Session of 2019

## HOUSE BILL No. 2066

By Committee on Health and Human Services

1-23

 AN ACT concerning advanced practice registered nurses; board of nursing; relating to definition of practice; prescribing authority; licensure requirements; rules and regulations; amending K.S.A. 65-1130 and 65-4101 and K.S.A. 2017 Supp. 65-1113, as amended by section 2 of of chapter 42 of the 2018 Session Laws of Kansas and repealing the existing sections.

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

9 Section 1. K.S.A. 2017 Supp. 65-1113, as amended by section 2 of 10 chapter 42 of the 2018 Session Laws of Kansas, is hereby amended to read 11 as follows: 65-1113. When used in this act and the act of which this 12 section is amendatory:

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(a) "Board" means the board of nursing.

(b) "Diagnosis" in the context of nursing practice means<u>that</u> the
identification of and discrimination between physical and psychosocial
signs and symptoms essential to effective<u>execution</u> implementation and
management of the nursing regimen and shall be construed as distinct from
a medical diagnosis patient's healthcare, determined by the nurse's level of
education.

(c) "Treatment" means the selection and performance of those
 therapeutic measures essential to effective-execution *implementation* and
 management of the nursing regimen, and any prescribed medical regimen
 *patient's healthcare, determined by the nurse's level of education.*

(d) Practice of nursing. (1) The practice of professional nursing as 24 25 performed by a registered professional nurse for compensation or 26 gratuitously, except as permitted by K.S.A. 65-1124, and amendments 27 thereto, means the process in which substantial specialized knowledge 28 derived from the biological, physical, and behavioral sciences is applied 29 to: the: Care, diagnosis, treatment, counsel and health teaching of persons 30 who are experiencing changes in the normal health processes or who 31 require assistance in the maintenance of health or the prevention or 32 management of illness, injury or infirmity; administration, supervision or 33 teaching of the process as defined in this section; and the execution of the 34 medical *treatment* regimen as prescribed by a person licensed to practice 35 medicine and surgery-or, a person licensed to practice dentistry or a 36 person licensed to practice advanced practice registered nursing.

1 (2) The practice of nursing as a licensed practical nurse means the performance for compensation or gratuitously, except as permitted by 2 K.S.A. 65-1124, and any amendments thereto, of tasks and responsibilities 3 defined in paragraph (1), which tasks and responsibilities are based on 4 acceptable educational preparation within the framework of supportive and 5 6 restorative care under the direction of a registered professional nurse, a 7 person licensed to practice medicine and surgery or a person licensed to 8 practice dentistry.

9 (3) The practice of professional nursing as an advanced practice registered nurse as defined in subsection (g) within the APRN role means, 10 in addition to the practice and responsibilities of professional nursing as 11 12 defined in paragraph (1): Conducting an advanced assessment; ordering and interpreting diagnostic procedures; establishing primary and 13 differential diagnoses; prescribing, ordering, administering and furnishing 14 therapeutic measures as set forth by the board; delegating and assigning 15 16 therapeutic measures to assistive personnel; collaborating and consulting with physicians and other healthcare providers; providing referrals to 17 healthcare providers, agencies and community resources; and other acts 18 19 that require education and training consistent with the professional 20 standards and commensurate with the APRN's education, certification, 21 demonstrated competencies and experience.

(e) A "professional nurse" means a person who is licensed to practiceprofessional nursing as defined in subsection (d)(1).

(f) A "practical nurse" means a person who is licensed to practicepractical nursing as defined in subsection (d)(2).

26 (g) "Advanced practice registered nurse" or "APRN" means a 27 professional nurse who holds a license from the board to-function practice 28 advanced practice registered nursing as defined in subsection (d)(3) as a 29 professional nurse in an advanced role, and this advanced role-shall may be 30 further defined by rules and regulations consistent with the Kansas nurse 31 practice act adopted by the board in accordance with K.S.A. 65-1130, and 32 amendments thereto.

(h) "Continuing nursing education" means learning experiences
intended to build upon the educational and experiential bases of the
registered professional and licensed practical nurse for the enhancement of
practice, education, administration, research or theory development to the
end of improving the health of the public.

(i) "Collaboration" means the process in which two or more
 healthcare professionals work together to meet the healthcare needs of a
 patient, as warranted by the patient.

41 *(j)* "Consultation" means the process in which an advanced practice 42 registered nurse who maintains primary management responsibility for a 43 patient's care seeks advice or opinion of a physician or another member of 1 the healthcare team.

Sec. 2. K.S.A. 65-1130 is hereby amended to read as follows: 65-1130. (a) No professional nurse shall announce or represent to the public that such person is an advanced practice registered nurse unless such professional nurse has complied with requirements established by the board and holds a valid license as an advanced practice registered nurse in accordance with the provisions of this section.

8 (b) (1) The board shall establish standards and requirements for any 9 professional nurse who desires to obtain licensure as an advanced practice 10 registered nurse. Such standards and requirements shall include, but not be 11 limited to, standards and requirements relating to the education of 12 advanced practice registered nurses. The board may give such 13 examinations and secure such assistance as it deems necessary to 14 determine the qualifications of applicants.

15 (2) On and after July 1, 2020, for an applicant, an initial advanced 16 practice registered nurse license shall have a current advanced practice 17 registered nurse certification in such applicant's specific role granted by a 18 national certifying organization recognized by the board whose 19 certification standards are approved by the board as equal to or greater 20 than the corresponding standards established by the board.

(c) The board shall adopt rules and regulations *consistent with the Kansas nurse practice act* applicable to advanced practice registered
 nurses which that:

(1) Establish roles and identify titles and abbreviations of advanced
 practice registered nurses-which *that* are consistent with nursing practice
 specialties recognized by the nursing profession *including titles describing the four APRN roles of certified registered nurse anesthetist, clinical nurse specialist, certified nurse midwife and certified nurse practitioner.*

29 (2) Establish education and qualifications necessary for licensure for each role of advanced practice registered nurse established by the board-at 30 31 a level adequate to assure the competent performance by advanced-32 practice registered nurses of functions and procedures which advanced-33 practice registered nurses are authorized to perform. Advanced practice-34 registered nursing is based on knowledge and skills acquired in. Education 35 and qualifications for APRN licensure established by the board shall 36 include completion of basic nursing education, licensure as a registered 37 nurse and graduation from or completion of a master's or higher degree an 38 accredited graduate or post-graduate level APRN program in one of the 39 advanced practice registered nurse roles approved by the board of nursing.

40 (3) Define the role of advanced practice registered nurses and 41 establish limitations and restrictions on such role *consistent with the* 42 *Kansas nurse practice act.* The board shall adopt a definition of the role 43 <del>under this paragraph which *that* is consistent with the education and</del>

qualifications required to obtain a license as an advanced practice 1 2 registered nurse, which that protects the public from persons performing 3 functions and procedures as advanced practice registered nurses for which 4 they lack adequate education and gualifications and which that authorizes 5 advanced practice registered nurses to perform acts generally recognized 6 by the profession of nursing as capable of being performed, in a manner 7 consistent with the public health and safety, by persons with postbasic 8 education in nursing. In defining such role the board shall consider:

9 (A) The education required for a licensure as an advanced practice 10 registered nurse;

(B) the type of nursing practice and preparation in specialized
 advanced practice skills involved in each role of advanced practice
 registered nurse established by the board;

14 (C) the scope and limitations of advanced practice nursing prescribed 15 by national advanced practice organizations. *Advanced practice nursing is* 16 *built on the practice of health promotion, health maintenance, illness* 17 *prevention, diagnosis, treatment and management of common health* 18 *problems and acute and chronic conditions*; and

(D) acts recognized by the nursing profession as appropriate to beperformed by persons with postbasic education in nursing.

(4) Require an advanced practice registered nurse to wear
identification that clearly identifies the nurse as such when providing
direct patient care, unless wearing identification creates a safety or health
risk to the nurse or patient.

25 (d) (1) An advanced practice registered nurse may prescribe drugs pursuant to a written protocol as authorized by a responsible physician. 26 Each written protocol shall contain a precise and detailed medical plan of 27 28 eare for each elassification of disease or injury for which the advanced 29 practice registered nurse is authorized to prescribe and shall specify all-30 drugs which may be prescribed by the advanced practice registered nurse. 31 Any written, procure and administer prescription drugs and controlled 32 substances in schedules II through V pursuant to applicable federal and 33 state laws

*(2)* A prescription order shall include the name, address and telephone
 number of the responsible physician. The advanced practice registered
 *nurse*. An advanced practice registered nurse may not dispense drugs, but
 may request, receive and sign for professional samples and may distribute
 professional samples to patients pursuant to a written protocol as authorized by a responsible physician.

40 (3) In order to prescribe controlled substances, the advanced practice 41 registered nurse shall: (1)

42 (A) Register with the federal drug enforcement administration; and 43 (2)

1 (B) notify the board of the name and address of the responsible 2 physician or physicians. In no case shall the scope of authority of the advanced practice registered nurse exceed the normal and customary-3 practice of the responsible physician federal drug enforcement 4 administration registration as prescribed by the rules and regulations of 5 6 the board. An advanced practice registered nurse shall comply with the 7 federal drug enforcement administration requirements related to 8 controlled substances.

9 (4) An advanced practice registered nurse certified in the role of registered nurse anesthetist while functioning as a registered nurse 10 anesthetist under K.S.A. 65-1151 through 65-1164, and amendments 11 thereto, shall be subject to the provisions of K.S.A. 65-1151 through 65-12 1164, and amendments thereto, with respect to drugs and anesthetic agents 13 and shall not be subject to the provisions of this subsection. For the 14 purposes of this subsection, "responsible physician" means a person-15 16 licensed to practice medicine and surgery in Kansas who has accepted-17 responsibility for the protocol and the actions of the advanced practice-18 registered nurse when prescribing drugs.

19 (5) An advanced practice registered nurse shall maintain malpractice 20 insurance coverage in effect as a condition of rendering professional 21 service as an advanced practice registered nurse in this state and shall 22 provide proof of insurance at the time of licensure and renewal of license. 23 The requirements of this paragraph shall not apply to an advanced practice registered nurse who: Practices solely in employment for which 24 25 the advanced practice registered nurse is covered under the federal tort claims act or Kansas tort claims act; practices solely as a charitable 26 27 healthcare provider under K.S.A. 75-6102, and amendments thereto; or is 28 serving on active duty in the military service of the United States.

(e) As used in this section, "drug" means those articles and substances
 defined as drugs in K.S.A. 65-1626 and 65-4101, and amendments thereto.

(f) A person registered to practice as an advanced registered nurse practitioner in the state of Kansas immediately prior to the effective date of this act shall be deemed to be licensed to practice as an advanced practice registered nurse under this act and such person shall not be required to file an original application for licensure under this act. Any application for registration filed-which *that* has not been granted prior to the effective date of this act shall be processed as an application for licensure under this act.

(g) An advanced practice registered nurse certified in the role of certified nurse-midwife and engaging in the independent practice of midwifery under the independent practice of midwifery act with respect to prescribing drugs shall be subject to the provisions of the independent practice of midwifery act and shall not be subject to the provisions of this section. 1 Sec. 3. K.S.A. 65-4101 is hereby amended to read as follows: 65-2 4101. As used in this act: (a) "Administer" means the direct application of 3 a controlled substance, whether by injection, inhalation, ingestion or any 4 other means, to the body of a patient or research subject by:

5 (1) A practitioner or pursuant to the lawful direction of a practitioner; 6 or

7 8 (2) the patient or research subject at the direction and in the presence of the practitioner.

9 (b) "Agent" means an authorized person who acts on behalf of or at 10 the direction of a manufacturer, distributor or dispenser. It does not include 11 a common carrier, public warehouseman or employee of the carrier or 12 warehouseman.

(c) "Application service provider" means an entity that sells
 electronic prescription or pharmacy prescription applications as a hosted
 service where the entity controls access to the application and maintains
 the software and records on its server.

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(d) "Board" means the state board of pharmacy.

(e) "Bureau" means the bureau of narcotics and dangerous drugs,United States department of justice, or its successor agency.

(f) "Controlled substance" means any drug, substance or immediate
precursor included in any of the schedules designated in K.S.A. 65-4105,
65-4107, 65-4109, 65-4111 and 65-4113, and amendments thereto.

(g) (1) "Controlled substance analog" means a substance that isintended for human consumption, and at least one of the following:

(A) The chemical structure of the substance is substantially similar to
the chemical structure of a controlled substance listed in or added to the
schedules designated in K.S.A. 65-4105 or 65-4107, and amendments
thereto;

(B) the substance has a stimulant, depressant or hallucinogenic effect
on the central nervous system substantially similar to the stimulant,
depressant or hallucinogenic effect on the central nervous system of a
controlled substance included in the schedules designated in K.S.A. 654105 or 65-4107, and amendments thereto; or

(C) with respect to a particular individual, such individual represents
or intends the substance to have a stimulant, depressant or hallucinogenic
effect on the central nervous system substantially similar to the stimulant,
depressant or hallucinogenic effect on the central nervous system of a
controlled substance included in the schedules designated in K.S.A. 654105 or 65-4107, and amendments thereto.

(2) "Controlled substance analog" does not include:

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- (A) A controlled substance;
- 42 (B) a substance for which there is an approved new drug application;
- 43 or

1 (C) a substance with respect to which an exemption is in effect for 2 investigational use by a particular person under section 505 of the federal food, drug and cosmetic act, 21 U.S.C. § 355, to the extent conduct with 3 respect to the substance is permitted by the exemption. 4

5 (h) "Counterfeit substance" means a controlled substance-which that, 6 or the container or labeling of which, without authorization bears the 7 trademark, trade name or other identifying mark, imprint, number or device or any likeness thereof of a manufacturer, distributor or dispenser 8 9 other than the person who in fact manufactured, distributed or dispensed 10 the substance.

11 (i) "Cultivate" means the planting or promotion of growth of five or 12 more plants-which that contain or can produce controlled substances.

"DEA" means the U.S. department of justice, drug enforcement 13 (i) 14 administration

(k) "Deliver" or "delivery" means the actual, constructive or 15 16 attempted transfer from one person to another of a controlled substance, 17 whether or not there is an agency relationship.

(1) "Dispense" means to deliver a controlled substance to an ultimate 18 19 user or research subject by or pursuant to the lawful order of a practitioner, 20 including the packaging, labeling or compounding necessary to prepare the 21 substance for that delivery, or pursuant to the prescription of a mid-level 22 practitioner.

23 (m) "Dispenser" means a practitioner or pharmacist who dispenses, or 24 a physician assistant who has authority to dispense prescription-only drugs 25 in accordance with K.S.A. 65-28a08(b), and amendments thereto.

(n) "Distribute" means to deliver other than by administering or 26 27 dispensing a controlled substance.

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

29 (p) "Drug" means: (1) Substances recognized as drugs in the official United States pharmacopeia, official homeopathic pharmacopoeia of the 30 31 United States or official national formulary or any supplement to any of 32 them; (2) substances intended for use in the diagnosis, cure, mitigation, 33 treatment or prevention of disease in human or animals; (3) substances 34 (other than food) intended to affect the structure or any function of the 35 body of human or animals; and (4) substances intended for use as a 36 component of any article specified in paragraph (1), (2) or (3). It does not 37 include devices or their components, parts or accessories.

38 (q) "Immediate precursor" means a substance-which that the board 39 has found to be and by rule and regulation designates as being the principal compound commonly used or produced primarily for use and 40 41 which that is an immediate chemical intermediary used or likely to be used 42 in the manufacture of a controlled substance, the control of which is 43 necessary to prevent, curtail or limit manufacture.

1 (r) "Electronic prescription" means an electronically prepared 2 prescription that is authorized and transmitted from the prescriber to the 3 pharmacy by means of electronic transmission.

4 (s) "Electronic prescription application" means software that is used 5 to create electronic prescriptions and that is intended to be installed on the 6 prescriber's computers and servers where access and records are controlled 7 by the prescriber.

8 (t) "Electronic signature" means a confidential personalized digital 9 key, code, number or other method for secure electronic data transmissions 10 <del>which</del> *that* identifies a particular person as the source of the message, 11 authenticates the signatory of the message and indicates the person's 12 approval of the information contained in the transmission.

(u) "Electronic transmission" means the transmission of an electronic
 prescription, formatted as an electronic data file, from a prescriber's
 electronic prescription application to a pharmacy's computer, where the
 data file is imported into the pharmacy prescription application.

(v) "Electronically prepared prescription" means a prescription that isgenerated using an electronic prescription application.

19 (w) "Facsimile transmission" or "fax transmission" means the 20 transmission of a digital image of a prescription from the prescriber or the 21 prescriber's agent to the pharmacy. "Facsimile transmission" includes, but 22 is not limited to, transmission of a written prescription between the 23 prescriber's fax machine and the pharmacy's fax machine; transmission of 24 an electronically prepared prescription from the prescriber's electronic 25 prescription application to the pharmacy's fax machine, computer or printer; or transmission of an electronically prepared prescription from the 26 27 prescriber's fax machine to the pharmacy's fax machine, computer or 28 printer.

(x) "Intermediary" means any technology system that receives and
 transmits an electronic prescription between the prescriber and the
 pharmacy.

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(y) "Isomer" means all enantiomers and diastereomers.

33 "Manufacture" means the production, propagation, (z) 34 compounding, conversion or processing of a controlled substance either 35 directly or indirectly or by extraction from substances of natural origin or 36 independently by means of chemical synthesis or by a combination of 37 extraction and chemical synthesis and includes any packaging or 38 repackaging of the substance or labeling or relabeling of its container, 39 except that this term does not include the preparation or compounding of a 40 controlled substance by an individual for the individual's own lawful use 41 or the preparation, compounding, packaging or labeling of a controlled 42 substance:

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(1) By a practitioner or the practitioner's agent pursuant to a lawful

order of a practitioner as an incident to the practitioner's administering or
 dispensing of a controlled substance in the course of the practitioner's
 professional practice; or

4 (2) by a practitioner or by the practitioner's authorized agent under 5 such practitioner's supervision for the purpose of or as an incident to 6 research, teaching or chemical analysis or by a pharmacist or medical care 7 facility as an incident to dispensing of a controlled substance.

8 (aa) "Marijuana" means all parts of all varieties of the plant Cannabis 9 whether growing or not, the seeds thereof, the resin extracted from any 10 part of the plant and every compound, manufacture, salt, derivative, mixture or preparation of the plant, its seeds or resin. It does not include: 11 (1) The mature stalks of the plant, fiber produced from the stalks, oil or 12 cake made from the seeds of the plant, any other compound, manufacture, 13 salt, derivative, mixture or preparation of the mature stalks, except the 14 15 resin extracted therefrom, fiber, oil or cake or the sterilized seed of the 16 plant-which that is incapable of germination; (2) any substance listed in 17 schedules II through V of the uniform controlled substances act; or (3) cannabidiol (other trade name: 2-[(3-methyl-6-(1-methylethenyl)-2-18 19 cyclohexen-1-yl]-5-pentyl-1,3-benzenediol).

20 (bb) "Medical care facility" shall have the meaning ascribed to that 21 term in K.S.A. 65-425, and amendments thereto.

22 "Mid-level practitioner" means a certified nurse-midwife (cc)23 engaging in the independent practice of midwifery under the independent 24 practice of midwifery act, an advanced practice registered nurse issued a 25 license pursuant to K.S.A. 65-1131, and amendments thereto, who has authority to prescribe drugs pursuant to-a written protocol with a-26 27 responsible physician under K.S.A. 65-1130, and amendments thereto, or a 28 physician assistant licensed under the physician assistant licensure act who 29 has authority to prescribe drugs pursuant to a written agreement with a supervising physician under K.S.A. 65-28a08, and amendments thereto. 30

(dd) "Narcotic drug" means any of the following whether produced
 directly or indirectly by extraction from substances of vegetable 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 *that* is chemically equivalent or identical with any of the substances
referred to in paragraph (1) but not including the isoquinoline alkaloids of
opium;

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(3) opium poppy and poppy straw; *or* 

42 (4) coca leaves and any salt, compound, derivative or preparation of 43 coca leaves, and any salt, compound, isomer, derivative or preparation thereof which that is chemically equivalent or identical with any of these
 substances, but not including decocainized coca leaves or extractions of
 coca leaves which that do not contain cocaine or ecgonine.

4 (ee) "Opiate" means any substance having an addiction-forming or 5 addiction-sustaining liability similar to morphine or being capable of 6 conversion into a drug having addiction-forming or addiction-sustaining 7 liability. It does not include, unless specifically designated as controlled 8 under K.S.A. 65-4102, and amendments thereto, the dextrorotatory isomer 9 of 3-methoxy-n-methylmorphinan and its salts (dextromethorphan). It does 10 include its racemic and levorotatory forms.

11 (ff) "Opium poppy" means the plant of the species Papaver 12 somniferum l. except its seeds.

(gg) "Person" means an individual, corporation, government, or
 governmental subdivision or agency, business trust, estate, trust,
 partnership or association or any other legal entity.

(hh) "Pharmacist" means any natural person licensed under K.S.A.
65-1625 et seq., and amendments thereto, to practice pharmacy.

18 (ii) "Pharmacist intern" means: (1) A student currently enrolled in an 19 accredited pharmacy program; (2) a graduate of an accredited pharmacy 20 program serving such person's internship; or (3) a graduate of a pharmacy 21 program located outside of the United States-which *that* is not accredited, 22 and who had successfully passed equivalency examinations approved by 23 the board.

(jj) "Pharmacy prescription application" means software that is used
to process prescription information, is installed on a pharmacy's computers
and servers, and is controlled by the pharmacy.

(kk) "Poppy straw" means all parts, except the seeds, of the opiumpoppy, after mowing.

(II) "Practitioner" means a person licensed to practice medicine and
surgery, dentist, podiatrist, veterinarian, optometrist, or scientific
investigator or other person authorized by law to use a controlled
substance in teaching or chemical analysis or to conduct research with
respect to a controlled substance.

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(mm) "Prescriber" means a practitioner or a mid-level practitioner.

(nn) "Production" includes the manufacture, planting, cultivation,growing or harvesting of a controlled substance.

(oo) "Readily retrievable" means that records kept by automatic data processing applications or other electronic or mechanized recordkeeping systems can be separated out from all other records within a reasonable time not to exceed 48 hours of a request from the board or other authorized agent or that hard-copy records are kept on which certain items are asterisked, redlined or in some other manner visually identifiable apart from other items appearing on the records.

## HB 2066

1 (pp) "Ultimate user" means a person who lawfully possesses a 2 controlled substance for such person's own use or for the use of a member 3 of such person's household or for administering to an animal owned by 4 such person or by a member of such person's household.

5 Sec. 4. K.S.A. 65-1130 and 65-4101 and K.S.A. 2017 Supp. 65-1113, 6 as amended by section 2 of chapter 42 of the 2018 Session Laws of Kansas 7 are hereby repealed.

8 Sec. 5. This act shall take effect and be in force from and after July 1,
9 2020, and its publication in the statute book.