

1 SECTION 1. NEW LAW A new section of law not to be
2 codified in the Oklahoma Statutes reads as follows:

3 This act shall be known and may be cited as "Katie's Law".

4 SECTION 2. AMENDATORY 63 O.S. 2011, Section 2-101, as
5 last amended by Section 1, Chapter 154, O.S.L. 2014 (63 O.S. Supp.
6 2014, Section 2-101), is amended to read as follows:

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

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

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

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

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

1 required to register under the Uniform Controlled Dangerous
2 Substances Act;

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

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

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

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

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

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

23 9. "Counterfeit substance" means a controlled substance which,
24 or the container or labeling of which without authorization, bears

1 the trademark, trade name or other identifying marks, imprint,
2 number or device or any likeness thereof of a manufacturer,
3 distributor or dispenser other than the person who in fact
4 manufactured, distributed or dispensed the substance;

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

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

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

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

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

23 14. "Drug" means articles:
24

- 1 a. recognized in the official United States
2 Pharmacopoeia, official Homeopathic Pharmacopoeia of
3 the United States, or official National Formulary, or
4 any supplement to any of them,
5 b. intended for use in the diagnosis, cure, mitigation,
6 treatment or prevention of disease in man or other
7 animals,
8 c. other than food, intended to affect the structure or
9 any function of the body of man or other animals, and
10 d. intended for use as a component of any article
11 specified in this paragraph;

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

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

22 16. "Home care agency" means any sole proprietorship,
23 partnership, association, corporation, or other organization which
24 administers, offers, or provides home care services, for a fee or

1 pursuant to a contract for such services, to clients in their place
2 of residence;

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

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

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

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

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

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

1 an immediate chemical intermediary used, or likely to be used, in
2 the manufacture of a controlled dangerous substance, the control of
3 which is necessary to prevent, curtail or limit such manufacture;

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

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

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

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

- 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, ~~or~~
- 8 d. the sterilized seed of such plant which is incapable
9 of germination,
- 10 e. for persons eighteen (18) years of age or younger
11 participating in a clinical trial or in an expanded-
12 access program related to administering cannabidiol
13 for the treatment of severe forms of epilepsy pursuant
14 to Section 4 of this act, a drug or substance approved
15 by the federal Food and Drug Administration for use by
16 those participants, or
- 17 f. for persons eighteen (18) years of age or younger, or
18 the parents, legal guardians, or caretakers of the
19 person, who have received a written certification from
20 a physician licensed in this state that the person has
21 been diagnosed by a physician as having Lennox-Gastaut
22 Syndrome, Dravet Syndrome, also known as Severe
23 Myoclonic Epilepsy of Infancy, or any other severe
24 form of epilepsy that is not adequately treated by

1 traditional medical therapies, the substance
2 cannabidiol, a nonpsychoactive cannabinoid, or any
3 compound, manufacture, salt, derivative, mixture, or
4 preparation of any plant of the Cannabis sativa L. or
5 Cannabis indica that is essentially free from plant
6 material, and has a tetrahydrocannabinol concentration
7 of not more than three-tenths percent (.3%) on a dry
8 weight basis;

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

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

23 26. "Narcotic drug" means any of the following, whether
24 produced directly or indirectly by extraction from substances of

1 vegetable origin, or independently by means of chemical synthesis,
2 or by a combination of extraction and chemical synthesis:

- 3 a. opium, coca leaves and opiates,
- 4 b. a compound, manufacture, salt, derivative or
5 preparation of opium, coca leaves or opiates,
- 6 c. cocaine, its salts, optical and geometric isomers, and
7 salts of isomers,
- 8 d. ecgonine, its derivatives, their salts, isomers and
9 salts of isomers, and
- 10 e. a substance, and any compound, manufacture, salt,
11 derivative or preparation thereof, which is chemically
12 identical with any of the substances referred to in
13 subparagraphs a through d of this paragraph, except
14 that the words "narcotic drug" as used in Section 2-
15 101 et seq. of this title shall not include
16 decocainized coca leaves or extracts of coca leaves,
17 which extracts do not contain cocaine or ecgonine;

18 27. "Opiate" means any substance having an addiction-forming or
19 addiction-sustaining liability similar to morphine or being capable
20 of conversion into a drug having such addiction-forming or
21 addiction-sustaining liability. It does not include, unless
22 specifically designated as controlled under the Uniform Controlled
23 Dangerous Substances Act, the dextrorotatory isomer of 3-methoxy-n-

1 methyl-morphinan and its salts (dextromethorphan). It does include
2 its racemic and levorotatory forms;

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

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

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

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

15 32. "Practitioner" means:

- 16 a. (1) a medical doctor or osteopathic physician,
17 (2) a dentist,
18 (3) a podiatrist,
19 (4) an optometrist,
20 (5) a veterinarian,
21 (6) a physician assistant under the supervision of a
22 licensed medical doctor or osteopathic physician,
23 (7) a scientific investigator, or
24 (8) any other person,

1 licensed, registered or otherwise permitted to
2 prescribe, distribute, dispense, conduct research with
3 respect to, use for scientific purposes or administer
4 a controlled dangerous substance in the course of
5 professional practice or research in this state, or
6 b. a pharmacy, hospital, laboratory or other institution
7 licensed, registered or otherwise permitted to
8 distribute, dispense, conduct research with respect
9 to, use for scientific purposes or administer a
10 controlled dangerous substance in the course of
11 professional practice or research in this state;

12 33. "Production" includes the manufacture, planting,
13 cultivation, growing or harvesting of a controlled dangerous
14 substance;

15 34. "State" means the State of Oklahoma or any other state of
16 the United States;

17 35. "Ultimate user" means a person who lawfully possesses a
18 controlled dangerous substance for the person's own use or for the
19 use of a member of the person's household or for administration to
20 an animal owned by the person or by a member of the person's
21 household;

22 36. "Drug paraphernalia" means all equipment, products and
23 materials of any kind which are used, intended for use, or fashioned
24 specifically for use in planting, propagating, cultivating, growing,

1 harvesting, manufacturing, compounding, converting, producing,
2 processing, preparing, testing, analyzing, packaging, repackaging,
3 storing, containing, concealing, injecting, ingesting, inhaling or
4 otherwise introducing into the human body, a controlled dangerous
5 substance in violation of the Uniform Controlled Dangerous
6 Substances Act including, but not limited to:

- 7 a. kits used, intended for use, or fashioned specifically
8 for use in planting, propagating, cultivating, growing
9 or harvesting of any species of plant which is a
10 controlled dangerous substance or from which a
11 controlled dangerous substance can be derived,
- 12 b. kits used, intended for use, or fashioned specifically
13 for use in manufacturing, compounding, converting,
14 producing, processing or preparing controlled
15 dangerous substances,
- 16 c. isomerization devices used, intended for use, or
17 fashioned specifically for use in increasing the
18 potency of any species of plant which is a controlled
19 dangerous substance,
- 20 d. testing equipment used, intended for use, or fashioned
21 specifically for use in identifying, or in analyzing
22 the strength, effectiveness or purity of controlled
23 dangerous substances,

- 1 e. scales and balances used, intended for use, or
2 fashioned specifically for use in weighing or
3 measuring controlled dangerous substances,
- 4 f. diluents and adulterants, such as quinine
5 hydrochloride, mannitol, mannite, dextrose and
6 lactose, used, intended for use, or fashioned
7 specifically for use in cutting controlled dangerous
8 substances,
- 9 g. separation gins and sifters used, intended for use, or
10 fashioned specifically for use in removing twigs and
11 seeds from, or in otherwise cleaning or refining,
12 marihuana,
- 13 h. blenders, bowls, containers, spoons and mixing devices
14 used, intended for use, or fashioned specifically for
15 use in compounding controlled dangerous substances,
- 16 i. capsules, balloons, envelopes and other containers
17 used, intended for use, or fashioned specifically for
18 use in packaging small quantities of controlled
19 dangerous substances,
- 20 j. containers and other objects used, intended for use,
21 or fashioned specifically for use in parenterally
22 injecting controlled dangerous substances into the
23 human body,
24

1 k. hypodermic syringes, needles and other objects used,
2 intended for use, or fashioned specifically for use in
3 parenterally injecting controlled dangerous substances
4 into the human body,

5 l. objects used, intended for use, or fashioned
6 specifically for use in ingesting, inhaling or
7 otherwise introducing marihuana, cocaine, hashish or
8 hashish oil into the human body, such as:

- 9 (1) metal, wooden, acrylic, glass, stone, plastic or
10 ceramic pipes with or without screens, permanent
11 screens, hashish heads or punctured metal bowls,
12 (2) water pipes,
13 (3) carburetion tubes and devices,
14 (4) smoking and carburetion masks,
15 (5) roach clips, meaning objects used to hold burning
16 material, such as a marihuana cigarette, that has
17 become too small or too short to be held in the
18 hand,
19 (6) miniature cocaine spoons and cocaine vials,
20 (7) chamber pipes,
21 (8) carburetor pipes,
22 (9) electric pipes,
23 (10) air-driven pipes,
24 (11) chillums,

1 (12) bonges, or

2 (13) ice pipes or chillers,

3 m. all hidden or novelty pipes, and

4 n. any pipe that has a tobacco bowl or chamber of less
5 than one-half (1/2) inch in diameter in which there is
6 any detectable residue of any controlled dangerous
7 substance as defined in this section or any other
8 substances not legal for possession or use;

9 provided, however, the term "drug paraphernalia" shall not include
10 separation gins intended for use in preparing tea or spice, clamps
11 used for constructing electrical equipment, water pipes designed for
12 ornamentation in which no detectable amount of an illegal substance
13 is found or pipes designed and used solely for smoking tobacco,
14 traditional pipes of an American Indian tribal religious ceremony,
15 or antique pipes that are thirty (30) years of age or older;

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

17 (1) the chemical structure of which is substantially
18 similar to the chemical structure of a controlled
19 dangerous substance in Schedule I or II,

20 (2) which has a stimulant, depressant, or
21 hallucinogenic effect on the central nervous
22 system that is substantially similar to or
23 greater than the stimulant, depressant or
24 hallucinogenic effect on the central nervous

1 system of a controlled dangerous substance in
2 Schedule I or II, or

3 (3) with respect to a particular person, which such
4 person represents or intends to have a stimulant,
5 depressant, or hallucinogenic effect on the
6 central nervous system that is substantially
7 similar to or greater than the stimulant,
8 depressant, or hallucinogenic effect on the
9 central nervous system of a controlled dangerous
10 substance in Schedule I or II.

11 b. The designation of gamma butyrolactone or any other
12 chemical as a precursor, pursuant to Section 2-322 of
13 this title, does not preclude a finding pursuant to
14 subparagraph a of this paragraph that the chemical is
15 a synthetic controlled substance.

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

- 17 (1) a controlled dangerous substance,
18 (2) any substance for which there is an approved new
19 drug application,
20 (3) with respect to a particular person any
21 substance, if an exemption is in effect for
22 investigational use, for that person under the
23 provisions of Section 505 of the Federal Food,
24 Drug and Cosmetic Act, Title 21 of the United

1 States Code, Section 355, to the extent conduct
2 with respect to such substance is pursuant to
3 such exemption, or

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

7 d. Prima facie evidence that a substance containing
8 salvia divinorum has been enhanced, concentrated or
9 chemically or physically altered shall give rise to a
10 rebuttable presumption that the substance is a
11 synthetic controlled substance;

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

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

22 40. "Hazardous materials" means materials, whether solid,
23 liquid or gas, which are toxic to human, animal, aquatic or plant
24

1 life, and the disposal of which materials is controlled by state or
2 federal guidelines; and

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

5 SECTION 3. NEW LAW A new section of law to be codified
6 in the Oklahoma Statutes as Section 2-801 of Title 63, unless there
7 is created a duplication in numbering, reads as follows:

8 As used in this act:

9 1. "Academic medical center" means a medical school and its
10 affiliated teaching hospitals and clinics that:

11 a. operate a medical residency program for physicians,
12 and

13 b. conduct research that is overseen by the federal
14 Department of Health and Human Services and involves
15 human subjects;

16 2. "Approved source" means a provider approved by the United
17 States Food and Drug Administration which produces cannabidiol that:

18 a. has been manufactured and tested in a facility
19 approved or certified by the United States Food and
20 Drug Administration or similar national regulatory
21 agency in another country which has been approved by
22 the United States Food and Drug Administration, and
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24

1 b. has been tested on animals to demonstrate preliminary
2 effectiveness and to ensure that it is safe to
3 administer to humans;

4 3. "Cannabidiol" means a nonpsychoactive cannabinoid, or any
5 compound, manufacture, salt, derivative, mixture, or preparation of
6 any plant of the Cannabis sativa L. or Cannabis indica that is
7 essentially free from plant material, and has a tetrahydrocannabinol
8 concentration of not more than three-tenths percent (.3%) on a dry
9 weight basis;

10 4. "Physician" means a doctor of medicine or doctor of
11 osteopathic medicine licensed by the Oklahoma Board of Medical
12 Examiners; and

13 5. "Qualifying patient" means any person eighteen (18) years of
14 age or younger who suffers from Lennox-Gastaut Syndrome, Dravet
15 Syndrome, also known as Severe Myoclonic Epilepsy of Infancy, or any
16 other form of refractory epilepsy that is not adequately treated by
17 traditional medical therapies.

18 SECTION 4. NEW LAW A new section of law to be codified
19 in the Oklahoma Statutes as Section 2-802 of Title 63, unless there
20 is created a duplication in numbering, reads as follows:

21 A. A statewide investigational new drug application may be
22 established in this state, if approved by the United States Food and
23 Drug Administration, to conduct expanded-access clinical trials
24

1 using cannabidiol on qualifying patients with severe forms of
2 epilepsy.

3 B. Any physician who is board certified, practicing in an
4 academic medical center in this state, and treating patients with
5 severe forms of epilepsy may serve as the principal investigator for
6 such clinical trials if such physician:

7 1. Applies to and is approved by the United States Food and
8 Drug Administration as the principal investigator in a statewide
9 investigational new drug application; and

10 2. Receives a license from the United States Drug Enforcement
11 Administration.

12 C. Such physician, acting as principal investigator, may
13 include subinvestigators who are also board certified, practice in
14 an academic medical center in this state, and treat patients with
15 severe forms of epilepsy. Such subinvestigators shall be required
16 to comply with the licensing requirement provided in paragraph 2 of
17 subsection B of this section.

18 D. The principal investigator and all subinvestigators shall
19 adhere to the rules and regulations established by the relevant
20 institutional review board for each participating academic medical
21 center and by the United States Food and Drug Administration, the
22 United States Drug Enforcement Administration and the National
23 Institute on Drug Abuse.
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1 E. Nothing in this section shall be construed to prohibit a
2 physician licensed in Oklahoma from applying for Investigational New
3 Drug authorization from the United States Food and Drug
4 Administration.

5 SECTION 5. NEW LAW A new section of law to be codified
6 in the Oklahoma Statutes as Section 2-803 of Title 63, unless there
7 is created a duplication in numbering, reads as follows:

8 A. Expanded-access clinical trials conducted pursuant to a
9 statewide investigational new drug application established pursuant
10 to the provisions of this act shall only utilize cannabidiol which
11 is:

12 1. From an approved source; and

13 2. Approved by the United States Food and Drug Administration
14 to be used for treatment of a condition specified in an
15 investigational new drug application.

16 B. The principal investigator and any subinvestigator may
17 receive cannabidiol directly from an approved source or authorized
18 distributor for an approved source for use in the expanded-access
19 clinical trials.

20 SECTION 6. NEW LAW A new section of law to be codified
21 in the Oklahoma Statutes as Section 2-804 of Title 63, unless there
22 is created a duplication in numbering, reads as follows:

23 A person acting in compliance with the provisions of this act
24 shall not be subject to arrest, prosecution, or any civil or

1 administrative penalty, including a civil penalty or disciplinary
2 action by a professional licensing board, or be denied any right or
3 privilege, for the use, prescription, administration, possession,
4 manufacture, or distribution of medical cannabidiol.

5 SECTION 7. This act shall become effective November 1, 2015.

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7 COMMITTEE REPORT BY: COMMITTEE ON ALCOHOL, TOBACCO, AND CONTROLLED
8 SUBSTANCES, dated 02/04/2015 - DO PASS, As Amended and Coauthored.
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