

1 **SENATE FLOOR VERSION**

2 February 17, 2022

3 SENATE BILL NO. 1338

By: Bullard

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5  
6 An Act relating to the Uniformed Controlled Dangerous  
7 Substances Act; amending 2 O.S. 2021, Section 3-601,  
8 which relates to the Oklahoma Industrial Hemp  
9 Remediation Program; modifying definition; amending  
10 63 O.S. 2021, Section 2-101, as last amended by  
11 Section 1, Chapter 222, O.S.L. 2021, which relates to  
12 definitions; modifying definition; amending 63 O.S.  
13 2021, Section 427.2, as last amended by Section 4,  
14 Chapter 584, O.S.L. 2021, which relates to  
15 definitions; modifying definition; updating statutory  
16 language; and providing an effective date.

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

18 SECTION 1. AMENDATORY 2 O.S. 2021, Section 3-601, is  
19 amended to read as follows:

20 Section 3-601. A. This act shall be known and may be cited as  
21 the "Oklahoma Industrial Hemp Remediation Program".

22 B. As used in the Oklahoma Industrial Hemp Remediation Program,  
23 the following words and terms, and any derivative of such words or  
24 terms, shall have the following meanings, unless the context clearly  
25 indicates otherwise:

26 1. "Cannabis" means a genus of flowering plants in the family  
27 Cannabaceae of which Cannabis sativa is a species and Cannabis

1 indica and Cannabis ruderalis are subspecies thereof. Cannabis  
2 refers to any form of the plant in which the delta-9  
3 tetrahydrocannabinol concentration on a dry-weight basis has not yet  
4 been determined;

5 2. "Certified laboratory" means the laboratory operated by the  
6 Oklahoma Department of Agriculture, Food, and Forestry or a  
7 laboratory located in Oklahoma that is certified by the Department;

8 3. "Commercial sale" means the sale of a product in the stream  
9 of commerce at retail, at wholesale or on the Internet;

10 4. "CSA" means the federal Controlled Substances Act;

11 5. "DEA" means the United States Drug Enforcement  
12 Administration;

13 6. "Department" means the Oklahoma Department of Agriculture,  
14 Food, and Forestry;

15 7. "Hemp" means the plant Cannabis sativa L. and any part of  
16 such plant including, but not limited to, the seeds and all  
17 derivatives, extracts, cannabinoids, isomers, acids, salts and salts  
18 of isomers, whether growing or not, and grown from a certified seed  
19 with a delta-9 or delta-8 tetrahydrocannabinol concentration of not  
20 more than three-tenths of one percent (0.3%) on a dry-weight basis.  
21 Hemp and hemp-derived cannabinoids, including cannabidiol, shall be  
22 considered an agricultural commodity and not a controlled substance  
23 due to the presence of hemp or hemp-derived cannabinoids;

24

1           8. "Hemp Program" means the Oklahoma Industrial Hemp  
2 Remediation Program and any final ruling from the USDA;

3           9. "Law enforcement" means any federal, state or local agencies  
4 responsible for maintaining public order and enforcing the law;

5           10. "License" means the written authorization by the Department  
6 for any person to grow, process, handle or transport certified seeds  
7 or hemp in this state;

8           11. "Person" means any natural person or any corporation,  
9 general partnership, limited partnership, limited liability  
10 partnership, limited liability company, trust, estate, charitable  
11 organization, joint stock company, joint venture, association or any  
12 other business or similar organization recognized by the state;

13           12. "Processor" means any person who is licensed by the  
14 Department to process hemp in this state;

15           13. "State" means the State of Oklahoma;

16           14. "THC" means delta-9 tetrahydrocannabinol, which is a  
17 psychoactive component in cannabis plants;

18           15. "Tracking software" means software that is approved by the  
19 Department and is capable of transparently tracking hemp in any  
20 state or form whatsoever including, but not limited to, a certified  
21 seed, any stage of growth, processing or handling, and any hemp  
22 product; and

23           16. "USDA" means the United States Department of Agriculture.  
24

1 C. In the event that any hemp produced under the Hemp Program  
2 is determined by testing results to be noncompliant with the Hemp  
3 Program, the person holding the license for the noncompliant hemp  
4 may request approval from the Department to remediate the  
5 noncompliant hemp.

6 D. If the Department approves the remediation of the  
7 noncompliant hemp, the person holding the license shall promptly  
8 have the noncompliant hemp extracted by a licensed processor into  
9 concentrated form and the hemp concentrate shall be sampled by a  
10 certified laboratory for compliance with USDA levels for THC in  
11 concentrated form.

12 E. If the samples of the hemp concentrate are below USDA levels  
13 for THC, the hemp concentrate shall be compliant as a hemp product  
14 with the Hemp Program and may be used in commercial sales.

15 F. If the samples of the hemp concentrate are above the USDA  
16 levels for THC, the hemp concentrate shall be noncompliant with the  
17 Hemp Program and shall be destroyed in accordance with the CSA and  
18 DEA regulations found at 21 C.F.R., Section 1317.15, as enforced by  
19 federal, state and local law enforcement. The person holding the  
20 license for the noncompliant hemp concentrate shall promptly notify  
21 the Department and USDA of its intent to destroy the noncompliant  
22 hemp concentrate and verify destruction by submitting required  
23 documentation using the tracking software.

24

1 SECTION 2. AMENDATORY 63 O.S. 2021, Section 2-101, as  
2 last amended by Section 1, Chapter 222, O.S.L. 2021, is amended to  
3 read as follows:

4 Section 2-101. As used in the Uniform Controlled Dangerous  
5 Substances Act:

6 1. "Administer" means the direct application of a controlled  
7 dangerous substance, whether by injection, inhalation, ingestion or  
8 any other means, to the body of a patient, animal or research  
9 subject by:

10 a. a practitioner (or, in the presence of the  
11 practitioner, by the authorized agent of the  
12 practitioner), or

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

15 2. "Agent" means a peace officer appointed by and who acts on  
16 behalf of the Director of the Oklahoma State Bureau of Narcotics and  
17 Dangerous Drugs Control or an authorized person who acts on behalf  
18 of or at the direction of a person who manufactures, distributes,  
19 dispenses, prescribes, administers or uses for scientific purposes  
20 controlled dangerous substances but does not include a common or  
21 contract carrier, public warehouse or employee thereof, or a person  
22 required to register under the Uniform Controlled Dangerous  
23 Substances Act;

24

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

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

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

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

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

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

21           9. "Counterfeit substance" means a controlled substance which,  
22 or the container or labeling of which without authorization, bears  
23 the trademark, trade name or other identifying marks, imprint,  
24 number or device or any likeness thereof of a manufacturer,

1 distributor or dispenser other than the person who in fact  
2 manufactured, distributed or dispensed the substance;

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

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

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

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

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

21 14. "Drug" means articles:

22 a. recognized in the official United States Pharmacopeia,  
23 official Homeopathic Pharmacopoeia of the United  
24

1 States, or official National Formulary, or any  
2 supplement to any of them,

3 b. intended for use in the diagnosis, cure, mitigation,  
4 treatment or prevention of disease in man or other  
5 animals,

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

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

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

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

20 16. "Home care agency" means any sole proprietorship,  
21 partnership, association, corporation, or other organization which  
22 administers, offers, or provides home care services, for a fee or  
23 pursuant to a contract for such services, to clients in their place  
24 of residence;



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

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

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

1 authority concerned should consider, in addition to all other  
2 factors, the following factors as related to "representations made"  
3 in determining whether the substance is an "imitation controlled  
4 substance":

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

21 20. "Immediate precursor" means a substance which the Director  
22 has found to be and by regulation designates as being the principal  
23 compound commonly used or produced primarily for use, and which is  
24 an immediate chemical intermediary used, or likely to be used, in

1 the manufacture of a controlled dangerous substance, the control of  
2 which is necessary to prevent, curtail or limit such manufacture;

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

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

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

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

- 1           b.   oil or cake made from the seeds of such plant,  
2                   including cannabidiol derived from the seeds of the  
3                   marijuana 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 drug  
10           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 or delta-8 tetrahydrocannabinol concentration  
14           of not more than three-tenths of one percent (0.3%) on  
15           a dry weight basis which shall only be grown pursuant  
16           to the Oklahoma Industrial Hemp Program and may be  
17           shipped intrastate and interstate;

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

23           25. "Mid-level practitioner" means an Advanced Practice  
24           Registered Nurse as defined and within parameters specified in

1 Section 567.3a of Title 59 of the Oklahoma Statutes, or a certified  
2 animal euthanasia technician as defined in Section 698.2 of Title 59  
3 of the Oklahoma Statutes, or an animal control officer registered by  
4 the Oklahoma State Bureau of Narcotics and Dangerous Drugs Control  
5 under subsection B of Section 2-301 of this title within the  
6 parameters of such officer's duties under Sections 501 through 508  
7 of Title 4 of the Oklahoma Statutes;

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

- 12 a. opium, coca leaves and opiates,
- 13 b. a compound, manufacture, salt, derivative or  
14 preparation of opium, coca leaves or opiates,
- 15 c. cocaine, its salts, optical and geometric isomers, and  
16 salts of isomers,
- 17 d. ecgonine, its derivatives, their salts, isomers and  
18 salts of isomers, and
- 19 e. a substance, and any compound, manufacture, salt,  
20 derivative or preparation thereof, which is chemically  
21 identical with any of the substances referred to in  
22 subparagraphs a through d of this paragraph, except  
23 that the words "narcotic drug" as used in Section 2-  
24 101 et seq. of this title shall not include

1           decocainized coca leaves or extracts of coca leaves,  
2           which extracts do not contain cocaine or ecgonine;

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

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

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

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

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

24       32. "Practitioner" means:

- 1 a. (1) a medical doctor or osteopathic physician,  
2 (2) a dentist,  
3 (3) a podiatrist,  
4 (4) an optometrist,  
5 (5) a veterinarian,  
6 (6) a physician assistant or Advanced Practice  
7 Registered Nurse 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~~ this state or any other  
2 state of 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,

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 marijuana,

- 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 marijuana, 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 marijuana 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 marijuana;

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;

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

17 42. "Acute pain" means pain, whether resulting from disease,  
18 accidental or intentional trauma or other cause, that the  
19 practitioner reasonably expects to last only a short period of time.  
20 "Acute pain" does not include chronic pain, pain being treated as  
21 part of cancer care, hospice or other end-of-life care, or pain  
22 being treated as part of palliative care;

23 43. "Chronic pain" means pain that persists beyond the usual  
24 course of an acute disease or healing of an injury. "Chronic pain"

1 may or may not be associated with an acute or chronic pathologic  
2 process that causes continuous or intermittent pain over months or  
3 years;

4 44. "Initial prescription" means a prescription issued to a  
5 patient who:

6 a. has never previously been issued a prescription for  
7 the drug or its pharmaceutical equivalent in the past  
8 year, or

9 b. requires a prescription for the drug or its  
10 pharmaceutical equivalent due to a surgical procedure  
11 or new acute event and has previously had a  
12 prescription for the drug or its pharmaceutical  
13 equivalent within the past year.

14 When determining whether a patient was previously issued a  
15 prescription for a drug or its pharmaceutical equivalent, the  
16 practitioner shall consult with the patient and review the medical  
17 record and prescription monitoring information of the patient;

18 45. "Patient-provider agreement" means a written contract or  
19 agreement that is executed between a practitioner and a patient,  
20 prior to the commencement of treatment for chronic pain using an  
21 opioid drug as a means to:

22 a. explain the possible risk of development of physical  
23 or psychological dependence in the patient and prevent  
24 the possible development of addiction,



- 1           b.    document the understanding of both the practitioner  
2                    and the patient regarding the patient-provider  
3                    agreement of the patient,
- 4           c.    establish the rights of the patient in association  
5                    with treatment and the obligations of the patient in  
6                    relation to the responsible use, discontinuation of  
7                    use, and storage of opioid drugs, including any  
8                    restrictions on the refill of prescriptions or the  
9                    acceptance of opioid prescriptions from practitioners,
- 10          d.    identify the specific medications and other modes of  
11                    treatment, including physical therapy or exercise,  
12                    relaxation or psychological counseling, that are  
13                    included as a part of the patient-provider agreement,
- 14          e.    specify the measures the practitioner may employ to  
15                    monitor the compliance of the patient including, but  
16                    not limited to, random specimen screens and pill  
17                    counts, and
- 18          f.    delineate the process for terminating the agreement,  
19                    including the consequences if the practitioner has  
20                    reason to believe that the patient is not complying  
21                    with the terms of the agreement.  Compliance with the  
22                    "consent items" shall constitute a valid, informed  
23                    consent for opioid therapy.  The practitioner shall be  
24                    held harmless from civil litigation for failure to

1            treat pain if the event occurs because of nonadherence  
2            by the patient with any of the provisions of the  
3            patient-provider agreement;

4            46. "Serious illness" means a medical illness or physical  
5 injury or condition that substantially affects quality of life for  
6 more than a short period of time. "Serious illness" includes, but  
7 is not limited to, Alzheimer's disease or related dementias, lung  
8 disease, cancer, heart failure, renal failure, liver failure or  
9 chronic, unremitting or intractable pain such as neuropathic pain;  
10 and

11           47. "Surgical procedure" means a procedure that is performed  
12 for the purpose of structurally altering the human body by incision  
13 or destruction of tissues as part of the practice of medicine. This  
14 term includes the diagnostic or therapeutic treatment of conditions  
15 or disease processes by use of instruments such as lasers,  
16 ultrasound, ionizing, radiation, scalpels, probes or needles that  
17 cause localized alteration or transportation of live human tissue by  
18 cutting, burning, vaporizing, freezing, suturing, probing or  
19 manipulating by closed reduction for major dislocations or  
20 fractures, or otherwise altering by any mechanical, thermal, light-  
21 based, electromagnetic or chemical means.

22           SECTION 3.            AMENDATORY            63 O.S. 2021, Section 427.2, as  
23 last amended by Section 4, Chapter 584, O.S.L. 2021, is amended to  
24 read as follows:

1 Section 427.2. As used in the Oklahoma Medical Marijuana and  
2 Patient Protection Act:

3 1. "Advertising" means the act of providing consideration for  
4 the publication, dissemination, solicitation or circulation, of  
5 visual, oral or written communication to induce directly or  
6 indirectly any person to patronize a particular medical marijuana  
7 business, or to purchase particular medical marijuana or a medical  
8 marijuana product. Advertising includes marketing, but does not  
9 include packaging and labeling;

10 2. "Authority" means the Oklahoma Medical Marijuana Authority;

11 3. "Batch number" means a unique numeric or alphanumeric  
12 identifier assigned prior to testing to allow for inventory tracking  
13 and traceability;

14 4. "Cannabinoid" means any of the chemical compounds that are  
15 active principles of marijuana;

16 5. "Caregiver" means a family member or assistant who regularly  
17 looks after a medical marijuana license holder whom a physician  
18 attests needs assistance;

19 6. "Child-resistant" means special packaging that is:

20 a. designed or constructed to be significantly difficult  
21 for children under five (5) years of age to open and  
22 not difficult for normal adults to use properly as  
23 defined by 16 C.F.R. 1700.15 (1995) and 16 C.F.R.  
24 1700.20 (1995),

1           b.   opaque so that the outermost packaging does not allow  
2                   the product to be seen without opening the packaging  
3                   material, and

4           c.   resealable to maintain its child-resistant  
5                   effectiveness for multiple openings for any product  
6                   intended for more than a single use or containing  
7                   multiple servings;

8           7.   "Clone" means a nonflowering plant cut from a mother plant  
9           that is capable of developing into a new plant and has shown no  
10           signs of flowering;

11           8.   "Commissioner" means the State Commissioner of Health;

12           9.   "Complete application" means a document prepared in  
13           accordance with the provisions set forth in the Oklahoma Medical  
14           Marijuana and Patient Protection Act, rules promulgated pursuant  
15           thereto, and the forms and instructions provided by the Department  
16           including any supporting documentation required and the applicable  
17           license application fee;

18           10.  "Department" means the State Department of Health;

19           11.  "Director" means the Executive Director of the Oklahoma  
20           Medical Marijuana Authority;

21           12.  "Dispense" means the selling of medical marijuana or a  
22           medical marijuana product to a qualified patient or the designated  
23           caregiver of the patient that is packaged in a suitable container  
24

1 appropriately labeled for subsequent administration to or use by a  
2 qualifying patient;

3 13. "Dispensary" means a medical marijuana dispensary, an  
4 entity that has been licensed by the Department pursuant to the  
5 Oklahoma Medical Marijuana and Patient Protection Act to purchase  
6 medical marijuana or medical marijuana products from a licensed  
7 medical marijuana commercial grower or medical marijuana processor,  
8 sell medical marijuana or medical marijuana products to patients and  
9 caregivers as defined under the Oklahoma Medical Marijuana and  
10 Patient Protection Act, or sell or transfer products to another  
11 dispensary;

12 14. "Edible medical marijuana product" means any medical-  
13 marijuana-infused product for which the intended use is oral  
14 consumption including, but not limited to, any type of food, drink  
15 or pill;

16 15. "Entity" means an individual, general partnership, limited  
17 partnership, limited liability company, trust, estate, association,  
18 corporation, cooperative or any other legal or commercial entity;

19 16. "Flower" means the reproductive organs of the marijuana or  
20 cannabis plant referred to as the bud or parts of the plant that are  
21 harvested and used to consume in a variety of medical marijuana  
22 products;

23

24

1 17. "Flowering" means the reproductive state of the marijuana  
2 or cannabis plant in which there are physical signs of flower or  
3 budding out of the nodes of the stem;

4 18. "Food-based medical marijuana concentrate" means a medical  
5 marijuana concentrate that was produced by extracting cannabinoids  
6 from medical marijuana through the use of propylene glycol,  
7 glycerin, butter, olive oil, coconut oil or other typical food-safe  
8 cooking fats;

9 19. "Good cause" for purposes of an initial, renewal or  
10 reinstatement license application, or for purposes of discipline of  
11 a licensee, means:

- 12 a. the licensee or applicant has violated, does not meet,  
13 or has failed to comply with any of the terms,  
14 conditions or provisions of the act, any rules  
15 promulgated pursuant thereto, or any supplemental  
16 relevant state or local law, rule or regulation,
- 17 b. the licensee or applicant has failed to comply with  
18 any special terms or conditions that were placed upon  
19 the license pursuant to an order of the State  
20 Department of Health, Oklahoma Medical Marijuana  
21 Authority or the municipality, or
- 22 c. the licensed premises of a medical marijuana business  
23 or applicant have been operated in a manner that  
24 adversely affects the public health or welfare or the

1 safety of the immediate vicinity in which the  
2 establishment is located;

3 20. "Harvest batch" means a specifically identified quantity of  
4 medical marijuana that is uniform in strain, cultivated utilizing  
5 the same cultivation practices, harvested at the same time from the  
6 same location and cured under uniform conditions;

7 21. "Harvested marijuana" means post-flowering medical  
8 marijuana not including trim, concentrate or waste;

9 22. "Heat- or pressure-based medical marijuana concentrate"  
10 means a medical marijuana concentrate that was produced by  
11 extracting cannabinoids from medical marijuana through the use of  
12 heat or pressure;

13 23. "Immature plant" means a nonflowering marijuana plant that  
14 has not demonstrated signs of flowering;

15 24. "Inventory tracking system" means the required tracking  
16 system that accounts for medical marijuana from either the seed or  
17 immature plant stage until the medical marijuana or medical  
18 marijuana product is sold to a patient at a medical marijuana  
19 dispensary, transferred to a medical marijuana research facility,  
20 destroyed by a medical marijuana business or used in a research  
21 project by a medical marijuana research facility;

22 25. "Licensed patient" or "patient" means a person who has been  
23 issued a medical marijuana patient license by the State Department  
24 of Health or Oklahoma Medical Marijuana Authority;

1        26. "Licensed premises" means the premises specified in an  
2 application for a medical marijuana business license, medical  
3 marijuana research facility license or medical marijuana education  
4 facility license pursuant to the Oklahoma Medical Marijuana and  
5 Patient Protection Act that are owned or in possession of the  
6 licensee and within which the licensee is authorized to cultivate,  
7 manufacture, distribute, sell, store, transport, test or research  
8 medical marijuana or medical marijuana products in accordance with  
9 the provisions of the Oklahoma Medical Marijuana and Patient  
10 Protection Act and rules promulgated pursuant thereto;

11        27. "Manufacture" means the production, propagation,  
12 compounding or processing of a medical marijuana product, excluding  
13 marijuana plants, either directly or indirectly by extraction from  
14 substances of natural or synthetic origin, or independently by means  
15 of chemical synthesis, or by a combination of extraction and  
16 chemical synthesis;

17        28. "Marijuana" shall have the same meaning as such term is  
18 defined in Section 2-101 of this title ~~and shall not include any~~  
19 ~~plant or material containing delta-8 or delta-10~~  
20 ~~tetrahydrocannabinol which is grown, processed or sold pursuant to~~  
21 ~~the provisions of the Oklahoma Industrial Hemp Program;~~

22        29. "Material change" means any change that would require a  
23 substantive revision to the standard operating procedures of a  
24



1 licensee for the cultivation or production of medical marijuana,  
2 medical marijuana concentrate or medical marijuana products;

3 30. "Mature plant" means a harvestable female marijuana plant  
4 that is flowering;

5 31. "Medical marijuana business (MMB)" means a licensed medical  
6 marijuana dispensary, medical marijuana processor, medical marijuana  
7 commercial grower, medical marijuana laboratory, medical marijuana  
8 business operator or a medical marijuana transporter;

9 32. "Medical marijuana concentrate" or "concentrate" means a  
10 specific subset of medical marijuana that was produced by extracting  
11 cannabinoids from medical marijuana. Categories of medical  
12 marijuana concentrate include water-based medical marijuana  
13 concentrate, food-based medical marijuana concentrate, solvent-based  
14 medical marijuana concentrate, and heat- or pressure-based medical  
15 marijuana concentrate;

16 33. "Medical marijuana commercial grower" or "commercial  
17 grower" means an entity licensed to cultivate, prepare and package  
18 medical marijuana and transfer or contract for transfer medical  
19 marijuana to a medical marijuana dispensary, medical marijuana  
20 processor, any other medical marijuana commercial grower, medical  
21 marijuana research facility, medical marijuana education facility  
22 and pesticide manufacturers. A commercial grower may sell seeds,  
23 flower or clones to commercial growers pursuant to the Oklahoma  
24 Medical Marijuana and Patient Protection Act;

1       34. "Medical marijuana education facility" or "education  
2 facility" means a person or entity approved pursuant to the Oklahoma  
3 Medical Marijuana and Patient Protection Act to operate a facility  
4 providing training and education to individuals involving the  
5 cultivation, growing, harvesting, curing, preparing, packaging or  
6 testing of medical marijuana, or the production, manufacture,  
7 extraction, processing, packaging or creation of medical-marijuana-  
8 infused products or medical marijuana products as described in the  
9 Oklahoma Medical Marijuana and Patient Protection Act;

10       35. "Medical-marijuana-infused product" means a product infused  
11 with medical marijuana including, but not limited to, edible  
12 products, ointments and tinctures;

13       36. "Medical marijuana product" or "product" means a product  
14 that contains cannabinoids that have been extracted from plant  
15 material or the resin therefrom by physical or chemical means and is  
16 intended for administration to a qualified patient including, but  
17 not limited to, oils, tinctures, edibles, pills, topical forms,  
18 gels, creams, vapors, patches, liquids and forms administered by a  
19 nebulizer, excluding live plant forms which are considered medical  
20 marijuana;

21       37. "Medical marijuana processor" means a person or entity  
22 licensed pursuant to the Oklahoma Medical Marijuana and Patient  
23 Protection Act to operate a business including the production,  
24 manufacture, extraction, processing, packaging or creation of

1 concentrate, medical-marijuana-infused products or medical marijuana  
2 products as described in the Oklahoma Medical Marijuana and Patient  
3 Protection Act;

4 38. "Medical marijuana research facility" or "research  
5 facility" means a person or entity approved pursuant to the Oklahoma  
6 Medical Marijuana and Patient Protection Act to conduct medical  
7 marijuana research. A medical marijuana research facility is not a  
8 medical marijuana business;

9 39. "Medical marijuana testing laboratory" or "laboratory"  
10 means a public or private laboratory licensed pursuant to the  
11 Oklahoma Medical Marijuana and Patient Protection Act, to conduct  
12 testing and research on medical marijuana and medical marijuana  
13 products;

14 40. "Medical marijuana transporter" or "transporter" means a  
15 person or entity that is licensed pursuant to the Oklahoma Medical  
16 Marijuana and Patient Protection Act. A medical marijuana  
17 transporter does not include a medical marijuana business that  
18 transports its own medical marijuana, medical marijuana concentrate  
19 or medical marijuana products to a property or facility adjacent to  
20 or connected to the licensed premises if the property is another  
21 licensed premises of the same medical marijuana business;

22 41. "Medical marijuana waste" or "waste" means unused, surplus,  
23 returned or out-of-date marijuana, plant debris of the plant of the  
24 genus Cannabis including dead plants and all unused plant parts and

1 roots, except the term shall not include roots, stems, stalks and  
2 fan leaves;

3 42. "Medical use" means the acquisition, possession, use,  
4 delivery, transfer or transportation of medical marijuana, medical  
5 marijuana products, medical marijuana devices or paraphernalia  
6 relating to the administration of medical marijuana to treat a  
7 licensed patient;

8 43. "Mother plant" means a marijuana plant that is grown or  
9 maintained for the purpose of generating clones, and that will not  
10 be used to produce plant material for sale to a medical marijuana  
11 processor or medical marijuana dispensary;

12 44. "Oklahoma physician" or "physician" means a physician  
13 licensed by and in good standing with the State Board of Medical  
14 Licensure and Supervision, the State Board of Osteopathic Examiners  
15 or the Board of Podiatric Medical Examiners;

16 45. "Oklahoma resident" means an individual who can provide  
17 proof of residency as required by the Oklahoma Medical Marijuana and  
18 Patient Protection Act;

19 46. "Owner" means, except where the context otherwise requires,  
20 a direct beneficial owner including, but not limited to, all persons  
21 or entities as follows:

22 a. all shareholders owning an interest of a corporate  
23 entity and all officers of a corporate entity,

24 b. all partners of a general partnership,

- 1 c. all general partners and all limited partners that own  
2 an interest in a limited partnership,  
3 d. all members that own an interest in a limited  
4 liability company,  
5 e. all beneficiaries that hold a beneficial interest in a  
6 trust and all trustees of a trust,  
7 f. all persons or entities that own interest in a joint  
8 venture,  
9 g. all persons or entities that own an interest in an  
10 association,  
11 h. the owners of any other type of legal entity, and  
12 i. any other person holding an interest or convertible  
13 note in any entity which owns, operates or manages a  
14 licensed facility;

15 47. "Package" or "packaging" means any container or wrapper  
16 that may be used by a medical marijuana business to enclose or  
17 contain medical marijuana;

18 48. "Person" means a natural person, partnership, association,  
19 business trust, company, corporation, estate, limited liability  
20 company, trust or any other legal entity or organization, or a  
21 manager, agent, owner, director, servant, officer or employee  
22 thereof, except that "person" does not include any governmental  
23 organization;

1 49. "Pesticide" means any substance or mixture of substances  
2 intended for preventing, destroying, repelling or mitigating any  
3 pest or any substance or mixture of substances intended for use as a  
4 plant regulator, defoliant or desiccant, except that the term  
5 "pesticide" shall not include any article that is a "new animal  
6 drug" as designated by the United States Food and Drug  
7 Administration;

8 50. "Production batch" means:

- 9 a. any amount of medical marijuana concentrate of the  
10 same category and produced using the same extraction  
11 methods, standard operating procedures and an  
12 identical group of harvest batch of medical marijuana,  
13 or  
14 b. any amount of medical marijuana product of the same  
15 exact type, produced using the same ingredients,  
16 standard operating procedures and the same production  
17 batch of medical marijuana concentrate;

18 51. "Public institution" means any entity established or  
19 controlled by the federal government, state government, or a local  
20 government or municipality including, but not limited to,  
21 institutions of higher education or related research institutions;

22 52. "Public money" means any funds or money obtained by the  
23 holder from any governmental entity including, but not limited to,  
24 research grants;

1       53. "Recommendation" means a document that is signed or  
2 electronically submitted by a physician on behalf of a patient for  
3 the use of medical marijuana pursuant to the Oklahoma Medical  
4 Marijuana and Patient Protection Act;

5       54. "Registered to conduct business" means a person that has  
6 provided proof that the business applicant is in good standing with  
7 the ~~Oklahoma~~ Secretary of State and Oklahoma Tax Commission;

8       55. "Remediation" means the process by which the medical  
9 marijuana flower or trim, which has failed microbial testing, is  
10 processed into solvent-based medical marijuana concentrate and  
11 retested as required by the Oklahoma Medical Marijuana and Patient  
12 Protection Act;

13       56. "Research project" means a discrete scientific endeavor to  
14 answer a research question or a set of research questions related to  
15 medical marijuana and is required for a medical marijuana research  
16 license. A research project shall include a description of a  
17 defined protocol, clearly articulated goals, defined methods and  
18 outputs, and a defined start and end date. The description shall  
19 demonstrate that the research project will comply with all  
20 requirements in the Oklahoma Medical Marijuana and Patient  
21 Protection Act and rules promulgated pursuant thereto. All research  
22 and development conducted by a medical marijuana research facility  
23 shall be conducted in furtherance of an approved research project;

24

1           57. "Revocation" means the final decision by the Department  
2 that any license issued pursuant to the Oklahoma Medical Marijuana  
3 and Patient Protection Act is rescinded because the individual or  
4 entity does not comply with the applicable requirements set forth in  
5 the Oklahoma Medical Marijuana and Patient Protection Act or rules  
6 promulgated pursuant thereto;

7           58. "School" means a public or private preschool or a public or  
8 private elementary or secondary school which is primarily used for  
9 classroom instruction. A homeschool, daycare or child-care facility  
10 shall not be considered a "school" as used in the Oklahoma Medical  
11 Marijuana and Patient Protection Act;

12           59. "Shipping container" means a hard-sided container with a  
13 lid or other enclosure that can be secured in place. A shipping  
14 container is used solely for the transport of medical marijuana,  
15 medical marijuana concentrate, or medical marijuana products between  
16 medical marijuana businesses, a medical marijuana research facility,  
17 or a medical marijuana education facility;

18           60. "Solvent-based medical marijuana concentrate" means a  
19 medical marijuana concentrate that was produced by extracting  
20 cannabinoids from medical marijuana through the use of a solvent  
21 approved by the Department;

22           61. "State Question" means Oklahoma State Question No. 788,  
23 Initiative Petition No. 412, approved by a majority vote of the  
24 citizens of Oklahoma on June 26, 2018;



1       62. "Strain" means the classification of marijuana or cannabis  
2 plants in either pure sativa, indica, afghanica, ruderalis or hybrid  
3 varieties;

4       63. "THC" means tetrahydrocannabinol, which is the primary  
5 psychotropic cannabinoid in marijuana formed by decarboxylation of  
6 naturally tetrahydrocannabinolic acid, which generally occurs by  
7 exposure to heat;

8       64. "Test batch" means with regard to usable marijuana, a  
9 homogenous, identified quantity of usable marijuana by strain, no  
10 greater than ten (10) pounds, that is harvested during a seven-day  
11 period from a specified cultivation area, and with regard to oils,  
12 vapors and waxes derived from usable marijuana, means an identified  
13 quantity that is uniform, that is intended to meet specifications  
14 for identity, strength and composition, and that is manufactured,  
15 packaged and labeled during a specified time period according to a  
16 single manufacturing, packaging and labeling protocol;

17       65. "Transporter agent" means a person who transports medical  
18 marijuana or medical marijuana products for a licensed transporter  
19 and holds a transporter agent license pursuant to the Oklahoma  
20 Medical Marijuana and Patient Protection Act;

21       66. "Universal symbol" means the image established by the State  
22 Department of Health or Oklahoma Medical Marijuana Authority and  
23 made available to licensees through its website indicating that the  
24 medical marijuana or the medical marijuana product contains THC;

1       67. "Usable marijuana" means the dried leaves, flowers, oils,  
2 vapors, waxes and other portions of the marijuana plant and any  
3 mixture or preparation thereof, excluding seeds, roots, stems,  
4 stalks and fan leaves; and

5       68. "Water-based medical marijuana concentrate" means a  
6 concentrate that was produced by extracting cannabinoids from  
7 medical marijuana through the use of only water, ice or dry ice.

8       SECTION 4. This act shall become effective November 1, 2022.

9       COMMITTEE REPORT BY: COMMITTEE ON BUSINESS, COMMERCE AND TOURISM  
10       February 17, 2022 - DO PASS

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