

General Assembly

Amendment

January Session, 2023

LCO No. **7520**



Offered by:

REP. ANDERSON, 62nd Dist.

To: Subst. House Bill No. 6699

File No. 201

Cal. No. 150

(As Amended)

"AN ACT CONCERNING CANNABIS REGULATION."

- 1 Strike section 1 in its entirety and substitute the following in lieu
- 2 thereof:
- 3 "Section 1. Section 21a-240 of the general statutes is repealed and the
- 4 following is substituted in lieu thereof (*Effective July 1, 2023*):
- 5 The following words and phrases, as used in this chapter, shall have
- 6 the following meanings, unless the context otherwise requires:
- 7 (1) "Abuse of drugs" means the use of controlled substances solely for
- 8 their stimulant, depressant or hallucinogenic effect upon the higher
- 9 functions of the central nervous system and not as a therapeutic agent
- 10 prescribed in the course of medical treatment or in a program of
- 11 research operated under the direction of a physician or pharmacologist.
- 12 [;]
- 13 (2) "Administer" means the direct application of a controlled

substance, whether by injection, inhalation, ingestion or any other means, to the body of a patient or research subject by: (A) A practitioner, or, in [his] the practitioner's presence, by [his] the practitioner's authorized agent, or (B) the patient or research subject at the direction and in the presence of the practitioner, or (C) a nurse or intern under the direction and supervision of a practitioner. [;]

- (3) "Agent" means an authorized person who acts on behalf of or at the direction of a manufacturer, distributor, dispenser or prescribing practitioner. [. It] <u>but</u> does not include a common or contract carrier, public warehouseman, or employee of the carrier or warehouseman. [;]
- (4) "Amphetamine-type substances" include amphetamine, optical isomers thereof, salts of amphetamine and its isomers, and chemical compounds which are similar thereto in chemical structure or which are similar thereto in physiological effect, and which show a like potential for abuse, which are controlled substances under this chapter unless modified. [;]
- (5) "Barbiturate-type drugs" include barbituric acid and its salts, derivatives thereof and chemical compounds which are similar thereto in chemical structure or which are similar thereto in physiological effect, and which show a like potential for abuse, which are controlled substances under this chapter unless modified. [;]
- (6) "Bureau" means the Bureau of Narcotics and Dangerous Drugs, United States Department of Justice, or its successor agency. [;]
- (7) "Cannabis-type substances" include all parts of any plant, or species of the genus cannabis or any infra specific taxon thereof whether growing or not; the seeds thereof; the resin extracted from any part of such a plant; and every compound, manufacture, salt, derivative, mixture or preparation of such plant, its seeds or resin; but shall not include the mature stalks of such plant, fiber produced from such stalks, oil or cake made from the seeds of such plant, any other compound, manufacture, salt, derivative, mixture or preparation of such mature stalks, except the resin extracted therefrom, fiber, oil or cake, the

sterilized seed of such plant which is incapable of germination, or hemp, as defined in 7 USC 16390, as amended from time to time. Included are cannabinon, cannabinol, cannabidiol and chemical compounds which are similar to cannabinon, cannabinol or cannabidiol in chemical structure or which are similar thereto in physiological effect, and which show a like potential for abuse, which are controlled substances under this chapter unless derived from hemp, as defined in section 22-61*l*, as amended by this act. [;]

- (8) "Controlled drugs" are those drugs which contain any quantity of a substance which has been designated as subject to the federal Controlled Substances Act, or which has been designated as a depressant or stimulant drug pursuant to federal food and drug laws, or which has been designated by the Commissioner of Consumer Protection pursuant to section 21a-243, as having a stimulant, depressant or hallucinogenic effect upon the higher functions of the central nervous system and as having a tendency to promote abuse or psychological or physiological dependence, or both. Such controlled drugs are classifiable as amphetamine-type, barbiturate-type, cannabistype, cocaine-type, hallucinogenic, morphine-type and other stimulant and depressant drugs. Specifically excluded from controlled drugs and controlled substances are alcohol, nicotine and caffeine. [;]
- (9) "Controlled substance" means a drug, substance, or immediate precursor in schedules I to V, inclusive, of the Connecticut controlled substance scheduling regulations adopted pursuant to section 21a-243.

 [;]
- (10) "Counterfeit substance" means a controlled substance which, or the container or labeling of which, without authorization, bears the trademark, trade name or other identifying mark, imprint, number or device, or any likeness thereof, of a manufacturer, distributor or dispenser other than the person who in fact manufactured, distributed or dispensed the substance. [;]
- 77 (11) "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. [;]

- 80 (12) "Dentist" means a person authorized by law to practice dentistry 81 in this state. [;]
- (13) "Dispense" means to deliver a controlled substance to an ultimate user or research subject by or pursuant to the lawful order of a practitioner, including the prescribing, administering, packaging, labeling or compounding necessary to prepare the substance for the delivery. [;]
- 87 (14) "Dispenser" means a practitioner who dispenses. [;]
- 88 (15) "Distribute" means to deliver other than by administering or dispensing a controlled substance. [;]
- (16) "Distributor" means a person who distributes and includes a wholesaler who is a person supplying or distributing controlled drugs which [he himself] the person personally has not produced or prepared to hospitals, clinics, practitioners, pharmacies, other wholesalers, manufacturers and federal, state and municipal agencies. [;]
 - (17) "Drug" means (A) substances recognized as drugs in the official United States Pharmacopoeia, official Homeopathic Pharmacopoeia of the United States, or official National Formulary, or any supplement to any of them; (B) substances intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in man or animals; (C) substances, other than food, intended to affect the structure or any function of the body of man or animals; and (D) substances intended for use as a component of any article specified in subparagraph (A), (B) or (C) of this subdivision. It does not include devices or their components, parts or accessories. [;]
- (18) "Drug dependence" means a psychoactive substance dependence
 on drugs as that condition is defined in the most recent edition of the
 "Diagnostic and Statistical Manual of Mental Disorders" of the American

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Psychiatric Association. [;]

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109 (19) "Drug-dependent person" means a person who has a 110 psychoactive substance dependence on drugs as that condition is 111 defined in the most recent edition of the "Diagnostic and Statistical 112 Manual of Mental Disorders" of the American Psychiatric Association. 113 [;]

(20) (A) "Drug paraphernalia" means equipment, products and materials of any kind that are used, intended for use or designed for use planting, propagating, cultivating, growing, harvesting, manufacturing, compounding, converting, producing, processing, testing, analyzing, packaging, repackaging, containing or concealing, or ingesting, inhaling or otherwise introducing into the human body, any controlled substance contrary to the provisions of this chapter, including, but not limited to: (i) Kits intended for use or designed for use in planting, propagating, cultivating, growing or harvesting of any species of plant that is a controlled substance or from which a controlled substance can be derived; (ii) kits used, intended for use or designed for use in manufacturing, compounding, converting, producing, processing or preparing controlled substances; (iii) isomerization devices used or intended for use in increasing the potency of any species of plant that is a controlled substance; (iv) testing equipment used, intended for use or designed for use in identifying or analyzing the strength, effectiveness or purity of controlled substances; (v) dilutents and adulterants, including, but not limited to, quinine hydrochloride, mannitol, mannite, dextrose and lactose used, intended for use or designed for use in cutting controlled substances; (vi) separation gins and sifters used, intended for use or designed for use in removing twigs and seeds from, or in otherwise cleaning or refining, marijuana; (vii) capsules and other containers used, intended for use or designed for use in packaging small quantities of controlled substances; (viii) containers and other objects used, intended for use or designed for use in storing or concealing controlled substances; and (ix) objects used, intended for use or designed for use in ingesting, inhaling, or otherwise introducing

142 marijuana, cocaine, hashish, or hashish oil into the human body, 143 including, but not limited to, wooden, acrylic, glass, stone, plastic or 144 ceramic pipes with screens, permanent screens, hashish heads or 145 punctured metal bowls; water pipes; carburetion tubes and devices; 146 smoking and carburetion masks; roach clips; miniature cocaine spoons 147 and cocaine vials; chamber pipes; carburetor pipes; electric pipes; air-148 driven pipes; chillums; bongs; ice pipes and chillers. "Drug 149 paraphernalia" does not include a product used by a manufacturer 150 licensed pursuant to this chapter for the activities permitted under the 151 license or by an individual to test any substance prior to injection, 152 inhalation or ingestion of the substance to prevent accidental overdose 153 by injection, inhalation or ingestion of the substance, provided the 154 licensed manufacturer or individual is not using the product to engage 155 in the unlicensed manufacturing or distribution of controlled 156 substances. As used in this subdivision, "roach clip" means an object 157 used to hold burning material, including, but not limited to, a marijuana 158 cigarette, that has become too small or too short to be held between the 159 fingers. [;]

- 160 (B) "Factory" means any place used for the manufacturing, mixing, compounding, refining, processing, packaging, distributing, storing, keeping, holding, administering or assembling illegal substances contrary to the provisions of this chapter, or any building, rooms or location which contains equipment or paraphernalia used for this purpose. [;]
- (21) "Federal Controlled Substances Act, 21 USC 801 et seq." means 166 167 Public Law 91-513, the Comprehensive Drug Abuse Prevention and 168 Control Act of 1970. [;]
- 169 (22) "Federal food and drug laws" means the federal Food, Drug and 170 Cosmetic Act, as amended, Title 21 USC 301 et seq. [;]
- 171 (23) "Hallucinogenic substances" are psychodysleptic substances, 172 other than cannabis-type substances, which assert a confusional or 173 disorganizing effect upon mental processes or behavior and mimic

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acute psychotic disturbances. Exemplary of such drugs are mescaline, peyote, psilocyn and d-lysergic acid diethylamide, which are controlled substances under this chapter unless modified. [;]

- (24) "Hospital", as used in sections 21a-243 to 21a-283, inclusive, means an institution for the care and treatment of the sick and injured, approved by the Department of Public Health or the Department of Mental Health and Addiction Services as proper to be entrusted with the custody of controlled drugs and substances and professional use of controlled drugs and substances under the direction of a licensed practitioner. [;]
- (25) "Intern" means a person who holds a degree of doctor of medicine or doctor of dental surgery or medicine and whose period of service has been recorded with the Department of Public Health and who has been accepted and is participating in training by a hospital or institution in this state. Doctors meeting the foregoing requirements and commonly designated as "residents" and "fellows" shall be regarded as interns for purposes of this chapter. [;]
 - (26) "Immediate precursor" means a substance which the Commissioner of Consumer Protection has found to be, and by regulation designates as being, the principal compound commonly used or produced primarily for use, and which is an immediate chemical intermediary used or likely to be used, in the manufacture of a controlled substance, the control of which is necessary to prevent, curtail or limit manufacture. [;]
 - (27) "Laboratory" means a laboratory approved by the Department of Consumer Protection as proper to be entrusted with the custody of controlled substances and the use of controlled substances for scientific and medical purposes and for purposes of instruction, research or analysis. [;]
- (28) "Manufacture" means the production, preparation, cultivation, growing, propagation, compounding, conversion or processing of a controlled substance, either directly or indirectly by extraction from

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substances of natural origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis, and includes any packaging or repackaging of the substance or labeling or relabeling of its container, except that this term does not include the preparation or compounding of a controlled substance by an individual for [his] the individual's own use or the preparation, compounding, packaging or labeling of a controlled substance: (A) By a practitioner as an incident to [his] the practitioner administering or dispensing of a controlled substance in the course of [his] such practitioner's professional practice, or (B) by a practitioner, or by [his] the practitioner's authorized agent under [his] such practitioner's supervision, for the purpose of, or as an incident to, research, teaching or chemical analysis and not for sale. [;]

(29) "Marijuana" means all parts of any plant, or species of the genus cannabis or any infra specific taxon thereof, whether growing or not; the seeds thereof; the resin extracted from any part of the plant; every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds or resin, any [product made using hemp, as defined in section 22-61l, which exceeds three-tenths per cent total THC concentration on a dry-weight basis | high-THC hemp product; manufactured cannabinoids, synthetic cannabinoids, except as provided in subparagraph (E) of this subdivision; or cannabinon, cannabinol or cannabidiol and chemical compounds which are similar to cannabinon, cannabinol or cannabidiol in chemical structure or which are similar thereto in physiological effect, which are controlled substances under this chapter, except cannabidiol derived from hemp, as defined in section 22-61l, as amended by this act, [with a total THC concentration of not more than three-tenths per cent on a dry-weight basis] that is not a high-THC hemp product. "Marijuana" does not include: (A) The mature stalks of such plant, fiber produced from such stalks, oil or cake made from the seeds of such plant, any other compound, manufacture, salt, derivative, mixture or preparation of such mature stalks, except the resin extracted from such mature stalks or fiber, oil or cake; (B) the sterilized seed of such plant which is

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240 incapable of germination; (C) hemp, as defined in section 22-61l, as 241 amended by this act, (i) with a total THC concentration of not more than 242 three-tenths per cent on a dry-weight basis, and (ii) that is not a high-243 THC hemp product; (D) any substance approved by the federal Food 244 and Drug Administration or successor agency as a drug and reclassified 245 in any schedule of controlled substances or unscheduled by the federal 246 Drug Enforcement Administration or successor agency which is 247 included in the same schedule designated by the federal Drug 248 Enforcement Administration or successor agency; or (E) synthetic 249 cannabinoids which are controlled substances that are designated by the 250 Commissioner of Consumer Protection, by whatever official, common, 251 usual, chemical or trade name designation, as controlled substances and 252 are classified in the appropriate schedule in accordance with 253 subsections (i) and (j) of section 21a-243. [;]

(30) "Narcotic substance" means any of the following, whether produced directly or indirectly by extraction from a substance of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis: (A) Morphinetype: (i) Opium or opiate, or any salt, compound, derivative, or preparation of opium or opiate which is similar to any such substance in chemical structure or which is similar to any such substance in physiological effect and which shows a like potential for abuse, which is a controlled substance under this chapter unless modified; (ii) any salt, compound, isomer, derivative, or preparation of any such substance which is chemically equivalent or identical to any substance referred to in clause (i) of this subdivision, but not including the isoquinoline alkaloids of opium; (iii) opium poppy or poppy straw; or (iv) (I) fentanyl or any salt, compound, derivative or preparation of fentanyl which is similar to any such substance in chemical structure or which is similar to any such substance in physiological effect and which shows a like potential for abuse, which is a controlled substance under this chapter unless modified, or (II) any salt, compound, isomer, derivative or preparation of any such substance which is chemically equivalent or identical to any substance referred to in subclause (I) of

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274 this clause; or (B) cocaine-type; coca leaves or any salt, compound, 275 derivative or preparation of coca leaves, or any salt, compound, isomer, 276 derivatives or preparation of any such substance which is chemically 277 equivalent or identical to any such substance or which is similar to any 278 such substance in physiological effect and which shows a like potential 279 for abuse, but not including decocainized coca leaves or extractions of 280 coca leaves which do not contain cocaine or ecgonine. [;]

- 281 (31) "Nurse" means a person performing nursing as defined in section 282 20-87a. [;]
- 283 (32) "Official written order" means an order for controlled substances 284 written on a form provided by the bureau for that purpose under the 285 federal Controlled Substances Act. [;]
 - (33) "Opiate" means any substance having an addiction-forming or addiction-sustaining liability similar to morphine or being capable of conversion into a drug having addiction-forming or addictionsustaining liability; it does not include, unless specifically designated as controlled under this chapter, the dextrorotatory isomer of 3-methoxyn-methylmorthinan and its salts (dextro-methorphan) but shall include its racemic and levorotatory forms. [;]
 - (34) "Opium poppy" means the plant of the species papaver somniferum l., except its seed. [;]
- 295 (35) Repealed by P.A. 99-102, S. 51. [;]
- 296 (36) "Other stimulant and depressant drugs" means controlled 297 substances other than amphetamine-type, barbiturate-type, cannabis-298 type, cocaine-type, hallucinogenics and morphine-type which are found 299 to exert a stimulant and depressant effect upon the higher functions of 300 the central nervous system and which are found to have a potential for abuse and are controlled substances under this chapter. [;]
- 302 (37) "Person" includes any corporation, limited liability company, 303 association or partnership, or one or more individuals, government or

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governmental subdivisions or agency, business trust, estate, trust, or any other legal entity. Words importing the plural number may include the singular; words importing the masculine gender may be applied to females. [;]

- 308 (38) "Pharmacist" means a person authorized by law to practice pharmacy pursuant to section 20-590, 20-591, 20-592 or 20-593. [;]
- 310 (39) "Pharmacy" means an establishment licensed pursuant to section 311 20-594. [;]
- 312 (40) "Physician" means a person authorized by law to practice 313 medicine in this state pursuant to section 20-9. [;]
- (41) "Podiatrist" means a person authorized by law to practice podiatry in this state. [;]
- 316 (42) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing. [;]
- 318 (43) "Practitioner" means: (A) A physician, dentist, veterinarian, 319 podiatrist, scientific investigator or other person licensed, registered or 320 otherwise permitted to distribute, dispense, conduct research with 321 respect to or to administer a controlled substance in the course of 322 professional practice or research in this state; and (B) a pharmacy, 323 hospital or other institution licensed, registered or otherwise permitted 324 to distribute, dispense, conduct research with respect to or to administer 325 a controlled substance in the course of professional practice or research 326 in this state. [;]
- 327 (44) "Prescribe" means order or designate a remedy or any 328 preparation containing controlled substances. [;]
- 329 (45) "Prescription" means a written, oral or electronic order for any 330 controlled substance or preparation from a licensed practitioner to a 331 pharmacist for a patient. [;]
- 332 (46) "Production" includes the manufacture, planting, cultivation,

growing or harvesting of a controlled substance. [;]

334 (47) "Registrant" means any person licensed by this state and 335 assigned a current federal Bureau of Narcotics and Dangerous Drug 336 Registry Number as provided under the federal Controlled Substances 337 Act. [;]

- (48) "Registry number" means the alphabetical or numerical designation of identification assigned to a person by the federal Drug Enforcement Administration, or other federal agency, which is commonly known as the federal registry number. [;]
- (49) "Restricted drugs or substances" are the following substances without limitation and for all purposes: Datura stramonium; hyoscyamus niger; atropa belladonna, or the alkaloids atropine; hyoscyamine; belladonnine; apatropine; or any mixture of these alkaloids such as daturine, or the synthetic homatropine or any salts of these alkaloids, except that any drug or preparation containing any of the above-mentioned substances which is permitted by federal food and drug laws to be sold or dispensed without a prescription or written order shall not be a controlled substance; amyl nitrite; the following volatile substances to the extent that said chemical substances or compounds containing said chemical substances are sold, prescribed, dispensed, compounded, possessed or controlled or delivered or administered to another person with the purpose that said chemical substances shall be breathed, inhaled, sniffed or drunk to induce a stimulant, depressant or hallucinogenic effect upon the higher functions of the central nervous system: Acetone; benzene; butyl alcohol; butyl nitrate and its salts, isomers, esters, ethers or their salts; cyclohexanone; dichlorodifluoromethane; ether; ethyl acetate; formaldehyde; hexane; isopropanol; methanol; methyl cellosolve acetate; methyl ethyl ketone; methyl isobutyl ketone; nitrous oxide; pentochlorophenol; toluene; toluol; trichloroethane; trichloroethylene; 1,4 butanediol. [;]
- 363 (50) "Sale" is any form of delivery which includes barter, exchange or 364 gift, or offer therefor, and each such transaction made by any person

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365 whether as principal, proprietor, agent, servant or employee. [;]

366 (51) "State", when applied to a part of the United States, includes any 367 state, district, commonwealth, territory or insular possession thereof, 368 and any area subject to the legal authority of the United States of 369 America. [;]

- 370 (52) "State food, drug and cosmetic laws" means the Uniform Food, 371 Drug and Cosmetic Act, section 21a-91 et seq. [;]
- 372 (53) "Ultimate user" means a person who lawfully possesses a 373 controlled substance for [his] the person's own use or for the use of a 374 member of [his] such person's household or for administering to an 375 animal owned by [him] such person or by a member of [his] such 376 person's household. [;]
- 377 (54) "Veterinarian" means a person authorized by law to practice veterinary medicine in this state. [;]
- 379 (55) "Wholesaler" means a distributor or a person who supplies 380 controlled substances that [he himself] the person personally has not 381 produced or prepared to registrants. [as defined in subdivision (47) of 382 this section;]
- 383 (56) "Reasonable times" means the time or times any office, care-384 giving institution, pharmacy, clinic, wholesaler, manufacturer, 385 laboratory, warehouse, establishment, store or place of business, vehicle 386 or other place is open for the normal affairs or business or the practice 387 activities usually conducted by the registrant. [;]
 - (57) "Unit dose drug distribution system" means a drug distribution system used in a hospital or chronic and convalescent nursing home in which drugs are supplied in individually labeled unit of use packages, each patient's supply of drugs is exchanged between the hospital pharmacy and the drug administration area or, in the case of a chronic and convalescent nursing home between a pharmacy and the drug administration area, at least once each twenty-four hours and each

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patient's medication supply for this period is stored within a patientspecific container, all of which is conducted under the direction of a pharmacist licensed in Connecticut and, in the case of a hospital, directly involved in the provision and supervision of pharmaceutical services at such hospital at least thirty-five hours each week. [;]

- (58) "Cocaine in a free-base form" means any substance which contains cocaine, or any compound, isomer, derivative or preparation thereof, in a nonsalt form.
- 403 (59) "THC" means tetrahydrocannabinol, including, but not limited 404 to, delta-7, delta-8-tetrahydrocannabinol, delta-9-tetrahydrocannabinol 405 and delta-10-tetrahydrocannabinol, and any material, compound, 406 mixture or preparation which contain their salts, isomers and salts of 407 isomers, whenever the existence of such salts, isomers and salts of 408 isomers is possible within the specific chemical designation, regardless 409 of the source, except: (A) Dronabinol substituted in sesame oil and 410 encapsulated in a soft gelatin capsule in a federal Food and Drug 411 Administration or successor agency approved product, or (B) any 412 tetrahydrocannabinol product that has been approved by the federal 413 Food and Drug Administration or successor agency to have a medical 414 use and reclassified in any schedule of controlled substances or 415 unscheduled by the federal Drug Enforcement Administration or 416 successor agency.
 - (60) "Total THC" means the sum of the percentage by weight of tetrahydrocannabinolic acid, multiplied by eight hundred seventy-seven-thousandths, plus the percentage of weight of [tetrahydrocannabinol] <u>THC</u>.
 - (61) "Manufactured cannabinoid" means cannabinoids naturally occurring from a source other than marijuana that are similar in chemical structure or physiological effect to cannabinoids derived from marijuana, as defined in section 21a-243, but are derived by a chemical or biological process.
- 426 (62) "Synthetic cannabinoid" means any material, compound, mixture

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or preparation which contains any quantity of a substance having a psychotropic response primarily by agonist activity at cannabinoidspecific receptors affecting the central nervous system that is produced artificially and not derived from an organic source naturally containing cannabinoids, unless listed in another schedule pursuant to section 21a-

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- (63) (A) "High-THC hemp product" means a manufacturer hemp product, as defined in section 22-61*l*, as amended by this act, that has, or is advertised, labeled or offered for sale as having, total THC that exceeds (i) for a hemp edible, hemp topical or hemp transdermal patch (I) one milligram on a per-serving basis, or (II) five milligrams on a percontainer basis, (ii) for a hemp tincture, including, but not limited to, oil intended for ingestion by swallowing, buccal administration or sublingual absorption (I) one milligram on a per-serving basis, or (II) twenty-five milligrams on a per-container basis, (iii) for a hemp concentrate or extract, including, but not limited to, a vape oil, wax or shatter, twenty-five milligrams on a per-container basis, or (iv) for a manufacturer hemp product not described in clause (i), (ii) or (iii) of this subparagraph, (I) one milligram on a per-serving basis, (II) five milligrams on a per-container basis, or (III) three-tenths per cent on a dry-weight basis for cannabis flower or cannabis trim.
- (B) "High-THC hemp product" does not include any manufacturer hemp product, as defined in section 22-61*l*, as amended by this act, that (i) is a full-spectrum CBD product, (ii) has a ratio of THC to CBD that is less than one to twenty-five, and (iii) has total THC that does not exceed three-tenths per cent."
- After the last section, add the following and renumber sections and internal references accordingly:
- "Sec. 501. (NEW) (*Effective July 1, 2023*) No person shall sell any manufacturer hemp product, as defined in section 22-61*l* of the general statutes, as amended by this act, to any individual who is younger than twenty-one years of age."

This act shall take effect as follows and shall amend the following sections:

Section 1	July 1, 2023	21a-240
Sec. 501	July 1, 2023	New section