AN ACT in relation to animals.

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

Section 5. The Humane Euthanasia in Animal Shelters Act is amended by changing Sections 35, 55, and 57 as follows:

(510 ILCS 72/35)

Sec. 35. Technician certification; duties.

(a) An applicant for certification as a euthanasia technician shall file an application with the Department and shall:

(1) Be 18 years of age.

(2) Be of good moral character. In determining moral character under this Section, the Department may take into consideration whether the applicant has engaged in conduct or activities that would constitute grounds for discipline under this Act.

(3) Each applicant for certification as a euthanasia technician shall have his or her fingerprints submitted to the Department of State Police in an electronic format that complies with the form and manner for requesting and furnishing criminal history record information as prescribed by the Department of State Police. These fingerprints shall be checked against the Department of State Police and Federal Bureau of Investigation criminal history record databases now and hereafter filed. The Department of State Police shall charge applicants a fee for conducting the criminal history records check, which shall be deposited in the State Police Services Fund and shall not exceed the actual cost of the records check. The Department of State Police shall furnish, pursuant to positive

identification, records of Illinois convictions to the Department. ~~Submit--fingerprints--to--the-Illinois-State Police-or-its-designated-vendor-as--set--forth--by--rule. These--fingerprints-shall-be-checked-against-the-Illinois State-Police-and-Federal-Bureau-of-Investigation-criminal history--record--databases.---A--separate--fee--shall--be charged-to--the--applicant--for--fingerprinting,--payable either--to-the-Department-or-the-Illinois-State-Police-or its-designated-vendor.~~

(4) Hold a ~~current~~ license or certification from the American Humane Association, the National Animal Control Association, the Illinois Federation of Humane Societies, or the Humane Society of the United States issued within 3 years preceding the date of application.

For a period of 12 months after the adoption of final administrative rules for this Act, the Department may issue a certification to an applicant who holds a license or certification from the American Humane Association, the National Animal Control Association, the Illinois Federation of Humane Societies, or the Humane Society of the United States issued after January 1, 1997.

(5) Pay the required fee.

(b) The duties of a euthanasia technician shall include but are not limited to:

(1) preparing animals for euthanasia and scanning each animal, prior to euthanasia, for microchips;

(2) accurately recording the dosages administered and the amount of drugs wasted;

(3) ordering supplies;

(4) maintaining the security of all controlled substances and drugs;

(5) humanely euthanizing animals via intravenous injection by hypodermic needle, intraperitoneal injection by hypodermic needle, solutions or powder added to food

or by mouth, intracardiac injection only on comatose animals by hypodermic needle, or carbon monoxide in a commercially manufactured chamber; and

(6) properly disposing of euthanized animals after verification of death.

(c) A euthanasia technician employed by a euthanasia agency may perform euthanasia by the administration of a Schedule II or Schedule III nonnarcotic controlled substance. A euthanasia technician may not personally possess, order, or administer a controlled substance except as an agent of the euthanasia agency.

(d) Upon termination from a euthanasia agency, a euthanasia technician shall not perform animal euthanasia until he or she is employed by another certified euthanasia agency.

(e) A certified euthanasia technician or an instructor in an approved course does not engage in the practice of veterinary medicine when performing duties set forth in this Act.

(Source: P.A. 92-449, eff. 1-1-02.)

(510 ILCS 72/55)

Sec. 55. Endorsement. An applicant, who is a euthanasia technician registered or licensed under the laws of another state or territory of the United States that has requirements that are substantially similar to the requirements of this Act, may be granted certification as a euthanasia technician in this State without examination, upon presenting satisfactory proof to the Department that the applicant has been engaged in the practice of euthanasia for a period of not less than one year and upon payment of the required fee. <u>In addition, an applicant shall have his or her fingerprints submitted to the Department of State Police for purposes of a criminal history records check pursuant to clause (a)(3) of</u>

Section 35.

(Source: P.A. 92-449, eff. 1-1-02.)

(510 ILCS 72/57)

Sec. 57. Procedures for euthanasia.

(a) Only euthanasia drugs and commercially compressed carbon monoxide, subject to the limitations imposed under subsection (b) of this Section, shall be used for the purpose of humanely euthanizing injured, sick, homeless, or unwanted companion animals in an animal shelter or an animal control facility licensed under the Illinois Animal Welfare Act.

(b) Commercially compressed carbon monoxide may be used as a permitted method of euthanasia provided that it is performed in a commercially manufactured chamber pursuant to the guidelines set forth in the most recent report of the AVMA Panel on Euthanasia. A chamber that is designed to euthanize more than one animal at a time must be equipped with independent sections or cages to separate incompatible animals. The interior of the chamber must be well lit and equipped with view-ports, a regulator, and a flow meter. Monitoring equipment must be used at all times during the operation. Animals that are under 4 months of age, old, injured, or sick may not be euthanized by carbon monoxide. Animals shall remain in the chamber and be exposed for a minimum of 20 minutes. Staff members shall be fully notified of potential health risks.

(c) Animals cannot be transported beyond State lines for the sole purpose of euthanasia unless the euthanasia methods comply with subsection (a) or (b) of this Section and the euthanasia is performed by a certified euthanasia technician.

(Source: P.A. 92-449, eff. 1-1-02.)

(510 ILCS 72/50 rep.)

Section 10. The Humane Euthanasia in Animal Shelters Act

is amended by repealing Section 50.

Section 15. The Illinois Controlled Substances Act is amended by changing Sections 102, 302, 303, 303.05, 304, and 306 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, ~~or~~ 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), ~~or~~

(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, 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, 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, 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.)

(720 ILCS 570/302) (from Ch. 56 1/2, par. 1302)

Sec. 302. (a) Every person who manufactures, distributes, or dispenses any controlled substances, or engages in chemical analysis, and instructional activities which utilize controlled substances, <u>or who purchases, stores, or administers euthanasia drugs,</u> within this State or who proposes to engage in the manufacture, distribution, or dispensing of any controlled substance, or to engage in chemical analysis, and instructional activities which utilize controlled substances, <u>or to engage in purchasing, storing, or administering euthanasia drugs,</u> within this State, must obtain a registration issued by the Department of Professional Regulation in accordance with its rules. The rules shall include, but not be limited to, setting the expiration date and renewal period for each registration under this Act. The Department, and any facility or service licensed by the Department, shall be exempt from the regulation requirements of this Section.

(b) Persons registered by the Department of Professional Regulation under this Act to manufacture, distribute, or dispense controlled substances, or purchase, store, or administer euthanasia drugs, may possess, manufacture, distribute, or dispense those substances, or purchase, store, or administer euthanasia drugs, to the extent authorized by their registration and in conformity with the other provisions of this Article.

(c) The following persons need not register and may lawfully possess controlled substances under this Act:

(1) an agent or employee of any registered manufacturer, distributor, or dispenser of any controlled substance if he is acting in the usual course of his employer's lawful business or employment;

(2) a common or contract carrier or warehouseman, or an agent or employee thereof, whose possession of any controlled substance is in the usual lawful course of such business or employment;

(3) an ultimate user or a person in possession of any controlled substance pursuant to a lawful prescription of a practitioner or in lawful possession of a Schedule V substance;

(4) officers and employees of this State or of the United States while acting in the lawful course of their official duties which requires possession of controlled substances;

(5) a registered pharmacist who is employed in, or the owner of, a pharmacy licensed under this Act and the Federal Controlled Substances Act, at the licensed location, or if he is acting in the usual course of his lawful profession, business, or employment.

(d) A separate registration is required at each place of business or professional practice where the applicant manufactures, distributes, or dispenses controlled

substances, or purchases, stores, or administers euthanasia drugs. Persons are required to obtain a separate registration for each place of business or professional practice where controlled substances are located or stored. A separate registration is not required for every location at which a controlled substance may be prescribed.

(e) The Department of Professional Regulation or the Department of State Police may inspect the controlled premises, as defined in Section 502 of this Act, of a registrant or applicant for registration in accordance with this Act and the rules promulgated hereunder and with regard to persons licensed by the Department, in accordance with subsection (bb) of Section 30-5 of the Alcoholism and Other Drug Abuse and Dependency Act and the rules and regulations promulgated thereunder.

(Source: P.A. 87-711; 88-670, eff. 12-2-94.)

(720 ILCS 570/303) (from Ch. 56 1/2, par. 1303)

Sec. 303. (a) The Department of Professional Regulation shall license an applicant to manufacture, distribute or dispense controlled substances included in Sections 204, 206, 208, 210 and 212 of this Act or purchase, store, or administer euthanasia drugs unless it determines that the issuance of that license would be inconsistent with the public interest. In determining the public interest, the Department of Professional Regulation shall consider the following:

(1) maintenance of effective controls against diversion of controlled substances into other than lawful medical, scientific, or industrial channels;

(2) compliance with applicable Federal, State and local law;

(3) any convictions of the applicant under any law of the United States or of any State relating to any

controlled substance;

(4) past experience in the manufacture or distribution of controlled substances, and the existence in the applicant's establishment of effective controls against diversion;

(5) furnishing by the applicant of false or fraudulent material in any application filed under this Act;

(6) suspension or revocation of the applicant's Federal registration to manufacture, distribute, or dispense controlled substances, or purchase, store, or administer euthanasia drugs, as authorized by Federal law;

(7) whether the applicant is suitably equipped with the facilities appropriate to carry on the operation described in his application;

(8) whether the applicant is of good moral character or, if the applicant is a partnership, association, corporation or other organization, whether the partners, directors, governing committee and managing officers are of good moral character;

(9) any other factors relevant to and consistent with the public health and safety; and

(10) Evidence from court, medical disciplinary and pharmacy board records and those of State and Federal investigatory bodies that the applicant has not or does not prescribe controlled substances within the provisions of this Act.

(b) No license shall be granted to or renewed for any person who has within 5 years been convicted of a wilful violation of any law of the United States or any law of any State relating to controlled substances, or who is found to be deficient in any of the matters enumerated in subsections (a)(1) through (a)(8).

(c) Licensure under subsection (a) does not entitle a registrant to manufacture, distribute or dispense controlled substances in Schedules I or II other than those specified in the registration.

(d) Practitioners who are licensed to dispense any controlled substances in Schedules II through V are authorized to conduct instructional activities with controlled substances in Schedules II through V under the law of this State.

(e) If an applicant for registration is registered under the Federal law to manufacture, distribute or dispense controlled substances, or purchase, store, or administer euthanasia drugs, upon filing a completed application for licensure in this State and payment of all fees due hereunder, he shall be licensed in this State to the same extent as his Federal registration, unless, within 30 days after completing his application in this State, the Department of Professional Regulation notifies the applicant that his application has not been granted. A practitioner who is in compliance with the Federal law with respect to registration to dispense controlled substances in Schedules II through V need only send a current copy of that Federal registration to the Department of Professional Regulation and he shall be deemed in compliance with the registration provisions of this State.

(e-5) Beginning July 1, 2003, all of the fees and fines collected under this Section 303 shall be deposited into the Illinois State Pharmacy Disciplinary Fund.

(f) The fee for registration as a manufacturer or wholesale distributor of controlled substances shall be $50.00 per year, except that the fee for registration as a manufacturer or wholesale distributor of controlled substances that may be dispensed without a prescription under this Act shall be $15.00 per year. The expiration date and

renewal period for each controlled substance license issued under this Act shall be set by rule.

(Source: P.A. 93-32, eff. 7-1-03.)

(720 ILCS 570/303.05)

Sec. 303.05. Mid-level practitioner registration.

(a) The Department of Professional Regulation shall register licensed physician assistants and licensed advanced practice nurses to prescribe and dispense Schedule III, IV, or V controlled substances under Section 303 and euthanasia agencies to purchase, store, or administer euthanasia drugs under the following circumstances:

(1) with respect to physician assistants or advanced practice nurses,

(A) the physician assistant or advanced practice nurse has been delegated prescriptive authority by a physician licensed to practice medicine in all its branches in accordance with Section 7.5 of the Physician Assistant Practice Act of 1987 or Section 15-20 of the Nursing and Advanced Practice Nursing Act; and

(B) ~~(2)~~ the physician assistant or advanced practice nurse has completed the appropriate application forms and has paid the required fees as set by rule; or~~.~~

(2) with respect to euthanasia agencies, the euthanasia agency has obtained a license from the Department of Professional Regulation and obtained a registration number from the Department.

(b) The mid-level practitioner shall only be licensed to prescribe those schedules of controlled substances for which a licensed physician has delegated prescriptive authority~~,~~ except that a euthanasia agency does not have any prescriptive authority.

(c) Upon completion of all registration requirements, physician assistants, ~~and~~ advanced practice nurses, and euthanasia agencies shall be issued a mid-level practitioner controlled substances license for Illinois.

(Source: P.A. 90-818, eff. 3-23-99.)

(720 ILCS 570/304) (from Ch. 56 1/2, par. 1304)

Sec. 304. (a) A registration under Section 303 to manufacture, distribute, or dispense a controlled substance or purchase, store, or administer euthanasia drugs may be suspended or revoked by the Department of Professional Regulation upon a finding that the registrant:

(1) has furnished any false or fraudulent material information in any application filed under this Act; or

(2) has been convicted of a felony under any law of the United States or any State relating to any controlled substance; or

(3) has had suspended or revoked his Federal registration to manufacture, distribute, or dispense controlled substances or purchase, store, or administer euthanasia drugs; or

(4) has been convicted of bribery, perjury, or other infamous crime under the laws of the United States or of any State; or

(5) has violated any provision of this Act or any rules promulgated hereunder, whether or not he has been convicted of such violation; or

(6) has failed to provide effective controls against the diversion of controlled substances in other than legitimate medical, scientific or industrial channels.

(b) The Department of Professional Regulation may limit revocation or suspension of a registration to the particular controlled substance with respect to which grounds for revocation or suspension exist.

(c) The Department of Professional Regulation shall promptly notify the Administration, the Department and the Department of State Police or their successor agencies, of all orders denying, suspending or revoking registration, all forfeitures of controlled substances, and all final court dispositions, if any, of such denials, suspensions, revocations or forfeitures.

(d) If Federal registration of any registrant is suspended, revoked, refused renewal or refused issuance, then the Department of Professional Regulation shall issue a notice and conduct a hearing in accordance with Section 305 of this Act.

(Source: P.A. 85-1209.)

(720 ILCS 570/306) (from Ch. 56 1/2, par. 1306)

Sec. 306. Every practitioner and person who is required under this Act to be registered to manufacture, distribute or dispense controlled substances <u>or purchase, store, or administer euthanasia drugs</u> under this Act shall keep records and maintain inventories in conformance with the recordkeeping and inventory requirements of the laws of the United States and with any additional rules and forms issued by the Department of Professional Regulation.

(Source: P.A. 89-202, eff. 10-1-95.)

Section 99. Effective date. This Act takes effect upon becoming law.