AN ACT concerning regulation.

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

Section 5. The Nurse Practice Act is amended by changing Section 65-40 as follows:

(225 ILCS 65/65-40) (was 225 ILCS 65/15-20)

(Section scheduled to be repealed on January 1, 2018)

Sec. 65-40. Prescriptive authority.

(a) A collaborating physician or podiatrist may, but is not required to, delegate prescriptive authority to an advanced practice nurse as part of a written collaborative agreement. This authority may, but is not required to, include prescription of, selection of, orders for, administration of, storage of, acceptance of samples of, and dispensing over the counter medications, legend drugs, medical gases, and controlled substances categorized as any Schedule III through~~, III N, IV, or~~ V controlled substances, as defined in Article II of the Illinois Controlled Substances Act, and other preparations, including, but not limited to, botanical and herbal remedies. The collaborating physician or podiatrist must have a valid current Illinois controlled substance license and federal registration to delegate authority to prescribe delegated controlled substances.

(b) To prescribe controlled substances under this Section, an advanced practice nurse must obtain a mid-level practitioner controlled substance license. Medication orders shall be reviewed periodically by the collaborating physician or podiatrist.

(c) The collaborating physician or podiatrist shall file with the Department notice of delegation of prescriptive authority and termination of such delegation, in accordance with rules of the Department. Upon receipt of this notice delegating authority to prescribe any Schedule III through~~, III-N, IV, or~~ V controlled substances, the licensed advanced practice nurse shall be eligible to register for a mid-level practitioner controlled substance license under Section 303.05 of the Illinois Controlled Substances Act.

(d) In addition to the requirements of subsections (a), (b), and (c) of this Section, a collaborating physician may, but is not required to, delegate authority to an advanced practice nurse to prescribe any Schedule II ~~or II-N~~ controlled substances, if all of the following conditions apply:

(1) No more than 5 Schedule II ~~or II-N~~ controlled substances by oral dosage may be delegated.

(2) Any delegation must be controlled substances that the collaborating physician prescribes.

(3) Any prescription must be limited to no more than a 30-day oral dosage, with any continuation authorized only after prior approval of the collaborating physician.

(4) The advanced practice nurse must discuss the condition of any patients for whom a controlled substance is prescribed monthly with the delegating physician.

(e) Nothing in this Act shall be construed to limit the delegation of tasks or duties by a physician to a licensed practical nurse, a registered professional nurse, or other persons.

(Source: P.A. 95-639, eff. 10-5-07.)

Section 10. The Pharmacy Practice Act is amended by changing Section 4 as follows:

(225 ILCS 85/4) (from Ch. 111, par. 4124)

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

Sec. 4. Exemptions. Nothing contained in any Section of this Act shall apply to, or in any manner interfere with:

(a) the lawful practice of any physician licensed to practice medicine in all of its branches, dentist, podiatrist, veterinarian, or therapeutically or diagnostically certified optometrist within the limits of his or her license, or prevent him or her from supplying to his or her bona fide patients such drugs, medicines, or poisons as may seem to him appropriate;

(b) the sale of compressed gases;

(c) the sale of patent or proprietary medicines and household remedies when sold in original and unbroken packages only, if such patent or proprietary medicines and household

remedies be properly and adequately labeled as to content and usage and generally considered and accepted as harmless and nonpoisonous when used according to the directions on the label, and also do not contain opium or coca leaves, or any compound, salt or derivative thereof, or any drug which, according to the latest editions of the following authoritative pharmaceutical treatises and standards, namely, The United States Pharmacopoeia/National Formulary (USP/NF), the United States Dispensatory, and the Accepted Dental Remedies of the Council of Dental Therapeutics of the American Dental Association or any or either of them, in use on the effective date of this Act, or according to the existing provisions of the Federal Food, Drug, and Cosmetic Act and Regulations of the Department of Health and Human Services, Food and Drug Administration, promulgated thereunder now in effect, is designated, described or considered as a narcotic, hypnotic, habit forming, dangerous, or poisonous drug;

(d) the sale of poultry and livestock remedies in original and unbroken packages only, labeled for poultry and livestock medication;

(e) the sale of poisonous substances or mixture of poisonous substances, in unbroken packages, for nonmedicinal use in the arts or industries or for insecticide purposes; provided, they are properly and adequately labeled as to content and such nonmedicinal usage, in conformity with the provisions of all applicable federal, state and local laws and

regulations promulgated thereunder now in effect relating thereto and governing the same, and those which are required under such applicable laws and regulations to be labeled with the word "Poison", are also labeled with the word "Poison" printed thereon in prominent type and the name of a readily obtainable antidote with directions for its administration;

(f) the delegation of limited prescriptive authority by a physician licensed to practice medicine in all its branches to a physician assistant under Section 7.5 of the Physician Assistant Practice Act of 1987. This delegated authority under Section 7.5 of the Physician Assistant Practice Act of 1987 may but is not required to include prescription of controlled substances, as defined in Article II of the Illinois Controlled Substances Act, in accordance with written guidelines; and

(g) The delegation of prescriptive authority by a physician licensed to practice medicine in all its branches or a licensed podiatrist to an advanced practice nurse in accordance with a written collaborative agreement under Sections ~~Section~~ 65-35 and 65-40 of the Nurse Practice Act. ~~This authority, which is delegated under Section 65-40 of the Nurse Practice Act, may but is not required to include the prescription of Schedule III, IV, or V controlled substances as defined in Article II of the Illinois Controlled Substances Act.~~

(Source: P.A. 95-639, eff. 10-5-07.)

Section 15. The Illinois Controlled Substances Act is

amended by changing Sections 102 and 303.05 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 other than methamphetamine, 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) (Blank).

(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 Nurse Practice 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 license or certificate of registration as a registered pharmacist, a local registered pharmacist or a registered assistant pharmacist under the Pharmacy Practice Act.

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

(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 delegated under Section 65-40 of the Nurse Practice Act and in accordance with Section 303.05, a written delegation, and a written

collaborative agreement under Section 65-35 of the Nurse Practice 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 an optometrist for a Schedule III, IV, or V controlled substance in accordance with Section 15.1 of the Illinois Optometric Practice Act of 1987, 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 with prescriptive authority delegated under Section 65-40 of the Nurse Practice Act who issues a prescription for a ~~Schedule III, IV, or V~~ controlled substance in accordance with Section 303.05, a written delegation, and a written collaborative agreement under Section 65-35 of the Nurse Practice Act.

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

(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. 94-556, eff. 9-11-05; 95-242, eff. 1-1-08; 95-639, eff. 10-5-07; 95-689, eff. 10-29-07; revised 11-19-07.)

(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 animal 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 to prescribe any Schedule III through V controlled substances 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 65-40 of the Nurse Practice Act~~; and

(B) 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 advanced practice nurses,

(A) the advanced practice nurse has been delegated authority to prescribe any Schedule III through V controlled substances by a physician licensed to practice medicine in all its branches or a podiatrist in accordance with Section 65-40 of the Nurse Practice Act. The advanced practice nurse has completed the appropriate application forms and has paid the required fees; or

(B) the advanced practice nurse has been delegated authority by a collaborating physician licensed to practice medicine in all its branches to prescribe or dispense Schedule II controlled substances through a written delegation of authority and under the following conditions:

(i) no more than 5 Schedule II controlled substances by oral dosage may be delegated;

(ii) any delegation must be of controlled substances prescribed by the collaborating physician;

(iii) all prescriptions must be limited to no

more than a 30-day oral dosage, with any continuation authorized only after prior approval of the collaborating physician;

(iv) the advanced practice nurse must discuss the condition of any patients for whom a controlled substance is prescribed monthly with the delegating physician; and

(v) the advanced practice nurse must have completed the appropriate application forms and paid the required fees as set by rule; or

(3) ~~(2)~~ with respect to animal 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 or licensed podiatrist has delegated prescriptive authority, except that an animal ~~a~~ euthanasia agency does not have any prescriptive authority. A physician assistant and an advanced practice nurse are prohibited from prescribing medications and controlled substances not set forth in the required written delegation of authority.

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

(d) Notwithstanding any other rulemaking authority that may exist, neither the Governor nor any agency or agency head under the jurisdiction of the Governor has any authority to make or promulgate rules to implement or enforce the provisions of this amendatory Act of the 95th General Assembly. If, however, the Governor believes that rules are necessary to implement or enforce the provisions of this amendatory Act of the 95th General Assembly, the Governor may suggest rules to the General Assembly by filing them with the Clerk of the House and the Secretary of the Senate and by requesting that the General Assembly authorize such rulemaking by law, enact those suggested rules into law, or take any other appropriate action in the General Assembly's discretion. Nothing contained in this amendatory Act of the 95th General Assembly shall be interpreted to grant rulemaking authority under any other Illinois statute where such authority is not otherwise explicitly given. For the purposes of this amendatory Act of the 95th General Assembly, "rules" is given the meaning contained in Section 1-70 of the Illinois Administrative Procedure Act, and "agency" and "agency head" are given the meanings contained in Sections 1-20 and 1-25 of the Illinois Administrative Procedure Act to the extent that such definitions apply to agencies or agency heads under the jurisdiction of the Governor.

(Source: P.A. 95-639, eff. 10-5-07.)