1	STATE OF OKLAHOMA
2	1st Session of the 57th Legislature (2019)
3	SENATE BILL NO. 863 By: Allen
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6	AS INTRODUCED
7	An Act relating to industrial hemp; creating the
8	Industrial Hemp Production Act; providing short title; defining terms; requiring the Oklahoma
9	Department of Agriculture, Food, and Forestry to develop a plan to regulate and license industrial
10	hemp production; requiring the Department to consult with state agencies; requiring the Department to
11	submit a plan before a certain date; requiring resubmission of a plan under certain circumstances;
12	requiring the Department to promulgate rules and establish a fee; creating revolving fund; authorizing
13	expenditures of funds under certain conditions; amending 63 O.S. 2011, Section 2-101, as last amended
14	by Section 11, Chapter 64, O.S.L. 2018 (63 O.S. Supp. 2018, Section 2-101), which relates to definitions;
15	modifying definition; repealing Sections 1 through 10, Chapter 64, O.S.L. 2018 (63 O.S. Supp. 2018, Sections 2 401 through 2 410), which relate to the
16	Sections 3-401 through 3-410), which relate to the Oklahoma Industrial Hemp Agricultural Pilot Program, short title, definitions, licensee authorization,
17	criminal liability exemption, licensee application, promulgation of rules, certified seed program,
18	harvest report, inspection and sampling, license denial, feasibility study and Oklahoma Industrial
19	Hemp Agricultural Pilot Program Fund; providing for codification; and providing an effective date.
20	coullication, and providing an effective date.
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22	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:
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1 SECTION 1. NEW LAW A new section of law to be codified 2 in the Oklahoma Statutes as Section 3-420 of Title 2, unless there 3 is created a duplication in numbering, reads as follows: 4 This act shall be known and may be cited as the "Industrial Hemp 5 Production Act". 6 SECTION 2. NEW LAW A new section of law to be codified 7 in the Oklahoma Statutes as Section 3-421 of Title 2, unless there 8 is created a duplication in numbering, reads as follows: 9 As used in this act: 10 "Department" means the Oklahoma Department of Agriculture, 1. 11 Food, and Forestry; and 12 2. "Industrial Hemp Production License" or "License" means 13 authorization by the Department to grow and cultivate industrial 14 hemp. 15 A new section of law to be codified SECTION 3. NEW LAW 16 in the Oklahoma Statutes as Section 3-422 of Title 2, unless there 17 is created a duplication in numbering, reads as follows: 18 The Oklahoma Department of Agriculture, Food, and Forestry Α. 19 shall develop a plan to license and regulate industrial hemp 20 production. 21 The Department shall consult with the Office of the Attorney в. 22 General and the Office of the Governor regarding the development of 23 the plan. 24 _ _

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C. The Department shall submit the plan to the United States
 Secretary of Agriculture for approval. Submission of the plan shall
 occur no later than January 1, 2020.

D. If the United States Secretary of Agriculture disapproves of
the plan, the Department shall consult with the Office of the
Attorney General and the Office of the Governor and submit a revised
plan. The revised plan shall be submitted within ninety (90) days
of receipt of the notice of disapproval.

9 SECTION 4. NEW LAW A new section of law to be codified 10 in the Oklahoma Statutes as Section 3-423 of Title 2, unless there 11 is created a duplication in numbering, reads as follows:

12 A. Upon the receipt of approval from the United States 13 Secretary of Agriculture for the plan to license and regulate 14 industrial hemp production, the Oklahoma Department of Agriculture, 15 Food, and Forestry shall promulgate rules to implement the plan and 16 issue licenses.

B. The Department shall establish a fee for an industrial hemp
 license that shall cost no more than One Dollar and twenty-five
 cents (\$1.25) per acre of land and no more than nine cents (\$0.09)
 per square foot of greenhouse area.

SECTION 5. NEW LAW A new section of law to be codified in the Oklahoma Statutes as Section 3-424 of Title 2, unless there is created a duplication in numbering, reads as follows:

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1 There is hereby created in the State Treasury a revolving fund 2 for the State Board of Agriculture to be designated the "Oklahoma 3 Industrial Hemp Production Fund". The fund shall be a continuing 4 fund, not subject to fiscal year limitations and shall consist of 5 all monies received by the State Board of Agriculture from fees 6 received and collected pursuant to the Oklahoma Industrial Hemp 7 Production Act, donations, grants, contributions and gifts from any 8 public or private source. The Board may expend funds for the 9 purposes set forth in the Oklahoma Industrial Hemp Production Act. 10 Expenditures from the fund shall be made upon warrants issued by the 11 State Treasurer against claims filed as prescribed by law with the 12 Director of the Office of Management and Enterprise Services for 13 approval and payment. 14 SECTION 6. AMENDATORY 63 O.S. 2011, Section 2-101, as 15 last amended by Section 11, Chapter 64, O.S.L. 2018 (63 O.S. Supp. 16 2018, Section 2-101), is amended to read as follows: 17 Section 2-101. As used in the Uniform Controlled Dangerous 18 Substances Act: 19 "Administer" means the direct application of a controlled 1. 20 dangerous substance, whether by injection, inhalation, ingestion or 21 any other means, to the body of a patient, animal or research 22 subject by:

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- a. a practitioner (or, in the presence of the practitioner, by the authorized agent of the practitioner), or
- 4 5

b. the patient or research subject at the direction and in the presence of the practitioner;

6 2. "Agent" means a peace officer appointed by and who acts on 7 behalf of the Director of the Oklahoma State Bureau of Narcotics and 8 Dangerous Drugs Control or an authorized person who acts on behalf 9 of or at the direction of a person who manufactures, distributes, 10 dispenses, prescribes, administers or uses for scientific purposes 11 controlled dangerous substances but does not include a common or 12 contract carrier, public warehouser or employee thereof, or a person 13 required to register under the Uniform Controlled Dangerous 14 Substances Act;

¹⁵ 3. "Board" means the Advisory Board to the Director of the
 ¹⁶ Oklahoma State Bureau of Narcotics and Dangerous Drugs Control;

¹⁷ 4. "Bureau" means the Oklahoma State Bureau of Narcotics and
 ¹⁸ Dangerous Drugs Control;

¹⁹ 5. "Coca leaves" includes cocaine and any compound, ²⁰ manufacture, salt, derivative, mixture or preparation of coca ²¹ leaves, except derivatives of coca leaves which do not contain ²² cocaine or ecgonine;

6. "Commissioner" or "Director" means the Director of the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control;

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¹ 7. "Control" means to add, remove or change the placement of a ² drug, substance or immediate precursor under the Uniform Controlled ³ Dangerous Substances Act;

8. "Controlled dangerous substance" means a drug, substance or
immediate precursor in Schedules I through V of the Uniform
Controlled Dangerous Substances Act or any drug, substance or
immediate precursor listed either temporarily or permanently as a
federally controlled substance. Any conflict between state and
federal law with regard to the particular schedule in which a
substance is listed shall be resolved in favor of state law;

9. "Counterfeit substance" means a controlled substance which, or the container or labeling of which without authorization, bears the trademark, trade name or other identifying marks, 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;

17 10. "Deliver" or "delivery" means the actual, constructive or 18 attempted transfer from one person to another of a controlled 19 dangerous substance or drug paraphernalia, whether or not there is 20 an agency relationship;

21 11. "Dispense" means to deliver a controlled dangerous 22 substance to an ultimate user or human research subject by or 23 pursuant to the lawful order of a practitioner, including the 24 prescribing, administering, packaging, labeling or compounding

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¹ necessary to prepare the substance for such distribution.
² "Dispenser" is a practitioner who delivers a controlled dangerous
³ substance to an ultimate user or human research subject;

⁴ 12. "Distribute" means to deliver other than by administering
 ⁵ or dispensing a controlled dangerous substance;

13. "Distributor" means a commercial entity engaged in the
distribution or reverse distribution of narcotics and dangerous
drugs and who complies with all regulations promulgated by the
federal Drug Enforcement Administration and the Oklahoma State
Bureau of Narcotics and Dangerous Drugs Control;

- 11 14. "Drug" means articles:
- a. recognized in the official United States
 Pharmacopoeia, official Homeopathic Pharmacopoeia of
 the United States, or official National Formulary, or
 any supplement to any of them,
- b. intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in man or other animals,
- 19c. other than food, intended to affect the structure or20any function of the body of man or other animals, and
- 21 d. intended for use as a component of any article
 22 specified in this paragraph;

²³ provided, however, the term "drug" does not include devices or their ²⁴ components, parts or accessories;

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1 15. "Drug-dependent person" means a person who is using a 2 controlled dangerous substance and who is in a state of psychic or 3 physical dependence, or both, arising from administration of that 4 controlled dangerous substance on a continuous basis. Drug 5 dependence is characterized by behavioral and other responses which 6 include a strong compulsion to take the substance on a continuous 7 basis in order to experience its psychic effects, or to avoid the 8 discomfort of its absence;

9 16. "Home care agency" means any sole proprietorship, 10 partnership, association, corporation, or other organization which 11 administers, offers, or provides home care services, for a fee or 12 pursuant to a contract for such services, to clients in their place 13 of residence;

14 17. "Home care services" means skilled or personal care 15 services provided to clients in their place of residence for a fee; 16 18. "Hospice" means a centrally administered, nonprofit or 17 profit, medically directed, nurse-coordinated program which provides 18 a continuum of home and inpatient care for the terminally ill 19 patient and the patient's family. Such term shall also include a 20 centrally administered, nonprofit or profit, medically directed, 21 nurse-coordinated program if such program is licensed pursuant to 22 the provisions of this act. A hospice program offers palliative and 23 supportive care to meet the special needs arising out of the 24 physical, emotional and spiritual stresses which are experienced _ _

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¹ during the final stages of illness and during dying and bereavement.
² This care is available twenty-four (24) hours a day, seven (7) days
³ a week, and is provided on the basis of need, regardless of ability
⁴ to pay. "Class A" Hospice refers to Medicare certified hospices.
⁵ "Class B" refers to all other providers of hospice services;

6 19. "Imitation controlled substance" means a substance that is 7 not a controlled dangerous substance, which by dosage unit 8 appearance, color, shape, size, markings or by representations made, 9 would lead a reasonable person to believe that the substance is a 10 controlled dangerous substance. In the event the appearance of the 11 dosage unit is not reasonably sufficient to establish that the 12 substance is an "imitation controlled substance", the court or 13 authority concerned should consider, in addition to all other 14 factors, the following factors as related to "representations made" 15 in determining whether the substance is an "imitation controlled 16 substance":

statements made by an owner or by any other person in a. 18 control of the substance concerning the nature of the 19 substance, or its use or effect, 20 b. statements made to the recipient that the substance 21 may be resold for inordinate profit, 22 whether the substance is packaged in a manner normally с. 23 used for illicit controlled substances, 24

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- 1 d. evasive tactics or actions utilized by the owner or 2 person in control of the substance to avoid detection 3 by law enforcement authorities,
- e. prior convictions, if any, of an owner, or any other
 person in control of the object, under state or
 federal law related to controlled substances or fraud,
 and
- 8 f. the proximity of the substances to controlled 9 dangerous substances;

10 20. "Immediate precursor" means a substance which the Director 11 has found to be and by regulation designates as being the principal 12 compound commonly used or produced primarily for use, and which is 13 an immediate chemical intermediary used, or likely to be used, in 14 the manufacture of a controlled dangerous substance, the control of 15 which is necessary to prevent, curtail or limit such manufacture; 16 21. "Laboratory" means a laboratory approved by the Director as 17 proper to be entrusted with the custody of controlled dangerous

¹⁸ substances and the use of controlled dangerous substances for ¹⁹ scientific and medical purposes and for purposes of instruction;

20 22. "Manufacture" means the production, preparation,
 21 propagation, compounding or processing of a controlled dangerous
 22 substance, either directly or indirectly by extraction from
 23 substances of natural or synthetic origin, or independently by means
 24 of chemical synthesis or by a combination of extraction and chemical

1 synthesis. "Manufacturer" includes any person who packages, 2 repackages or labels any container of any controlled dangerous 3 substance, except practitioners who dispense or compound 4 prescription orders for delivery to the ultimate consumer; 5 "Marijuana" means all parts of the plant Cannabis sativa 23. 6 L., whether growing or not; the seeds thereof; the resin extracted 7 from any part of such plant; and every compound, manufacture, salt, 8 derivative, mixture or preparation of such plant, its seeds or 9 resin, but shall not include: 10 the mature stalks of such plant or fiber produced from a. 11 such stalks, 12 b. oil or cake made from the seeds of such plant, 13 including cannabidiol derived from the seeds of the 14 marijuana plant, 15 any other compound, manufacture, salt, derivative, с. 16 mixture or preparation of such mature stalks (except 17 the resin extracted therefrom), including cannabidiol 18 derived from mature stalks, fiber, oil or cake, 19 the sterilized seed of such plant which is incapable d. 20 of germination, 21 for any person participating in a clinical trial to e. 22 administer cannabidiol for the treatment of severe 23 forms of epilepsy pursuant to Section 2-802 of this 24 title, a drug or substance approved by the federal _ _

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Food and Drug Administration for use by those participants,

- 3 f. for any person or the parents, legal quardians or 4 caretakers of the person who have received a written 5 certification from a physician licensed in this state 6 that the person has been diagnosed by a physician as 7 having Lennox-Gastaut Syndrome, Dravet Syndrome, also 8 known as Severe Myoclonic Epilepsy of Infancy, or any 9 other severe form of epilepsy that is not adequately 10 treated by traditional medical therapies, spasticity 11 due to multiple sclerosis or due to paraplegia, 12 intractable nausea and vomiting, appetite stimulation 13 with chronic wasting diseases, the substance 14 cannabidiol, a nonpsychoactive cannabinoid, found in 15 the plant Cannabis sativa L. or any other preparation 16 thereof, that has a tetrahydrocannabinol concentration 17 of not more than three-tenths of one percent (0.3%)18 and that is delivered to the patient in the form of a 19 liquid,
- 20 g. any federal Food and Drug Administration-approved 21 cannabidiol drug or substance, or
- h. industrial hemp, from the plant Cannabis sativa L. and
 any part of such plant, whether growing or not, with a
 delta-9 tetrahydrocannabinol concentration of not more

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than three-tenths of one percent (0.3%) on a dry weight basis which shall only be grown pursuant to the Oklahoma Industrial Hemp Agricultural Pilot Program and may be shipped to Oklahoma pursuant to the provisions of subparagraph e or f of this paragraph;

6 24. "Medical purpose" means an intention to utilize a 7 controlled dangerous substance for physical or mental treatment, for 8 diagnosis, or for the prevention of a disease condition not in 9 violation of any state or federal law and not for the purpose of 10 satisfying physiological or psychological dependence or other abuse;

11 "Mid-level practitioner" means an advanced practice nurse 25. 12 as defined and within parameters specified in Section 567.3a of 13 Title 59 of the Oklahoma Statutes, or a certified animal euthanasia 14 technician as defined in Section 698.2 of Title 59 of the Oklahoma 15 Statutes, or an animal control officer registered by the Oklahoma 16 State Bureau of Narcotics and Dangerous Drugs Control under 17 subsection B of Section 2-301 of this title within the parameters of 18 such officer's duty under Sections 501 through 508 of Title 4 of the 19 Oklahoma Statutes;

20 26. "Narcotic drug" means any of the following, whether
21 produced directly or indirectly by extraction from substances of
22 vegetable origin, or independently by means of chemical synthesis,
23 or by a combination of extraction and chemical synthesis:

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a. opium, coca leaves and opiates,

1	b. a	compound, manufacture, salt, derivative or
2	pre	eparation of opium, coca leaves or opiates,
3	C. CO	caine, its salts, optical and geometric isomers, and
4	sa	lts of isomers,
5	d. ecc	gonine, its derivatives, their salts, isomers and
6	sa	lts of isomers, and
7	e.a:	substance, and any compound, manufacture, salt,
8	de:	rivative or preparation thereof, which is chemically
9	ide	entical with any of the substances referred to in
10	sul	oparagraphs a through d of this paragraph, except
11	tha	at the words "narcotic drug" as used in Section 2-
12	10:	l et seq. of this title shall not include
13	deo	cocainized coca leaves or extracts of coca leaves,
14	wh	ich extracts do not contain cocaine or ecgonine;
15	27. "Opiate"	" means any substance having an addiction-forming or
16	addiction-sustain	ning liability similar to morphine or being capable
17	of conversion in	to a drug having such addiction-forming or
18	addiction-sustain	ning liability. It does not include, unless
19	specifically des	ignated as controlled under the Uniform Controlled
20	Dangerous Substan	nces Act, the dextrorotatory isomer of 3-methoxy-n-
21	methyl-morphinan	and its salts (dextromethorphan). It does include
22	its racemic and i	levorotatory forms;

23 28. "Opium poppy" means the plant of the species Papaver
24 somniferum L., except the seeds thereof;

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1 29. "Peace officer" means a police officer, sheriff, deputy 2 sheriff, district attorney's investigator, investigator from the 3 Office of the Attorney General, or any other person elected or 4 appointed by law to enforce any of the criminal laws of this state 5 or of the United States;

⁶ 30. "Person" means an individual, corporation, government or ⁷ governmental subdivision or agency, business trust, estate, trust, ⁸ partnership or association, or any other legal entity;

9 31. "Poppy straw" means all parts, except the seeds, of the 10 opium poppy, after mowing;

11 32. "Practitioner" means:

a. (1) a medical doctor or osteopathic physician,

- (2) a dentist,
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 (3) a podiatrist,
- 15 (4) an optometrist,
 - (5) a veterinarian,
- (6) a physician assistant under the supervision of a
 licensed medical doctor or osteopathic physician,
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 (7) a scientific investigator, or
- 20 (8) any other person,

21 licensed, registered or otherwise permitted to 22 prescribe, distribute, dispense, conduct research with 23 respect to, use for scientific purposes or administer

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1 a controlled dangerous substance in the course of 2 professional practice or research in this state, or 3 b. a pharmacy, hospital, laboratory or other institution 4 licensed, registered or otherwise permitted to 5 distribute, dispense, conduct research with respect 6 to, use for scientific purposes or administer a 7 controlled dangerous substance in the course of 8 professional practice or research in this state; 9 "Production" includes the manufacture, planting, 33. 10 cultivation, growing or harvesting of a controlled dangerous 11 substance; 12 34. "State" means the State of Oklahoma or any other state of 13 the United States; 14 "Ultimate user" means a person who lawfully possesses a 35. 15 controlled dangerous substance for the person's own use or for the 16 use of a member of the person's household or for administration to 17 an animal owned by the person or by a member of the person's 18 household;

¹⁹ 36. "Drug paraphernalia" means all equipment, products and ²⁰ materials of any kind which are used, intended for use, or fashioned ²¹ specifically for use in planting, propagating, cultivating, growing, ²² harvesting, manufacturing, compounding, converting, producing, ²³ processing, preparing, testing, analyzing, packaging, repackaging, ²⁴ storing, containing, concealing, injecting, ingesting, inhaling or

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1 otherwise introducing into the human body, a controlled dangerous 2 substance in violation of the Uniform Controlled Dangerous 3 Substances Act including, but not limited to: 4 a. kits used, intended for use, or fashioned specifically 5 for use in planting, propagating, cultivating, growing 6 or harvesting of any species of plant which is a 7 controlled dangerous substance or from which a 8 controlled dangerous substance can be derived, 9 b. kits used, intended for use, or fashioned specifically 10 for use in manufacturing, compounding, converting, 11 producing, processing or preparing controlled 12 dangerous substances, 13 с. isomerization devices used, intended for use, or 14 fashioned specifically for use in increasing the 15 potency of any species of plant which is a controlled 16 dangerous substance, 17 d. testing equipment used, intended for use, or fashioned 18 specifically for use in identifying, or in analyzing 19 the strength, effectiveness or purity of controlled 20 dangerous substances, 21 scales and balances used, intended for use, or e. 22 fashioned specifically for use in weighing or 23 measuring controlled dangerous substances, 24 _ _

- 1 f. diluents and adulterants, such as quinine 2 hydrochloride, mannitol, mannite, dextrose and 3 lactose, used, intended for use, or fashioned 4 specifically for use in cutting controlled dangerous 5 substances,
 - g. separation gins and sifters used, intended for use, or fashioned specifically for use in removing twigs and seeds from, or in otherwise cleaning or refining, marijuana,
- h. blenders, bowls, containers, spoons and mixing devices used, intended for use, or fashioned specifically for use in compounding controlled dangerous substances,
- i. capsules, balloons, envelopes and other containers
 used, intended for use, or fashioned specifically for
 use in packaging small quantities of controlled
 dangerous substances,
- j. containers and other objects used, intended for use, or fashioned specifically for use in parenterally injecting controlled dangerous substances into the human body,
- k. hypodermic syringes, needles and other objects used, intended for use, or fashioned specifically for use in parenterally injecting controlled dangerous substances into the human body,

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- 1 1. objects used, intended for use, or fashioned 2 specifically for use in ingesting, inhaling or 3 otherwise introducing marijuana, cocaine, hashish or 4 hashish oil into the human body, such as: 5 metal, wooden, acrylic, glass, stone, plastic or (1) 6 ceramic pipes with or without screens, permanent 7 screens, hashish heads or punctured metal bowls, 8 (2) water pipes, 9 carburction tubes and devices, (3) 10 smoking and carburetion masks, (4) 11 roach clips, meaning objects used to hold burning (5) 12 material, such as a marijuana cigarette, that has 13 become too small or too short to be held in the 14 hand, 15 miniature cocaine spoons and cocaine vials, (6) 16 (7) chamber pipes, 17 (8) carburetor pipes, 18 (9) electric pipes, 19 air-driven pipes, (10)20 (11) chillums, 21 (12) bongs, or 22 (13) ice pipes or chillers, 23 all hidden or novelty pipes, and m. 24 _ _
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1 any pipe that has a tobacco bowl or chamber of less n. 2 than one-half (1/2) inch in diameter in which there is 3 any detectable residue of any controlled dangerous 4 substance as defined in this section or any other 5 substances not legal for possession or use; 6 provided, however, the term "drug paraphernalia" shall not include 7 separation gins intended for use in preparing tea or spice, clamps 8 used for constructing electrical equipment, water pipes designed for 9 ornamentation in which no detectable amount of an illegal substance 10 is found or pipes designed and used solely for smoking tobacco, 11 traditional pipes of an American Indian tribal religious ceremony, 12 or antique pipes that are thirty (30) years of age or older; 13 37. "Synthetic controlled substance" means a substance: a. 14 the chemical structure of which is substantially (1)15 similar to the chemical structure of a controlled 16 dangerous substance in Schedule I or II, 17 (2) which has a stimulant, depressant, or 18 hallucinogenic effect on the central nervous 19 system that is substantially similar to or 20 greater than the stimulant, depressant or 21 hallucinogenic effect on the central nervous 22 system of a controlled dangerous substance in 23 Schedule I or II, or 24

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1		(3)	with respect to a particular person, which such
2			person represents or intends to have a stimulant,
3			depressant, or hallucinogenic effect on the
4			central nervous system that is substantially
5			similar to or greater than the stimulant,
6			depressant, or hallucinogenic effect on the
7			central nervous system of a controlled dangerous
8			substance in Schedule I or II.
9	b.	The	designation of gamma butyrolactone or any other
10		chem	nical as a precursor, pursuant to Section 2-322 of
11		this	title, does not preclude a finding pursuant to
12		subp	paragraph a of this paragraph that the chemical is
13		a sy	nthetic controlled substance.
14	C.	"Syn	thetic controlled substance" does not include:
15		(1)	a controlled dangerous substance,
16		(2)	any substance for which there is an approved new
17			drug application,
18		(3)	with respect to a particular person any
19			substance, if an exemption is in effect for
20			investigational use, for that person under the
21			provisions of Section 505 of the Federal Food,
22			Drug and Cosmetic Act, Title 21 of the United
23			States Code, Section 355, to the extent conduct
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- 1 with respect to such substance is pursuant to
 2 such exemption, or
 - (4) any substance to the extent not intended for human consumption before such an exemption takes effect with respect to that substance.
- d. Prima facie evidence that a substance containing
 7 salvia divinorum has been enhanced, concentrated or
 8 chemically or physically altered shall give rise to a
 9 rebuttable presumption that the substance is a
 10 synthetic controlled substance;

¹¹ 38. "Tetrahydrocannabinols" means all substances that have been ¹² chemically synthesized to emulate the tetrahydrocannabinols of ¹³ marijuana;

¹⁴ 39. "Isomer" means the optical isomer, except as used in ¹⁵ subsections C and F of Section 2-204 of this title and paragraph 4 ¹⁶ of subsection A of Section 2-206 of this title. As used in ¹⁷ subsections C and F of Section 2-204 of this title, "isomer" means ¹⁸ the optical, positional or geometric isomer. As used in paragraph 4 ¹⁹ of subsection A of Section 2-206 of this title, the term "isomer" ²⁰ means the optical or geometric isomer;

40. "Hazardous materials" means materials, whether solid, liquid or gas, which are toxic to human, animal, aquatic or plant life, and the disposal of which materials is controlled by state or federal guidelines; and

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1	41. "Anhydrous ammonia" means any substance that exhibits
2	cryogenic evaporative behavior and tests positive for ammonia.
3	SECTION 7. REPEALER Sections 1 through 10, Chapter 64,
4	O.S.L. 2018 (2 O.S. Supp. 2018, Section 3-401 through 3-410), are
5	hereby repealed.
6	SECTION 8. This act shall become effective November 1, 2019.
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