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AM1542 LB316 AJC - 05/20/2025

AMENDMENTS TO LB316

(Amendments to Final Reading copy)

Introduced by Kauth, 31.

Strike the original sections and all amendments thereto and
 insert the following new sections:

3 Section 1. Section 2-501, Revised Statutes Cumulative Supplement,
4 2024, is amended to read:

5 2-501 Sections 2-501 to 2-518 <u>and sections 3 and 6 of this act</u>shall 6 be known and may be cited as the Nebraska Hemp Farming Act.

Sec. 2. Section 2-503, Revised Statutes Cumulative Supplement, 2024,
is amended to read:

2-503 For purposes of the Nebraska Hemp Farming Act:

(1) Agriculture Improvement Act of 2018 means section 10113 of the
federal Agriculture Improvement Act of 2018, Public Law 115-334, and any
regulations adopted and promulgated under such section, as such section,
act, and regulations existed on January 1, 2024;

(2) Cannabidiol product means a finished hemp consumer product that:
 (a) Contains, as a primary ingredient, cannabidiol extracted or
 derived from hemp;

(b) Complies with the THC limits provided in subdivision (5)(a)(ii)
 of this section; and

<u>(c) Does not contain any cannabinoids created through chemical</u>
 <u>conversion, modification, or synthesis, including, but not limited to,</u>
 <u>hexahydrocannabinol;</u>

22 <u>(3)</u> (2) Cultivate or cultivating means planting, watering, growing, 23 and harvesting a hemp plant or crop. The presence of plants of the plant 24 Cannabis sativa L. growing as uncultivated, naturalized plants in the 25 environment is not cultivating hemp for purposes of the Nebraska Hemp 26 Farming Act;

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(4) Federally compliant hemp means hemp that complies with the 1 2 requirements of the Agriculture Improvement Act of 2018; 3 (5)(a) (3) Hemp means the plant Cannabis sativa L. and any part of such plant, including the viable seeds of such plant and all derivatives, 4 5 extracts, cannabinoids, isomers, acids, salts, and salts of isomers, 6 whether growing or not, that complies with the following THC limits: 7 (i) For raw hemp, a total THC with a delta-9 tetrahydrocannabinol 8 concentration of not more than 0.3 percent on a dry weight basis; and -9 (ii) For processed hemp, including cannabidiol products, not more 10 than the lesser of: (A) A total THC concentration of 0.3 percent on a total weight 11 12 basis; or 13 (B) Ten milligrams of total THC per package. 14 (b) Hemp includes cannabidiol products. 15 (c) Hemp does not include the mature stalks of the plant Cannabis sativa L.; fiber produced from such stalks; oil or cake made from the 16 17 seeds of such plant; any other compound, manufacture, salt, derivative, mixture, or preparation of such mature stalks; or the sterilized seed of 18 19 such plant that is incapable of germination Hemp shall be considered an 20 agricultural commodity. Notwithstanding any other provision of law, hemp 21 shall not be considered a controlled substance under the Uniform 22 Controlled Substances Act; (6) (4) Person means an individual, partnership, corporation, 23 24 limited liability company, association, postsecondary institution, or 25 other legal entity; 26 (7) Raw hemp means hemp that has been harvested and dried but is 27 otherwise unprocessed;

(8) (5) State-program-licensed hemp producer means a person licensed
 under a USDA-approved state or tribal program as authorized under the
 Agriculture Improvement Act of 2018 and includes the authorized employees
 or agents of such person;

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1 <u>(9) THC means tetrahydrocannabinol;</u>

2 (10) THC concentration refers to the concentration of THC as
3 measured through procedures that use post-decarboxylation or other
4 similarly reliable measures to account for any chemical precursors to
5 cannabinoids, including tetrahydrocannabinolic acid. Such chemical
6 precursors, including tetrahydrocannabinolic acid, shall be included in
7 the total THC concentration measurement;

8 <u>(11)</u> (6) USDA means the United States Department of Agriculture; and 9 <u>(12)</u> (7) USDA-licensed hemp producer means a person licensed by the 10 USDA to produce hemp as provided in 7 C.F.R. part 990, subpart C, as such 11 regulations existed on January 1, 2024, and includes the authorized 12 employees or agents of such person.

Sec. 3. (1) Beginning January 1, 2026, an excise tax shall be
 levied on the retail sale of cannabidiol products to consumers. The tax
 shall be at a rate of ten percent of the retail purchase price.

16 (2) The excise tax imposed by this section shall be in addition to
 17 all other occupation or privilege taxes imposed by this state or by any
 18 political subdivision of the state.

<u>(3) Each retail seller of cannabidiol products shall maintain</u>
 <u>complete and accurate electronic records of sales of cannabidiol</u>
 <u>products, in the manner prescribed by the Department of Revenue. Such</u>
 <u>seller shall provide such records to the department upon request.</u>

(4)(a) Each retail seller of cannabidiol products shall file a
 return with the department by the twentieth day of the month following
 the month reported and with the report shall remit the amount of excise
 tax due.

(b) The return, which shall be upon forms prescribed and furnished by the department, shall contain, among other things, the total amount of cannabidiol products sold or transferred during the preceding month and the amount of tax due thereon.

31 (c) The department may require retail sellers to file tax returns

1 electronically and to remit payments due by electronic funds transfers. 2 (5) The department shall collect the excise tax and shall account 3 for and remit to the State Treasurer at least once each month all money 4 collected pursuant to such tax for credit to the Property Tax Credit Cash 5 Fund. 6 Sec. 4. Section 2-505, Revised Statutes Cumulative Supplement, 2024, 7 is amended to read: 2-505 (1)(a) This subsection applies to hemp other than cannabidiol 8 9 products. (b) Hemp shall not be cultivated, possessed, handled, transported, 10 processed, used, sold, or consumed in this state, except as provided in 11 this subsection. 12 (c) Federally compliant hemp (1) Hemp may be cultivated in this 13 14 state by a USDA-licensed hemp producer, in accordance with such 15 producer's USDA-issued license, or by a state-program-licensed hemp producer, in accordance with such producer's license under a USDA-16 17 approved tribal program. (d) (2) Hemp may only be transported pursuant to section 2-515. 18 19 (2) Any cannabidiol product shall be possessed, handled, 20 transported, used, sold, and consumed in accordance with: 21 (a) The Nebraska Pure Food Act; and 22 (b) The Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 301 et seq., 23 and any regulations adopted and promulgated under such act, as such act 24 and regulations existed on January 1, 2025. 25 (3)(a) For purposes of this subsection: 26 (i) Consumer safe harbor period means the period of time beginning on the effective date of this act and ending on December 31, 2025; and 27 (ii) Illegal hemp means hemp, hemp products, or cannabidiol products 28 29 that do not comply with the THC limits provided in subdivision (5)(a) of 30 section 2-503 or, except as provided in subdivision (3)(d) of this 31 section, that are otherwise not lawful under the Nebraska Hemp Farming

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1 <u>Act.</u>

2 (b) An individual consumer shall not be subject to prosecution for
3 possession of illegal hemp during the consumer safe harbor period.

4 <u>(c) The Attorney General may coordinate with law enforcement</u> 5 agencies to establish appropriate locations where, during the consumer 6 safe harbor period, consumers may drop off illegal hemp for destruction 7 without prosecution for transporting or handling such illegal hemp for 8 destruction.

9 <u>(d) After the consumer safe harbor period expires, an individual may</u> 10 assert as a defense to a prosecution for possession of illegal hemp that 11 he or she purchased the illegal hemp for personal use prior to the 12 effective date of this act. The burden shall be on the individual to 13 prove the elements of this defense by a preponderance of the evidence. 14 This defense shall only apply to conduct that occurred on or before 15 December 31, 2026.

(e) This subsection does not prevent prosecution or provide a
 defense for the cultivation, possession, handling, transportation,
 processing, use, sale, or other distribution of raw materials or products
 with a delta-9 THC concentration of more than 0.3 percent on a dry weight
 basis.

Sec. 5. Section 2-515, Revised Statutes Cumulative Supplement, 2024, is amended to read:

23 2-515 (1) Except as provided in subsection (3) of this section, any 24 USDA-licensed hemp producer or state-program-licensed hemp producer 25 transporting hemp shall carry with the hemp being transported a copy of 26 the USDA license or state program license under which it was cultivated 27 and a copy of the test results pertaining to each lot of hemp being 28 transported.

(2) A USDA-licensed hemp producer or state-program-licensed hemp
 producer under a USDA-approved tribal program cultivating hemp in this
 state shall maintain a record of shipments of hemp shipped from or

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1 received by such producer. Such record shall, for each shipment of hemp,
2 indicate the date of shipment, identify the point of origin and
3 destination, identify the name of the person sending and receiving the
4 shipment, and include the vehicle identification number of the vehicle
5 transporting the hemp.

6 (3) Any USDA-licensed hemp producer or state-program-licensed hemp 7 producer transporting hemp cultivated under such producer's USDA license 8 or state program license shall not be required to carry a copy of the 9 test results relating to such hemp as provided in subsection (1) of this 10 section if such producer carries with the hemp being transported a copy 11 of the applicable USDA license or state program license and is 12 transporting:

(a) Hemp between two registered sites listed on the producer's USDA
or state program license application;

15 (b) Samples of hemp for testing to determine the 16 tetrahydrocannabinol level; or

(c) Live hemp plants to a registered site listed on the producer's
USDA or state program license application prior to cultivating such hemp
plants.

(4) Federally compliant hemp may be transported in interstate
 commerce for any lawful purpose (4) Any person who is carrying or
 transporting hemp who is not a USDA-licensed hemp producer or state program-licensed hemp producer shall only carry or transport hemp if such
 hemp meets the following requirements:

(a) The hemp is carried or transported with a bill of lading stating
the owner of the hemp, the point of origin of the hemp, and the
destination of the hemp; and

(b) The hemp is carried or transported with a copy of the valid USDA
 or state program license under which the hemp was cultivated;

30 (c) The hemp is carried or transported with a copy of the test
31 results pertaining to each lot of hemp being transported; and

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1 <u>(b)</u> (d) The hemp is not unloaded or in any way removed from the 2 vehicle transporting such hemp unless authorized by state or federal law 3 enforcement.

4 (5) No person shall transport or carry hemp in this state
5 concurrently with any other plant material that is not hemp.

6 Sec. 6. (1) In the event of a conflict between the Nebraska Hemp
7 Farming Act and the Nebraska Medical Cannabis Patient Protection Act, the
8 latter shall control.

9 <u>(2) In the event of a conflict between the Nebraska Hemp Farming Act</u> 10 <u>and the Nebraska Medical Cannabis Regulation Act, the latter shall</u> 11 <u>control.</u>

Sec. 7. Section 28-401, Revised Statutes Cumulative Supplement, 2024, is amended to read:

14 28-401 As used in the Uniform Controlled Substances Act, unless the 15 context otherwise requires:

16 (1) Administer means to directly apply a controlled substance by
17 injection, inhalation, ingestion, or any other means to the body of a
18 patient or research subject;

19 (2) Agent means an authorized person who acts on behalf of or at the 20 direction of another person but does not include a common or contract 21 carrier, public warehouse keeper, or employee of a carrier or warehouse 22 keeper;

(3) Administration means the Drug Enforcement Administration of the
 United States Department of Justice;

(4) Controlled substance means a drug, biological, substance, or
immediate precursor in Schedules I through V of section 28-405.
Controlled substance does not include distilled spirits, wine, malt
beverages, tobacco, hemp, or any nonnarcotic substance if such substance
may, under the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 301 et
seq., as such act existed on January 1, 2014, and the law of this state,
be lawfully sold over the counter without a prescription;

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(5) Counterfeit substance means a controlled substance which, or the 1 2 container or labeling of which, without authorization, bears the 3 trademark, trade name, or other identifying mark, imprint, number, or device, or any likeness thereof, of a manufacturer, distributor, or 4 5 dispenser other than the person or persons who in fact manufactured, 6 distributed, or dispensed such substance and which thereby falsely 7 purports or is represented to be the product of, or to have been 8 distributed by, such other manufacturer, distributor, or dispenser;

(6) Department means the Department of Health and Human Services;

10 (7) Division of Drug Control means the personnel of the Nebraska
11 State Patrol who are assigned to enforce the Uniform Controlled
12 Substances Act;

(8) Dispense means to deliver a controlled substance to an ultimate
user or a research subject pursuant to a medical order issued by a
practitioner authorized to prescribe, including the packaging, labeling,
or compounding necessary to prepare the controlled substance for such
delivery;

18 (9) Distribute means to deliver other than by administering or
19 dispensing a controlled substance;

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(10) Prescribe means to issue a medical order;

21 (11) Drug means (a) articles recognized in the official United 22 States Pharmacopoeia, official Homeopathic Pharmacopoeia of the United 23 States, official National Formulary, or any supplement to any of them, 24 (b) substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in human beings or animals, and (c) 25 26 substances intended for use as a component of any article specified in 27 subdivision (a) or (b) of this subdivision, but does not include devices or their components, parts, or accessories; 28

(12) Deliver or delivery means the actual, constructive, or
attempted transfer from one person to another of a controlled substance,
whether or not there is an agency relationship;

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1 (13) Hemp has the same meaning as in section 2-503; 2 (14)(a) Marijuana means all parts of the plant of the genus 3 cannabis, whether growing or not, the seeds thereof, and every compound, manufacture, salt, derivative, mixture, or preparation of such plant or 4 5 its seeds. 6 (b) Marijuana does not include: 7 (i) The the mature stalks of such plant; τ 8 (ii) Hashish; 9 (iii) Tetrahydrocannabinols hashish, tetrahydrocannabinols extracted 10 or isolated from the plant; τ 11 (iv) Fiber fiber produced from such stalks; T (v) Oil oil or cake made from the seeds of such plant; τ 12 (vi) Any any other compound, manufacture, salt, derivative, mixture, 13 14 or preparation of such mature stalks; τ 15 (vii) The the sterilized seed of such plant which is incapable of germination; or , or 16 17 (viii) Cannabidiol cannabidiol contained in a drug product approved by the federal Food and Drug Administration. 18 19 (c) Marijuana includes does not include hemp, except for hemp possessed in compliance with the Nebraska Hemp Farming Act. 20 21 (d) When the weight of marijuana is referred to in the Uniform 22 Controlled Substances Act, it means its weight at or about the time it is 23 seized or otherwise comes into the possession of law enforcement authorities, whether cured or uncured at that time; 24 (15) Manufacture means the production, preparation, propagation, 25 26 conversion, or processing of a controlled substance, either directly or 27 indirectly, extraction from substances of by natural origin, independently by means of chemical synthesis, or by a combination of 28 29 extraction and chemical synthesis, and includes any packaging or 30 repackaging of the substance or labeling or relabeling of its container. Manufacture does not include the preparation or compounding of a 31

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controlled substance by an individual for his or her own use, except for 1 2 the preparation or compounding of components or ingredients used for or 3 intended to be used for the manufacture of methamphetamine, or the compounding, conversion, packaging, or labeling of a 4 preparation, 5 controlled substance: (a) By a practitioner as an incident to his or her 6 prescribing, administering, or dispensing of a controlled substance in 7 the course of his or her professional practice; or (b) by a practitioner, or by his or her authorized agent under his or her supervision, for the 8 9 purpose of, or as an incident to, research, teaching, or chemical analysis and not for sale; 10

11 (16) Narcotic drug means any of the following, whether produced 12 directly or indirectly by extraction from substances of vegetable origin, independently by means of chemical synthesis, or by a combination of 13 14 extraction and chemical synthesis: (a) Opium, opium poppy and poppy 15 straw, coca leaves, and opiates; (b) a compound, manufacture, salt, derivative, or preparation of opium, coca leaves, or opiates; or (c) a 16 17 substance and any compound, manufacture, salt, derivative, or preparation thereof which is chemically equivalent to or identical with any of the 18 substances referred to in subdivisions (a) and (b) of this subdivision, 19 except that the words narcotic drug as used in the Uniform Controlled 20 21 Substances Act does not include decocainized coca leaves or extracts of 22 coca leaves, which extracts do not contain cocaine or ecgonine, or 23 isoquinoline alkaloids of opium;

(17) Opiate means any substance having an addiction-forming or addiction-sustaining liability similar to morphine or being capable of conversion into a drug having such addiction-forming or addictionsustaining liability. Opiate does not include the dextrorotatory isomer of 3-methoxy-n methylmorphinan and its salts. Opiate includes its racemic and levorotatory forms;

30 (18) Opium poppy means the plant of the species Papaver somniferum
31 L., except the seeds thereof;

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(19) Poppy straw means all parts, except the seeds, of the opium
 poppy after mowing;

3 (20) Person means any corporation, association, partnership, limited
4 liability company, or one or more persons;

5 (21) Practitioner means a physician, a physician assistant, a 6 dentist, a veterinarian, a pharmacist, a podiatrist, an optometrist, a 7 certified nurse midwife, a certified registered nurse anesthetist, a nurse practitioner, a scientific investigator, a pharmacy, a hospital, or 8 9 any other person licensed, registered, or otherwise permitted to distribute, dispense, prescribe, conduct research with respect to, or 10 11 administer a controlled substance in the course of practice or research 12 in this state, including an emergency medical service as defined in section 38-1207; 13

14 (22) Production includes the manufacture, planting, cultivation, or
 15 harvesting of a controlled substance;

16 (23) Immediate precursor means a substance which is the principal 17 compound commonly used or produced primarily for use and which is an 18 immediate chemical intermediary used or likely to be used in the 19 manufacture of a controlled substance, the control of which is necessary 20 to prevent, curtail, or limit such manufacture;

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(24) State means the State of Nebraska;

(25) Ultimate user means a person who lawfully possesses a controlled substance for his or her own use, for the use of a member of his or her household, or for administration to an animal owned by him or her or by a member of his or her household;

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(26) Hospital has the same meaning as in section 71-419;

27 (27) Cooperating individual means any person, other than a 28 commissioned law enforcement officer, who acts on behalf of, at the 29 request of, or as agent for a law enforcement agency for the purpose of 30 gathering or obtaining evidence of offenses punishable under the Uniform 31 Controlled Substances Act;

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(28) Cannabid

(28) Cannabidiol product has the same meaning as in section 2-503;

(29)(a) (28)(a) Hashish or concentrated cannabis means (i) the
 separated resin, whether crude or purified, obtained from a plant of the
 genus cannabis or (ii) any material, preparation, mixture, compound, or
 other substance which contains ten percent or more by weight of
 tetrahydrocannabinols.

7 (b) When resins extracted from hemp as defined in section 2-503 are 8 in the possession of a person as authorized under the Nebraska Hemp 9 Farming Act, they are not considered hashish or concentrated cannabis for 10 purposes of the Uniform Controlled Substances Act.

(c) Hashish or concentrated cannabis does not include <u>any</u>
 <u>cannabidiol product or cannabidiol contained in a drug product approved</u>
 by the federal Food and Drug Administration;

(30) (29) Exceptionally hazardous drug means (a) a narcotic drug,
 (b) thiophene analog of phencyclidine, (c) phencyclidine, (d)
 amobarbital, (e) secobarbital, (f) pentobarbital, (g) amphetamine, or (h)
 methamphetamine;

(31) (30) Imitation controlled substance means a substance which is 18 not a controlled substance or controlled substance analogue but which, by 19 20 way of express or implied representations and consideration of other 21 relevant factors including those specified in section 28-445, would lead 22 a reasonable person to believe the substance is a controlled substance or 23 controlled substance analogue. A placebo or registered investigational 24 drug manufactured, distributed, possessed, or delivered in the ordinary course of practice or research by a health care professional shall not be 25 26 deemed to be an imitation controlled substance;

(32)(a) (31)(a) Controlled substance analogue means a substance (i)
 the chemical structure of which is substantially similar to the chemical
 structure of a Schedule I or Schedule II controlled substance as provided
 in section 28-405 or (ii) which has a stimulant, depressant, analgesic,
 or hallucinogenic effect on the central nervous system that is

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substantially similar to or greater than the stimulant, depressant, analgesic, or hallucinogenic effect on the central nervous system of a Schedule I or Schedule II controlled substance as provided in section 28-405. A controlled substance analogue shall, to the extent intended for human consumption, be treated as a controlled substance under Schedule I of section 28-405 for purposes of the Uniform Controlled Substances Act; and

8 (b) Controlled substance analogue does not include (i) a controlled 9 substance, (ii) any substance generally recognized as safe and effective within the meaning of the Federal Food, Drug, and Cosmetic Act, 21 U.S.C. 10 11 301 et seq., as such act existed on January 1, 2014, (iii) any substance for which there is an approved new drug application, or (iv) with respect 12 to a particular person, any substance if an exemption is in effect for 13 14 investigational use for that person, under section 505 of the Federal 15 Food, Drug, and Cosmetic Act, 21 U.S.C. 355, as such section existed on January 1, 2014, to the extent conduct with respect to such substance is 16 pursuant to such exemption; 17

(33) (32) Anabolic steroid means any drug or hormonal substance, 18 chemically and pharmacologically related to testosterone (other than 19 estrogens, progestins, and corticosteroids), that promotes muscle growth 20 21 and includes any controlled substance in Schedule III(d) of section 22 28-405. Anabolic steroid does not include any anabolic steroid which is 23 expressly intended for administration through implants to cattle or other 24 nonhuman species and has been approved by the Secretary of Health and Human Services for such administration, but if any person prescribes, 25 26 dispenses, or distributes such a steroid for human use, such person shall 27 be considered to have prescribed, dispensed, or distributed an anabolic steroid within the meaning of this subdivision; 28

29 <u>(34)</u> (33) Chart order means an order for a controlled substance 30 issued by a practitioner for a patient who is in the hospital where the 31 chart is stored or for a patient receiving detoxification treatment or

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1 maintenance treatment pursuant to section 28-412. Chart order does not 2 include a prescription;

3 (35) (34) Medical order means a prescription, a chart order, or an
 4 order for pharmaceutical care issued by a practitioner;

5 (36) (35) Prescription means an order for a controlled substance
6 issued by a practitioner. Prescription does not include a chart order;

7 (37) (36) Registrant means any person who has a controlled
8 substances registration issued by the state or the Drug Enforcement
9 Administration of the United States Department of Justice;

10 <u>(38)</u> (37) Reverse distributor means a person whose primary function 11 is to act as an agent for a pharmacy, wholesaler, manufacturer, or other 12 entity by receiving, inventorying, and managing the disposition of 13 outdated, expired, or otherwise nonsaleable controlled substances;

14 (39) (38) Signature means the name, word, or mark of a person 15 written in his or her own hand with the intent to authenticate a writing 16 or other form of communication or a digital signature which complies with 17 section 86-611 or an electronic signature;

(40) (39) Facsimile means a copy generated by a system that encodes
 a document or photograph into electrical signals, transmits those signals
 over telecommunications lines, and reconstructs the signals to create an
 exact duplicate of the original document at the receiving end;

22 (41) (40) Electronic signature has the definition found in section 23 86-621;

(42) (41) Electronic transmission means transmission of information
 in electronic form. Electronic transmission includes computer-to-computer
 transmission or computer-to-facsimile transmission;

27 (43) (42) Long-term care facility means an intermediate care
28 facility, an intermediate care facility for persons with developmental
29 disabilities, a long-term care hospital, a mental health substance use
30 treatment center, a nursing facility, or a skilled nursing facility, as
31 such terms are defined in the Health Care Facility Licensure Act;

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1 (44) (43) Compounding has the same meaning as in section 38-2811;

2 <u>(45)</u> (44) Cannabinoid receptor agonist means any chemical compound 3 or substance that, according to scientific or medical research, study, 4 testing, or analysis, demonstrates the presence of binding activity at 5 one or more of the CB1 or CB2 cell membrane receptors located within the 6 human body. Cannabinoid receptor agonist does not include <u>any cannabidiol</u> 7 <u>product or cannabidiol</u> contained in a drug product approved by the 8 federal Food and Drug Administration; and

9 <u>(46)</u> (45) Lookalike substance means a product or substance, not 10 specifically designated as a controlled substance in section 28-405, that 11 is either portrayed in such a manner by a person to lead another person 12 to reasonably believe that it produces effects on the human body that 13 replicate, mimic, or are intended to simulate the effects produced by a 14 controlled substance or that possesses one or more of the following 15 indicia or characteristics:

(a) The packaging or labeling of the product or substance suggests
that the user will achieve euphoria, hallucination, mood enhancement,
stimulation, or another effect on the human body that replicates or
mimics those produced by a controlled substance;

(b) The name or packaging of the product or substance uses images or
labels suggesting that it is a controlled substance or produces effects
on the human body that replicate or mimic those produced by a controlled
substance;

(c) The product or substance is marketed or advertised for a
particular use or purpose and the cost of the product or substance is
disproportionately higher than other products or substances marketed or
advertised for the same or similar use or purpose;

(d) The packaging or label on the product or substance contains
words or markings that state or suggest that the product or substance is
in compliance with state and federal laws regulating controlled
substances;

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(e) The owner or person in control of the product or substance uses
 evasive tactics or actions to avoid detection or inspection of the
 product or substance by law enforcement authorities;

4 (f) The owner or person in control of the product or substance makes 5 a verbal or written statement suggesting or implying that the product or 6 substance is a synthetic drug or that consumption of the product or 7 substance will replicate or mimic effects on the human body to those 8 effects commonly produced through use or consumption of a controlled 9 substance;

10 (g) The owner or person in control of the product or substance makes 11 a verbal or written statement to a prospective customer, buyer, or 12 recipient of the product or substance implying that the product or 13 substance may be resold for profit; or

(h) The product or substance contains a chemical or chemical compound that does not have a legitimate relationship to the use or purpose claimed by the seller, distributor, packer, or manufacturer of the product or substance or indicated by the product name, appearing on the product's packaging or label or depicted in advertisement of the product or substance.

Sec. 8. Original sections 2-501, 2-503, 2-505, 2-515, and 28-401,
Revised Statutes Cumulative Supplement, 2024, are repealed.

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