such anabolic steroid for purposes of this Act.

(d) "Administration" means the Drug Enforcement Administration, United States Department of Justice, or its successor agency.

(e) "Control" means to add a drug or other substance, or immediate precursor, to a Schedule under Article II of this Act whether by transfer from another Schedule or otherwise.

(f) "Controlled Substance" means a drug, substance, or immediate precursor in the Schedules of Article II of this Act.

(g) "Counterfeit substance" means a controlled substance, which, or the container or labeling of which, without authorization bears the trademark, trade name, or other identifying mark, imprint, number or device, or any likeness thereof, of a manufacturer, distributor, or dispenser other than the person who in fact manufactured, distributed, or dispensed the substance.

(h) "Deliver" or "delivery" means the actual, constructive or attempted transfer of possession of a controlled substance,

with or without consideration, whether or not there is an agency relationship.

(i) "Department" means the Illinois Department of Human Services (as successor to the Department of Alcoholism and Substance Abuse) or its successor agency.

(j) "Department of State Police" means the Department of State Police of the State of Illinois or its successor agency.

(k) "Department of Corrections" means the Department of Corrections of the State of Illinois or its successor agency.

(l) "Department of Professional Regulation" means the Department of Professional Regulation of the State of Illinois or its successor agency.

(m) "Depressant" or "stimulant substance" means:

(1) a drug which contains any quantity of (i) barbituric acid or any of the salts of barbituric acid which has been designated as habit forming under section 502 (d) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 352 (d)); or

(2) a drug which contains any quantity of (i) amphetamine or methamphetamine and any of their optical isomers; (ii) any salt of amphetamine or methamphetamine or any salt of an optical isomer of amphetamine; or (iii) any substance which the Department, after investigation, has found to be, and by rule designated as, habit forming because of its depressant or stimulant effect on the central nervous system; or

(3) lysergic acid diethylamide; or

(4) any drug which contains any quantity of a substance which the Department, after investigation, has found to have, and by rule designated as having, a potential for abuse because of its depressant or stimulant effect on the central nervous system or its hallucinogenic effect.

(n) (Blank).

(o) "Director" means the Director of the Department of State Police or the Department of Professional Regulation or his designated agents.

(p) "Dispense" means to deliver a controlled substance to an ultimate user or research subject by or pursuant to the lawful order of a prescriber, including the prescribing, administering, packaging, labeling, or compounding necessary to prepare the substance for that delivery.

(q) "Dispenser" means a practitioner who dispenses.

(r) "Distribute" means to deliver, other than by administering or dispensing, a controlled substance.

(s) "Distributor" means a person who distributes.

(t) "Drug" means (1) substances recognized as drugs in the official United States Pharmacopoeia, Official Homeopathic Pharmacopoeia of the United States, or official National Formulary, or any supplement to any of them; (2) substances intended for use in diagnosis, cure, mitigation, treatment, or prevention of disease in man or animals; (3) substances (other than food) intended to affect the structure of any function of the body of man or animals and (4) substances intended for use as a component of any article specified in clause (1), (2), or (3) of this subsection. It does not include devices or their components, parts, or accessories.

(t-5) "Euthanasia agency" means an entity certified by the Department of Professional Regulation for the purpose of animal euthanasia that holds an animal control facility license or animal shelter license under the Animal Welfare Act. A euthanasia agency is authorized to purchase, store, possess, and utilize Schedule II nonnarcotic and Schedule III nonnarcotic drugs for the sole purpose of animal euthanasia.

(t-10) "Euthanasia drugs" means Schedule II or Schedule III substances (nonnarcotic controlled substances) that are used by a euthanasia agency for the purpose of animal euthanasia.

(u) "Good faith" means the prescribing or dispensing of a controlled substance by a practitioner in the regular course of professional treatment to or for any person who is under his treatment for a pathology or condition other than that individual's physical or psychological dependence upon or addiction to a controlled substance, except as provided herein:

and application of the term to a pharmacist shall mean the dispensing of a controlled substance pursuant to the prescriber's order which in the professional judgment of the pharmacist is lawful. The pharmacist shall be guided by accepted professional standards including, but not limited to the following, in making the judgment:

(1) lack of consistency of doctor-patient relationship,

(2) frequency of prescriptions for same drug by one prescriber for large numbers of patients,

(3) quantities beyond those normally prescribed,

(4) unusual dosages,

(5) unusual geographic distances between patient, pharmacist and prescriber,

(6) consistent prescribing of habit-forming drugs.

(u-1) "Home infusion services" means services provided by a pharmacy in compounding solutions for direct administration to a patient in a private residence, long-term care facility, or hospice setting by means of parenteral, intravenous, intramuscular, subcutaneous, or intraspinal infusion.

(v) "Immediate precursor" means a substance:

(1) which the Department has found to be and by rule designated as being a principal compound used, or produced primarily for use, in the manufacture of a controlled substance;

(2) which is an immediate chemical intermediary used or likely to be used in the manufacture of such controlled substance; and

(3) the control of which is necessary to prevent, curtail or limit the manufacture of such controlled substance.

(w) "Instructional activities" means the acts of teaching, educating or instructing by practitioners using controlled substances within educational facilities approved by the State Board of Education or its successor agency.

(x) "Local authorities" means a duly organized State,

County or Municipal peace unit or police force.

(y) "Look-alike substance" means a substance, other than a controlled substance which (1) by overall dosage unit appearance, including shape, color, size, markings or lack thereof, taste, consistency, or any other identifying physical characteristic of the substance, would lead a reasonable person to believe that the substance is a controlled substance, or (2) is expressly or impliedly represented to be a controlled substance or is distributed under circumstances which would lead a reasonable person to believe that the substance is a controlled substance. For the purpose of determining whether the representations made or the circumstances of the distribution would lead a reasonable person to believe the substance to be a controlled substance under this clause (2) of subsection (y), the court or other authority may consider the following factors in addition to any other factor that may be relevant:

(a) statements made by the owner or person in control of the substance concerning its nature, use or effect;

(b) statements made to the buyer or recipient that the substance may be resold for profit;

(c) whether the substance is packaged in a manner normally used for the illegal distribution of controlled substances;

(d) whether the distribution or attempted distribution included an exchange of or demand for money or other property as consideration, and whether the amount of the consideration was substantially greater than the reasonable retail market value of the substance.

Clause (1) of this subsection (y) shall not apply to a noncontrolled substance in its finished dosage form that was initially introduced into commerce prior to the initial introduction into commerce of a controlled substance in its finished dosage form which it may substantially resemble.

Nothing in this subsection (y) prohibits the dispensing or distributing of noncontrolled substances by persons authorized

to dispense and distribute controlled substances under this Act, provided that such action would be deemed to be carried out in good faith under subsection (u) if the substances involved were controlled substances.

Nothing in this subsection (y) or in this Act prohibits the manufacture, preparation, propagation, compounding, processing, packaging, advertising or distribution of a drug or drugs by any person registered pursuant to Section 510 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360).

(y-1) "Mail-order pharmacy" means a pharmacy that is located in a state of the United States, other than Illinois, that delivers, dispenses or distributes, through the United States Postal Service or other common carrier, to Illinois residents, any substance which requires a prescription.

(z) "Manufacture" means the production, preparation, propagation, compounding, conversion or processing of a controlled substance, either directly or indirectly, by extraction from substances of natural origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis, and includes any packaging or repackaging of the substance or labeling of its container, except that this term does not include:

(1) by an ultimate user, the preparation or compounding of a controlled substance for his own use; or

(2) by a practitioner, or his authorized agent under his supervision, the preparation, compounding, packaging, or labeling of a controlled substance:

(a) as an incident to his administering or dispensing of a controlled substance in the course of his professional practice; or

(b) as an incident to lawful research, teaching or chemical analysis and not for sale.

(z-1) "Methamphetamine manufacturing chemical" means any of the following chemicals or substances containing any of the following chemicals: benzyl methyl ketone, ephedrine, methyl benzyl ketone, phenylacetone, phenyl-2-propanone,

pseudoephedrine, or red phosphorous or any of the salts, optical isomers, or salts of optical isomers of the above-listed chemicals.

(aa) "Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of natural origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:

(1) opium and opiate, and any salt, compound, derivative, or preparation of opium or opiate;

(2) any salt, compound, isomer, derivative, or preparation thereof which is chemically equivalent or identical with any of the substances referred to in clause (1), but not including the isoquinoline alkaloids of opium;

(3) opium poppy and poppy straw;

(4) coca leaves and any salts, compound, isomer, salt of an isomer, derivative, or preparation of coca leaves including cocaine or ecgonine, and any salt, compound, isomer, derivative, or preparation thereof which is chemically equivalent or identical with any of these substances, but not including decocainized coca leaves or extractions of coca leaves which do not contain cocaine or ecgonine (for the purpose of this paragraph, the term "isomer" includes optical, positional and geometric isomers).

(bb) "Nurse" means a registered nurse licensed under the Nursing and Advanced Practice Nursing Act.

(cc) (Blank).

(dd) "Opiate" means any substance having an addiction forming or addiction sustaining liability similar to morphine or being capable of conversion into a drug having addiction forming or addiction sustaining liability.

(ee) "Opium poppy" means the plant of the species Papaver somniferum L., except its seeds.

(ff) "Parole and Pardon Board" means the Parole and Pardon Board of the State of Illinois or its successor agency.

(gg) "Person" means any individual, corporation, mail-order pharmacy, government or governmental subdivision or agency, business trust, estate, trust, partnership or association, or any other entity.

(hh) "Pharmacist" means any person who holds a certificate of registration as a registered pharmacist, a local registered pharmacist or a registered assistant pharmacist under the Pharmacy Practice Act of 1987.

(ii) "Pharmacy" means any store, ship or other place in which pharmacy is authorized to be practiced under the Pharmacy Practice Act of 1987.

(jj) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.

(kk) "Practitioner" means a physician licensed to practice medicine in all its branches, dentist, optometrist, podiatrist, veterinarian, scientific investigator, pharmacist, physician assistant, advanced practice nurse, licensed practical nurse, registered nurse, hospital, laboratory, or pharmacy, or other person licensed, registered, or otherwise lawfully permitted by the United States or this State to distribute, dispense, conduct research with respect to, administer or use in teaching or chemical analysis, a controlled substance in the course of professional practice or research.

(ll) "Pre-printed prescription" means a written prescription upon which the designated drug has been indicated prior to the time of issuance.

(mm) "Prescriber" means a physician licensed to practice medicine in all its branches, dentist, optometrist, podiatrist, or veterinarian who issues a prescription, a physician assistant who issues a prescription for a Schedule III, IV, or V controlled substance in accordance with Section 303.05 and the written guidelines required under Section 7.5 of the Physician Assistant Practice Act of 1987, or an advanced practice nurse with prescriptive authority in accordance with Section 303.05 and a written collaborative agreement under

Sections 15-15 and 15-20 of the Nursing and Advanced Practice Nursing Act.

(nn) "Prescription" means a lawful written, facsimile, or verbal order of a physician licensed to practice medicine in all its branches, dentist, optometrist, podiatrist, or veterinarian for any controlled substance, of a physician assistant for a Schedule III, IV, or V controlled substance in accordance with Section 303.05 and the written guidelines required under Section 7.5 of the Physician Assistant Practice Act of 1987, or of an advanced practice nurse who issues a prescription for a Schedule III, IV, or V controlled substance in accordance with Section 303.05 and a written collaborative agreement under Sections 15-15 and 15-20 of the Nursing and Advanced Practice Nursing Act.

(oo) "Production" or "produce" means manufacture, planting, cultivating, growing, or harvesting of a controlled substance.

(pp) "Registrant" means every person who is required to register under Section 302 of this Act.

(qq) "Registry number" means the number assigned to each person authorized to handle controlled substances under the laws of the United States and of this State.

(rr) "State" includes the State of Illinois and any state, district, commonwealth, territory, insular possession thereof, and any area subject to the legal authority of the United States of America.

(ss) "Ultimate user" means a person who lawfully possesses a controlled substance for his own use or for the use of a member of his household or for administering to an animal owned by him or by a member of his household.

(Source: P.A. 92-449, eff. 1-1-02; 93-596, eff. 8-26-03; 93-626, eff. 12-23-03.)

(720 ILCS 570/103) (from Ch. 56 1/2, par. 1103)

Sec. 103. Scope of Act. Nothing in this Act limits the lawful authority granted by the Medical Practice Act of 1987,

the Nursing and Advanced Practice Nursing Act, the Illinois Optometric Practice Act of 1987, or the Pharmacy Practice Act of 1987.

(Source: P.A. 90-742, eff. 8-13-98.)