

1 ENGROSSED HOUSE  
2 BILL NO. 2913

By: Dollens, Echols and  
Rosecrants of the House

3 and

4 Paxton of the Senate  
5  
6

7 An Act relating to industrial hemp; creating the  
8 Oklahoma Industrial Hemp Agricultural Pilot Program;  
9 defining terms; authorizing a licensee to engage in  
10 the growth and cultivation of industrial hemp for  
11 certain purposes; limiting liability; requiring  
12 application to the Oklahoma Department of  
13 Agriculture, Food, and Forestry; providing  
14 application content requirements; providing certain  
15 acknowledgements and agreements upon application  
16 submission; requiring certain application fee;  
17 directing the Department to establish certain fee  
18 schedule; providing length of valid license;  
19 providing license renewal process; requiring  
20 activities be done with a valid license; requiring  
21 certain plants not harvested or destroyed be  
22 declared; requiring submission of information for  
23 certain land alterations or changes to information;  
24 directing the Department to promulgate rules;  
directing the Department to establish a Certified  
Seed Program; allowing certain varieties of  
industrial hemp be approved; requiring the Department  
to maintain a list of certified seeds; requiring a  
harvest report; providing for routine inspection and  
sampling of plants of the licensee; providing for  
additional inspection and sampling under certain  
conditions and circumstances; providing inspection  
procedure requirements; requiring the licensee to pay  
for inspection and lab analysis with exception;  
directing the Department to promulgate rules;  
authorizing denial, revocation or suspension of  
license under certain circumstances; prohibiting  
penalty for certain sample testing levels; directing  
the Department to study certain funding  
possibilities; creating revolving fund; authorizing  
expenditures of funds under certain conditions;

1 amending 63 O.S. 2011, Section 2-101, as last amended  
2 by Section 1, Chapter 43, O.S.L. 2017 (63 O.S. Supp.  
3 2017, Section 2-101), which relates to the Uniform  
4 Controlled Dangerous Substances Act; amending  
5 definition; providing for codification; and declaring  
6 an emergency.

7 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

8 SECTION 1. NEW LAW A new section of law to be codified  
9 in the Oklahoma Statutes as Section 3-401 of Title 2, unless there  
10 is created a duplication in numbering, reads as follows:

11 This act shall be known and may be cited as the "Oklahoma  
12 Industrial Hemp Agricultural Pilot Program".

13 SECTION 2. NEW LAW A new section of law to be codified  
14 in the Oklahoma Statutes as Section 3-402 of Title 2, unless there  
15 is created a duplication in numbering, reads as follows:

16 As used in the Oklahoma Industrial Hemp Agricultural Pilot  
17 Program:

18 1. "Certified seed" means industrial hemp seed that has been  
19 certified by the Oklahoma Department of Agriculture, Food, and  
20 Forestry as having no more than three-tenths of one percent (0.3%)  
21 delta-9 tetrahydrocannabinol concentration on a dry-weight basis;

22 2. "Department" means the Oklahoma Department of Agriculture,  
23 Food, and Forestry;

24 3. "Industrial hemp" means the plant Cannabis sativa L. and any  
part of the plant, whether growing or not, with a delta-9

1 tetrahydrocannabinol concentration of not more than three-tenths of  
2 one percent (0.3%) on a dry-weight basis;

3 4. "Licensee" means a university or an institution of higher  
4 education located in Oklahoma which holds a valid Industrial Hemp  
5 License to grow industrial hemp under the Oklahoma Industrial Hemp  
6 Agricultural Pilot Program. Nothing in the Oklahoma Industrial Hemp  
7 Agricultural Pilot Program shall prevent the licensee from adopting  
8 policies and procedures to subcontract with persons or other legal  
9 entities to carry out the purposes of the program; provided, that  
10 the Oklahoma Department of Agriculture, Food, and Forestry shall  
11 ensure subcontractors comply with the program requirements; and

12 5. "Industrial Hemp License" or "License" means authorization  
13 by the Department for any university or an institution of higher  
14 education in Oklahoma to grow and cultivate industrial hemp on a  
15 registered land area for research and development purposes as part  
16 of the Oklahoma Industrial Hemp Agricultural Pilot Program.

17 SECTION 3. NEW LAW A new section of law to be codified  
18 in the Oklahoma Statutes as Section 3-403 of Title 2, unless there  
19 is created a duplication in numbering, reads as follows:

20 A. A licensee is authorized to:

21 1. Engage in the growth and cultivation of industrial hemp from  
22 certified seeds for agricultural plant research and development  
23 purposes; and

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1           2. Engage in the growth and cultivation of industrial hemp from  
2 certified seeds for marketing development purposes.

3           B. The activities performed under the Oklahoma Industrial Hemp  
4 Agricultural Pilot Program shall not subject the persons  
5 participating in the program to criminal liability under the Uniform  
6 Controlled Dangerous Substances Act. The exemption from criminal  
7 liability provided for in this subsection is a limited exemption  
8 that shall be strictly construed and shall not apply to an activity  
9 that is not expressly permitted under the Oklahoma Industrial Hemp  
10 Agricultural Pilot Program.

11           SECTION 4.       NEW LAW       A new section of law to be codified  
12 in the Oklahoma Statutes as Section 3-404 of Title 2, unless there  
13 is created a duplication in numbering, reads as follows:

14           A. A university or an institution of higher education located  
15 in Oklahoma wishing to engage in industrial hemp growth and  
16 cultivation authorized under the Oklahoma Industrial Hemp  
17 Agricultural Pilot Program shall apply to the Oklahoma Department of  
18 Agriculture, Food, and Forestry for a license prior to planting the  
19 industrial hemp.

20           1. The application shall include:

- 21           a. the name and address of the university or an  
22               institution of higher education,
- 23           b. the legal description, global positioning system  
24               location, and map of the land area on which the

1 licensee will engage in industrial hemp growth and  
2 cultivation operations,

3 c. a statement of intended end use, and

4 d. a statement that the licensee intends to plant only  
5 certified seeds.

6 2. By submitting an application, the licensee acknowledges and  
7 agrees that:

8 a. information provided to the Department may be provided  
9 to law enforcement agencies,

10 b. the licensee and any entities contracting with the  
11 licensee shall allow and fully cooperate with any  
12 inspection and sampling that the Department deems  
13 necessary,

14 c. the licensee will submit all required reports by the  
15 applicable due dates specified by the Department, and

16 d. the licensee has the legal right to cultivate  
17 industrial hemp from certified seeds on the registered  
18 land area and shall grant the Department access for  
19 inspection and sampling.

20 B. The Department shall collect a nonrefundable fee from the  
21 licensee at the time of application. The Department shall set a fee  
22 schedule based on the size and use of the land area on which the  
23 licensee will conduct industrial hemp growing or cultivation  
24 operations and shall set the fee at a level sufficient to generate

1 the amount of monies necessary to cover the Department's direct  
2 costs in implementing the Oklahoma Industrial Hemp Agricultural  
3 Pilot Program. Denied applications for a license may be resubmitted  
4 within a twelve-month period. The Department may waive the fee for  
5 resubmitted applications.

6 C. A license issued pursuant to this section is valid for one  
7 (1) year. In order to continue engaging in industrial hemp growth  
8 and cultivation operations in Oklahoma, the licensee must annually  
9 apply for a license in accordance with subsection A of this section.  
10 The Department may set a separate fee schedule for renewal of  
11 existing licenses in good standing.

12 D. All industrial hemp plant material shall be planted, grown  
13 and harvested under a valid license. Any plant material that is not  
14 harvested in the license period in which it was planted or volunteer  
15 plants that are not destroyed must be declared for inclusion in a  
16 subsequent license.

17 E. If the licensee wishes to alter the land area on which the  
18 licensee will conduct industrial hemp growth and cultivation  
19 operations within thirty (30) days of any new license, before  
20 altering the area, the licensee shall submit to the Department an  
21 updated legal description, global positioning system location, and  
22 map specifying the proposed alterations.

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1 F. Each licensee shall report any changes to information  
2 provided in the license application within ten (10) days of such  
3 change to the Department.

4 G. The Department shall promulgate rules necessary to implement  
5 the licensing program and to implement the Oklahoma Industrial Hemp  
6 Agricultural Pilot Program.

7 SECTION 5. NEW LAW A new section of law to be codified  
8 in the Oklahoma Statutes as Section 3-405 of Title 2, unless there  
9 is created a duplication in numbering, reads as follows:

10 A. The Department shall establish a Certified Seed Program to  
11 identify seeds that have been confirmed to produce industrial hemp.  
12 In accordance with all federal state laws and regulations, the  
13 Department may import seeds.

14 B. A variety of industrial hemp may be approved and certified  
15 by the Department if it is tested and confirmed to produce mature  
16 plants with a delta-9 tetrahydrocannabinol concentration of not more  
17 than three-tenths of one percent (0.3%) on a dry-weight basis.

18 C. The Department shall provide and maintain a list of  
19 certified seeds to be used by licensees.

20 SECTION 6. NEW LAW A new section of law to be codified  
21 in the Oklahoma Statutes as Section 3-406 of Title 2, unless there  
22 is created a duplication in numbering, reads as follows:

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1 A. At least thirty (30) days prior to harvest, each licensee  
2 shall file a harvest report on a form approved by the Department  
3 that includes:

4 1. A statement of intended disposition of its industrial hemp  
5 crop;

6 2. The harvest date or dates, location and yield of each  
7 variety cultivated within a registered land area;

8 3. The documented environmental impacts and viability of each  
9 variety; and

10 4. Research data that would assist the Department in future  
11 commercialization of industrial hemp.

12 B. A licensee shall notify the Department immediately of any  
13 changes in a reported harvest date by more than five (5) days.

14 SECTION 7. NEW LAW A new section of law to be codified  
15 in the Oklahoma Statutes as Section 3-407 of Title 2, unless there  
16 is created a duplication in numbering, reads as follows:

17 A. Any plants of the licensee are subject to routine inspection  
18 and sampling to verify that the delta-9 tetrahydrocannabinol  
19 concentration of the plants planted does not exceed three-tenths of  
20 one percent (0.3%) on a dry-weight basis. The Department shall  
21 notify each licensee of the scope of the inspection and the process  
22 by which the inspection will be conducted.



1 B. In addition to any routine inspection and sampling under  
2 subsection A of this section, the Department may inspect and take  
3 samples from any licensee's plants during normal business hours.

4 C. The Department shall make a good-faith attempt to have the  
5 licensee present at the time of inspection and sampling. The  
6 licensee or authorized representative shall provide the Department's  
7 inspector with complete and unrestricted access to all plants, parts  
8 and seeds, whether growing or harvested, and all land, buildings and  
9 other structures used for the growth, cultivation, harvesting or  
10 storage of industrial hemp, and all documents and records pertaining  
11 to the licensee's industrial hemp-growing and cultivation operation.

12 D. The licensee shall pay for any inspection and laboratory  
13 analysis costs that the Department deems necessary within thirty  
14 (30) days of the date of the receipt of an invoice for the costs.  
15 The Department shall waive all inspection or sampling costs if no  
16 inconsistencies or violations are identified during an inspection  
17 that is not part of the regular inspection process.

18 E. The Department shall promulgate rules to establish a process  
19 by which a licensee may contest the procedures, protocols and  
20 results or findings of the inspection.

21 SECTION 8. NEW LAW A new section of law to be codified  
22 in the Oklahoma Statutes as Section 3-408 of Title 2, unless there  
23 is created a duplication in numbering, reads as follows:  
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1       A. The Department may deny, revoke or suspend a license if the  
2 licensee:

3       1. Violates any provision of the Oklahoma Industrial Hemp  
4 Agricultural Pilot Program or rules adopted pursuant to the program;

5       2. Engages in fraud or deception in the procurement of or  
6 attempt to procure a license under this Oklahoma Industrial Hemp  
7 Agricultural Pilot Program or provides false information on a  
8 license application;

9       3. Refuses or fails to cooperate and assist the Department with  
10 the inspection process;

11       4. Refuses or fails to provide any information required or  
12 requested by the Department for purposes of the Oklahoma Industrial  
13 Hemp Agricultural Pilot Program;

14       5. Knowingly provides false, misleading or incorrect  
15 information pertaining to the licensee's cultivation of industrial  
16 hemp to the Department by any means, including information provided  
17 in any application form, report, record or inspection required or  
18 maintained for purposes of the Oklahoma Industrial Hemp Agricultural  
19 Pilot Program;

20       6. Fails to submit any report required by the Oklahoma  
21 Industrial Hemp Agricultural Pilot Program; or

22       7. Fails to pay fees required by the Oklahoma Industrial Hemp  
23 Agricultural Pilot Program.

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1 B. If a sample of a licensee's industrial hemp tests higher  
2 than three-tenths of one percent (0.3%) but less than one percent  
3 (1%) delta-9 tetrahydrocannabinol concentration, the licensee shall  
4 not be subject to any penalty under the Oklahoma Industrial Hemp  
5 Agricultural Pilot Program if the crop is destroyed or utilized on  
6 site in a manner approved of and verified by the Department.

7 SECTION 9. NEW LAW A new section of law to be codified  
8 in the Oklahoma Statutes as Section 3-409 of Title 2, unless there  
9 is created a duplication in numbering, reads as follows:

10 The Department shall study the feasibility of attracting federal  
11 and private funding to implement the Oklahoma Industrial Hemp  
12 Agricultural Pilot Program.

13 SECTION 10. NEW LAW A new section of law to be codified  
14 in the Oklahoma Statutes as Section 3-410 of Title 2, unless there  
15 is created a duplication in numbering, reads as follows:

16 There is hereby created in the State Treasury a revolving fund  
17 for the State Board of Agriculture to be designated the "Oklahoma  
18 Industrial Hemp Agricultural Pilot Program Fund". The fund shall be  
19 a continuing fund, not subject to fiscal year limitations and shall  
20 consist of all monies received by the State Board of Agriculture  
21 from fees received and collected pursuant to the Oklahoma Industrial  
22 Hemp Agricultural Pilot Program, donations, grants, contributions  
23 and gifts from any public or private source. The Board may expend  
24 funds for the purposes set forth in the Oklahoma Industrial Hemp

1 Agricultural Pilot Program. Expenditures from said fund shall be  
2 made upon warrants issued by the State Treasurer against claims  
3 filed as prescribed by law with the Director of the Office of  
4 Management and Enterprise Services for approval and payment.

5 SECTION 11. AMENDATORY 63 O.S. 2011, Section 2-101, as  
6 last amended by Section 1, Chapter 43, O.S.L. 2017 (63 O.S. Supp.  
7 2017, Section 2-101), is amended to read as follows:

8 Section 2-101. As used in the Uniform Controlled Dangerous  
9 Substances Act:

10 1. "Administer" means the direct application of a controlled  
11 dangerous substance, whether by injection, inhalation, ingestion or  
12 any other means, to the body of a patient, animal or research  
13 subject by:

14 a. a practitioner (or, in the presence of the  
15 practitioner, by the authorized agent of the  
16 practitioner), or

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

19 2. "Agent" means a peace officer appointed by and who acts on  
20 behalf of the Director of the Oklahoma State Bureau of Narcotics and  
21 Dangerous Drugs Control or an authorized person who acts on behalf  
22 of or at the direction of a person who manufactures, distributes,  
23 dispenses, prescribes, administers or uses for scientific purposes  
24 controlled dangerous substances but does not include a common or

1 contract carrier, public warehouser or employee thereof, or a person  
2 required to register under the Uniform Controlled Dangerous  
3 Substances Act;

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

6 4. "Bureau" means the Oklahoma State Bureau of Narcotics and  
7 Dangerous Drugs Control;

8 5. "Coca leaves" includes cocaine and any compound,  
9 manufacture, salt, derivative, mixture or preparation of coca  
10 leaves, except derivatives of coca leaves which do not contain  
11 cocaine or ecgonine;

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

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

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

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1       9. "Counterfeit substance" means a controlled substance which,  
2 or the container or labeling of which without authorization, bears  
3 the trademark, trade name or other identifying marks, imprint,  
4 number or device or any likeness thereof of a manufacturer,  
5 distributor or dispenser other than the person who in fact  
6 manufactured, distributed or dispensed the substance;

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

11       11. "Dispense" means to deliver a controlled dangerous  
12 substance to an ultimate user or human research subject by or  
13 pursuant to the lawful order of a practitioner, including the  
14 prescribing, administering, packaging, labeling or compounding  
15 necessary to prepare the substance for such distribution.

16 "Dispenser" is a practitioner who delivers a controlled dangerous  
17 substance to an ultimate user or human research subject;

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

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

1 14. "Drug" means articles:

2 a. recognized in the official United States

3 Pharmacopoeia, official Homeopathic Pharmacopoeia of  
4 the United States, or official National Formulary, or  
5 any supplement to any of them,

6 b. intended for use in the diagnosis, cure, mitigation,  
7 treatment or prevention of disease in man or other  
8 animals,

9 c. other than food, intended to affect the structure or  
10 any function of the body of man or other animals, and

11 d. intended for use as a component of any article  
12 specified in this paragraph;

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

15 15. "Drug-dependent person" means a person who is using a  
16 controlled dangerous substance and who is in a state of psychic or  
17 physical dependence, or both, arising from administration of that  
18 controlled dangerous substance on a continuous basis. Drug  
19 dependence is characterized by behavioral and other responses which  
20 include a strong compulsion to take the substance on a continuous  
21 basis in order to experience its psychic effects, or to avoid the  
22 discomfort of its absence;

23 16. "Home care agency" means any sole proprietorship,  
24 partnership, association, corporation, or other organization which

1 administers, offers, or provides home care services, for a fee or  
2 pursuant to a contract for such services, to clients in their place  
3 of residence;

4 17. "Home care services" means skilled or personal care  
5 services provided to clients in their place of residence for a fee;

6 18. "Hospice" means a centrally administered, nonprofit or  
7 profit, medically directed, nurse-coordinated program which provides  
8 a continuum of home and inpatient care for the terminally ill  
9 patient and the patient's family. Such term shall also include a  
10 centrally administered, nonprofit or profit, medically directed,  
11 nurse-coordinated program if such program is licensed pursuant to  
12 the provisions of this act. A hospice program offers palliative and  
13 supportive care to meet the special needs arising out of the  
14 physical, emotional and spiritual stresses which are experienced  
15 during the final stages of illness and during dying and bereavement.  
16 This care is available twenty-four (24) hours a day, seven (7) days  
17 a week, and is provided on the basis of need, regardless of ability  
18 to pay. "Class A" Hospice refers to Medicare certified hospices.  
19 "Class B" refers to all other providers of hospice services;

20 19. "Imitation controlled substance" means a substance that is  
21 not a controlled dangerous substance, which by dosage unit  
22 appearance, color, shape, size, markings or by representations made,  
23 would lead a reasonable person to believe that the substance is a  
24 controlled dangerous substance. In the event the appearance of the



1 dosage unit is not reasonably sufficient to establish that the  
2 substance is an "imitation controlled substance", the court or  
3 authority concerned should consider, in addition to all other  
4 factors, the following factors as related to "representations made"  
5 in determining whether the substance is an "imitation controlled  
6 substance":

- 7 a. statements made by an owner or by any other person in  
8 control of the substance concerning the nature of the  
9 substance, or its use or effect,
- 10 b. statements made to the recipient that the substance  
11 may be resold for inordinate profit,
- 12 c. whether the substance is packaged in a manner normally  
13 used for illicit controlled substances,
- 14 d. evasive tactics or actions utilized by the owner or  
15 person in control of the substance to avoid detection  
16 by law enforcement authorities,
- 17 e. prior convictions, if any, of an owner, or any other  
18 person in control of the object, under state or  
19 federal law related to controlled substances or fraud,  
20 and
- 21 f. the proximity of the substances to controlled  
22 dangerous substances;

23 20. "Immediate precursor" means a substance which the Director  
24 has found to be and by regulation designates as being the principal

1 compound commonly used or produced primarily for use, and which is  
2 an immediate chemical intermediary used, or likely to be used, in  
3 the manufacture of a controlled dangerous substance, the control of  
4 which is necessary to prevent, curtail or limit such manufacture;

5 21. "Laboratory" means a laboratory approved by the Director as  
6 proper to be entrusted with the custody of controlled dangerous  
7 substances and the use of controlled dangerous substances for  
8 scientific and medical purposes and for purposes of instruction;

9 22. "Manufacture" means the production, preparation,  
10 propagation, compounding or processing of a controlled dangerous  
11 substance, either directly or indirectly by extraction from  
12 substances of natural or synthetic origin, or independently by means  
13 of chemical synthesis or by a combination of extraction and chemical  
14 synthesis. "Manufacturer" includes any person who packages,  
15 repackages or labels any container of any controlled dangerous  
16 substance, except practitioners who dispense or compound  
17 prescription orders for delivery to the ultimate consumer;

18 23. "Marihuana" means all parts of the plant *Cannabis sativa*  
19 L., whether growing or not; the seeds thereof; the resin extracted  
20 from any part of such plant; and every compound, manufacture, salt,  
21 derivative, mixture or preparation of such plant, its seeds or  
22 resin, but shall not include:

23 a. the mature stalks of such plant or fiber produced from  
24 such stalks,

- 1           b.   oil or cake made from the seeds of such plant,  
2               including cannabidiol derived from the seeds of the  
3               marihuana plant,
- 4           c.   any other compound, manufacture, salt, derivative,  
5               mixture or preparation of such mature stalks (except  
6               the resin extracted therefrom), including cannabidiol  
7               derived from mature stalks, fiber, oil or cake,
- 8           d.   the sterilized seed of such plant which is incapable  
9               of germination,
- 10          e.   for any person participating in a clinical trial to  
11               administer cannabidiol for the treatment of severe  
12               forms of epilepsy pursuant to Section 2-802 of this  
13               title, a drug or substance approved by the federal  
14               Food and Drug Administration for use by those  
15               participants,
- 16          f.   for any person or the parents, legal guardians or  
17               caretakers of the person who have received a written  
18               certification from a physician licensed in this state  
19               that the person has been diagnosed by a physician as  
20               having Lennox-Gastaut Syndrome, Dravet Syndrome, also  
21               known as Severe Myoclonic Epilepsy of Infancy, or any  
22               other severe form of epilepsy that is not adequately  
23               treated by traditional medical therapies, spasticity  
24               due to multiple sclerosis or due to paraplegia,

1           intractable nausea and vomiting, appetite stimulation  
2           with chronic wasting diseases, the substance  
3           cannabidiol, a nonpsychoactive cannabinoid, found in  
4           the plant Cannabis sativa L. or any other preparation  
5           thereof, that has a tetrahydrocannabinol concentration  
6           of not more than three-tenths of one percent (0.3%)  
7           and that is delivered to the patient in the form of a  
8           liquid,

9           g. any federal Food and Drug Administration-approved  
10           cannabidiol drug or substance, or

11           h. industrial hemp, from the plant Cannabis sativa L. and  
12           any part of such plant, whether growing or not, with a  
13           delta-9 tetrahydrocannabinol concentration of not more  
14           than three-tenths of one percent (0.3%) on a dry  
15           weight basis which shall ~~not~~ only be grown ~~anywhere in~~  
16           ~~the State of Oklahoma but~~ pursuant to the Oklahoma  
17           Industrial Hemp Agricultural Pilot Program and may be  
18           shipped to Oklahoma pursuant to the provisions of  
19           subparagraph e or f of this paragraph;

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

1       25. "Mid-level practitioner" means an advanced practice nurse  
2 as defined and within parameters specified in Section 567.3a of  
3 Title 59 of the Oklahoma Statutes, or a certified animal euthanasia  
4 technician as defined in Section 698.2 of Title 59 of the Oklahoma  
5 Statutes, or an animal control officer registered by the Oklahoma  
6 State Bureau of Narcotics and Dangerous Drugs Control under  
7 subsection B of Section 2-301 of this title within the parameters of  
8 such officer's duty under Sections 501 through 508 of Title 4 of the  
9 Oklahoma Statutes;

10       26. "Narcotic drug" means any of the following, whether  
11 produced directly or indirectly by extraction from substances of  
12 vegetable origin, or independently by means of chemical synthesis,  
13 or by a combination of extraction and chemical synthesis:

- 14           a. opium, coca leaves and opiates,
- 15           b. a compound, manufacture, salt, derivative or  
16                preparation of opium, coca leaves or opiates,
- 17           c. cocaine, its salts, optical and geometric isomers, and  
18                salts of isomers,
- 19           d. ecgonine, its derivatives, their salts, isomers and  
20                salts of isomers, and
- 21           e. a substance, and any compound, manufacture, salt,  
22                derivative or preparation thereof, which is chemically  
23                identical with any of the substances referred to in  
24                subparagraphs a through d of this paragraph, except

1           that the words "narcotic drug" as used in Section 2-  
2           101 et seq. of this title shall not include  
3           decocainized coca leaves or extracts of coca leaves,  
4           which extracts do not contain cocaine or ecgonine;

5       27. "Opiate" means any substance having an addiction-forming or  
6 addiction-sustaining liability similar to morphine or being capable  
7 of conversion into a drug having such addiction-forming or  
8 addiction-sustaining liability. It does not include, unless  
9 specifically designated as controlled under the Uniform Controlled  
10 Dangerous Substances Act, the dextrorotatory isomer of 3-methoxy-n-  
11 methyl-morphinan and its salts (dextromethorphan). It does include  
12 its racemic and levorotatory forms;

13       28. "Opium poppy" means the plant of the species *Papaver*  
14 *somniferum* L., except the seeds thereof;

15       29. "Peace officer" means a police officer, sheriff, deputy  
16 sheriff, district attorney's investigator, investigator from the  
17 Office of the Attorney General, or any other person elected or  
18 appointed by law to enforce any of the criminal laws of this state  
19 or of the United States;

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

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

1 32. "Practitioner" means:

- 2 a. (1) a medical doctor or osteopathic physician,  
3 (2) a dentist,  
4 (3) a podiatrist,  
5 (4) an optometrist,  
6 (5) a veterinarian,  
7 (6) a physician assistant under the supervision of a  
8 licensed medical doctor or osteopathic physician,  
9 (7) a scientific investigator, or  
10 (8) any other person,

11 licensed, registered or otherwise permitted to  
12 prescribe, distribute, dispense, conduct research with  
13 respect to, use for scientific purposes or administer  
14 a controlled dangerous substance in the course of  
15 professional practice or research in this state, or

- 16 b. a pharmacy, hospital, laboratory or other institution  
17 licensed, registered or otherwise permitted to  
18 distribute, dispense, conduct research with respect  
19 to, use for scientific purposes or administer a  
20 controlled dangerous substance in the course of  
21 professional practice or research in this state;

22 33. "Production" includes the manufacture, planting,  
23 cultivation, growing or harvesting of a controlled dangerous  
24 substance;

1        34. "State" means the State of Oklahoma or any other state of  
2 the United States;

3        35. "Ultimate user" means a person who lawfully possesses a  
4 controlled dangerous substance for the person's own use or for the  
5 use of a member of the person's household or for administration to  
6 an animal owned by the person or by a member of the person's  
7 household;

8        36. "Drug paraphernalia" means all equipment, products and  
9 materials of any kind which are used, intended for use, or fashioned  
10 specifically for use in planting, propagating, cultivating, growing,  
11 harvesting, manufacturing, compounding, converting, producing,  
12 processing, preparing, testing, analyzing, packaging, repackaging,  
13 storing, containing, concealing, injecting, ingesting, inhaling or  
14 otherwise introducing into the human body, a controlled dangerous  
15 substance in violation of the Uniform Controlled Dangerous  
16 Substances Act including, but not limited to:

- 17            a. kits used, intended for use, or fashioned specifically  
18                    for use in planting, propagating, cultivating, growing  
19                    or harvesting of any species of plant which is a  
20                    controlled dangerous substance or from which a  
21                    controlled dangerous substance can be derived,
- 22            b. kits used, intended for use, or fashioned specifically  
23                    for use in manufacturing, compounding, converting,

24



1 producing, processing or preparing controlled  
2 dangerous substances,

3 c. isomerization devices used, intended for use, or  
4 fashioned specifically for use in increasing the  
5 potency of any species of plant which is a controlled  
6 dangerous substance,

7 d. testing equipment used, intended for use, or fashioned  
8 specifically for use in identifying, or in analyzing  
9 the strength, effectiveness or purity of controlled  
10 dangerous substances,

11 e. scales and balances used, intended for use, or  
12 fashioned specifically for use in weighing or  
13 measuring controlled dangerous substances,

14 f. diluents and adulterants, such as quinine  
15 hydrochloride, mannitol, mannite, dextrose and  
16 lactose, used, intended for use, or fashioned  
17 specifically for use in cutting controlled dangerous  
18 substances,

19 g. separation gins and sifters used, intended for use, or  
20 fashioned specifically for use in removing twigs and  
21 seeds from, or in otherwise cleaning or refining,  
22 marihuana,

- 1 h. blenders, bowls, containers, spoons and mixing devices  
2 used, intended for use, or fashioned specifically for  
3 use in compounding controlled dangerous substances,
- 4 i. capsules, balloons, envelopes and other containers  
5 used, intended for use, or fashioned specifically for  
6 use in packaging small quantities of controlled  
7 dangerous substances,
- 8 j. containers and other objects used, intended for use,  
9 or fashioned specifically for use in parenterally  
10 injecting controlled dangerous substances into the  
11 human body,
- 12 k. hypodermic syringes, needles and other objects used,  
13 intended for use, or fashioned specifically for use in  
14 parenterally injecting controlled dangerous substances  
15 into the human body,
- 16 l. objects used, intended for use, or fashioned  
17 specifically for use in ingesting, inhaling or  
18 otherwise introducing marihuana, cocaine, hashish or  
19 hashish oil into the human body, such as:
- 20 (1) metal, wooden, acrylic, glass, stone, plastic or  
21 ceramic pipes with or without screens, permanent  
22 screens, hashish heads or punctured metal bowls,
- 23 (2) water pipes,
- 24 (3) carburetion tubes and devices,

- 1 (4) smoking and carburetion masks,  
2 (5) roach clips, meaning objects used to hold burning  
3 material, such as a marihuana cigarette, that has  
4 become too small or too short to be held in the  
5 hand,  
6 (6) miniature cocaine spoons and cocaine vials,  
7 (7) chamber pipes,  
8 (8) carburetor pipes,  
9 (9) electric pipes,  
10 (10) air-driven pipes,  
11 (11) chillums,  
12 (12) bonges, or  
13 (13) ice pipes or chillers,  
14 m. all hidden or novelty pipes, and  
15 n. any pipe that has a tobacco bowl or chamber of less  
16 than one-half (1/2) inch in diameter in which there is  
17 any detectable residue of any controlled dangerous  
18 substance as defined in this section or any other  
19 substances not legal for possession or use;  
20 provided, however, the term "drug paraphernalia" shall not include  
21 separation gins intended for use in preparing tea or spice, clamps  
22 used for constructing electrical equipment, water pipes designed for  
23 ornamentation in which no detectable amount of an illegal substance  
24 is found or pipes designed and used solely for smoking tobacco,

1 traditional pipes of an American Indian tribal religious ceremony,  
2 or antique pipes that are thirty (30) years of age or older;

3 37. a. "Synthetic controlled substance" means a substance:

4 (1) the chemical structure of which is substantially  
5 similar to the chemical structure of a controlled  
6 dangerous substance in Schedule I or II,

7 (2) which has a stimulant, depressant, or  
8 hallucinogenic effect on the central nervous  
9 system that is substantially similar to or  
10 greater than the stimulant, depressant or  
11 hallucinogenic effect on the central nervous  
12 system of a controlled dangerous substance in  
13 Schedule I or II, or

14 (3) with respect to a particular person, which such  
15 person represents or intends to have a stimulant,  
16 depressant, or hallucinogenic effect on the  
17 central nervous system that is substantially  
18 similar to or greater than the stimulant,  
19 depressant, or hallucinogenic effect on the  
20 central nervous system of a controlled dangerous  
21 substance in Schedule I or II.

22 b. The designation of gamma butyrolactone or any other  
23 chemical as a precursor, pursuant to Section 2-322 of  
24 this title, does not preclude a finding pursuant to

1           subparagraph a of this paragraph that the chemical is  
2           a synthetic controlled substance.

3           c. "Synthetic controlled substance" does not include:

4           (1) a controlled dangerous substance,

5           (2) any substance for which there is an approved new  
6           drug application,

7           (3) with respect to a particular person any  
8           substance, if an exemption is in effect for  
9           investigational use, for that person under the  
10          provisions of Section 505 of the Federal Food,  
11          Drug and Cosmetic Act, Title 21 of the United  
12          States Code, Section 355, to the extent conduct  
13          with respect to such substance is pursuant to  
14          such exemption, or

15          (4) any substance to the extent not intended for  
16          human consumption before such an exemption takes  
17          effect with respect to that substance.

18          d. Prima facie evidence that a substance containing  
19          salvia divinorum has been enhanced, concentrated or  
20          chemically or physically altered shall give rise to a  
21          rebuttable presumption that the substance is a  
22          synthetic controlled substance;

1 38. "Tetrahydrocannabinols" means all substances that have been  
2 chemically synthesized to emulate the tetrahydrocannabinols of  
3 marihuana;

4 39. "Isomer" means the optical isomer, except as used in  
5 subsections C and F of Section 2-204 of this title and paragraph 4  
6 of subsection A of Section 2-206 of this title. As used in  
7 subsections C and F of Section 2-204 of this title, "isomer" means  
8 the optical, positional or geometric isomer. As used in paragraph 4  
9 of subsection A of Section 2-206 of this title, the term "isomer"  
10 means the optical or geometric isomer;

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

15 41. "Anhydrous ammonia" means any substance that exhibits  
16 cryogenic evaporative behavior and tests positive for ammonia.

17 SECTION 12. It being immediately necessary for the preservation  
18 of the public peace, health or safety, an emergency is hereby  
19 declared to exist, by reason whereof this act shall take effect and  
20 be in full force from and after its passage and approval.

1 Passed the House of Representatives the 5th day of March, 2018.

2  
3 \_\_\_\_\_  
4 Presiding Officer of the House  
of Representatives

5 Passed the Senate the \_\_\_\_ day of \_\_\_\_\_, 2018.

6  
7  
8 \_\_\_\_\_  
9 Presiding Officer of the Senate