23101465D

HOUSE BILL NO. 2294

Offered January 11, 2023
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A BILL to amend and reenact §§ 3.2-4113, 3.2-4114, 3.2-4118, 4.1-600, 9.1-1101, 18.2-247, 19.2-188.1, 54.1-3401, 54.1-3408.3, 54.1-3442.6, 54.1-3442.7, 59.1-200, and 59.1-206 of the Code of Virginia, relating to marijuana; tetrahydrocannabinol; hemp products; civil penalty.

Patrons—Kilgore, Leftwich and Rasoul

Referred to Committee for Courts of Justice

Be it enacted by the General Assembly of Virginia:

1. That §§ 3.2-4113, 3.2-4114, 3.2-4118, 4.1-600, 9.1-1101, 18.2-247, 19.2-188.1, 54.1-3401, 54.1-3408.3, 54.1-3442.6, 54.1-3442.7, 59.1-200, and 59.1-206 of the Code of Virginia are amended and reenacted as follows:

§ 3.2-4113. Production of industrial hemp lawful.

A. It is lawful for a grower, his agent, or a federally licensed hemp producer to grow, a dealer or his agent to deal in, or a processor or his agent to process industrial hemp in the Commonwealth for any lawful purpose. No federally licensed hemp producer or grower or his agent shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-247, 18.2-248, 18.2-248.01, 18.2-248.1, or 18.2-250 for the possession or growing of industrial hemp or any Cannabis sativa with a tetrahydrocannabinol concentration that does not exceed the total ~~delta-9~~ tetrahydrocannabinol concentration percentage established in federal regulations applicable to negligent violations located at 7 C.F.R. 990.6(b)(3). No dealer or his agent or processor or his agent shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-247, 18.2-248, 18.2-248.01, 18.2-248.1, or 18.2-250 or issued a summons or judgment for the possession, dealing, or processing of industrial hemp. In any complaint, information, or indictment, and in any action or proceeding brought for the enforcement of any provision of *Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1*, Article 1 (§ 18.2-247 et seq.) of Chapter 7 of Title 18.2, or the Drug Control Act (§ 54.1-3400 et seq.), it shall not be necessary to negate any exception, excuse, proviso, or exemption contained in this chapter or the Drug Control Act, and the burden of proof of any such exception, excuse, proviso, or exemption shall be on the defendant.

B. Nothing in this chapter shall be construed to authorize any person to violate any federal law or regulation.

C. No person shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-247, 18.2-248, 18.2-248.01, 18.2-248.1, or 18.2-250 for the involuntary growth of industrial hemp through the inadvertent natural spread of seeds or pollen as a result of proximity to a production field, dealership, or process site.

§ 3.2-4114. Regulations.

A. The Board may adopt regulations pursuant to this chapter as necessary to register persons to grow, deal in, or process industrial hemp or implement the provisions of this chapter. *Such regulations shall require that hemp products not intended for human consumption, orally or by inhalation, include a bittering agent that renders the product unpalatable.*

B. Upon publication by the U.S. Department of Agriculture in the Federal Register of any final rule regarding industrial hemp that materially expands opportunities for growing, producing, or dealing in industrial hemp in the Commonwealth, the Board shall immediately adopt amendments conforming Department regulations to such federal final rule. Such adoption of regulations by the Board shall be exempt from the provisions of the Administrative Process Act (§ 2.2-4000 et seq.).

§ 3.2-4118. Forfeiture of industrial hemp grower, dealer, or processor registration; violations.

A. The Commissioner shall deny the application, or suspend or revoke the registration, of any person who, with a culpable mental state greater than negligence, violates any provision of this chapter. The Commissioner shall provide reasonable notice of an informal fact-finding conference pursuant to § 2.2-4019 to any person in connection with the denial, suspension, or revocation of a registration.

B. If a registration is revoked as the result of an informal hearing, the decision may be appealed, and upon appeal an administrative hearing shall be conducted in accordance with the Administrative Process Act (§ 2.2-4000 et seq.). The grower, dealer, or processor may appeal a final order to the circuit court in accordance with the Administrative Process Act.

C. A person issued a registration pursuant to subsection A of § 3.2-4115 who negligently (i) fails to provide a description and geographic data sufficient for locating his production field, dealership, or process site; (ii) grows, deals in, or processes Cannabis sativa with a tetrahydrocannabinol concentration

greater than that allowed by federal law; or (iii) produces a Cannabis sativa product shall comply with any corrective action plan established by the Commissioner in accordance with the provisions of subsection E. The Commissioner shall not deem a grower negligent if such grower makes reasonable efforts to grow industrial hemp and grows Cannabis sativa with a tetrahydrocannabinol concentration that does not exceed the total ~~delta-9~~ tetrahydrocannabinol concentration percentage established in federal regulations applicable to negligent violations located at 7 C.F.R. 990.6(b)(3).

D. A person who grows, deals in, or processes industrial hemp and who negligently fails to register pursuant to subsection A of § 3.2-4115 shall comply with any corrective action plan established by the Commissioner in accordance with the provisions of subsection E.

E. A corrective action plan established by the Commissioner in response to a negligent violation of a provision of this chapter shall identify a reasonable date by which the person who is the subject of the plan shall correct the negligent violation and shall require such person to report periodically for not less than two calendar years to the Commissioner on the person's compliance with the provisions of this chapter.

F. No person who negligently violates the provisions of this chapter three times in a five-year period shall be eligible to grow, deal in, or process industrial hemp for a period of five years beginning on the date of the third violation.

§ 4.1-600. Definitions.

As used in this subtitle, unless the context requires a different meaning:

"Advertisement" or " advertising" means any written or verbal statement, illustration, or depiction that is calculated to induce sales of retail marijuana, retail marijuana products, marijuana plants, or marijuana seeds, including any written, printed, graphic, digital, electronic, or other material, billboard, sign, or other outdoor display, publication, or radio or television broadcast.

"Authority" means the Virginia Cannabis Control Authority created pursuant to this subtitle.

"Board" means the Board of Directors of the Virginia Cannabis Control Authority.

"Cannabis Control Act" means Subtitle II (§ 4.1-600 et seq.).

"Child-resistant" means, with respect to packaging or a container, (i) specially designed or constructed to be significantly difficult for a typical child under five years of age to open and not to be significantly difficult for a typical adult to open and reseal and (ii) for any product intended for more than a single use or that contains multiple servings, resealable.

"Cultivation" or "cultivate" means the planting, propagation, growing, harvesting, drying, curing, grading, trimming, or other similar processing of marijuana for use or sale. "Cultivation" or "cultivate" does not include manufacturing or testing.

"Edible marijuana product" means a marijuana product intended to be consumed orally, including marijuana intended to be consumed orally or marijuana concentrate intended to be consumed orally.

"Immature plant" means a nonflowering marijuana plant that is no taller than eight inches and no wider than eight inches, is produced from a cutting, clipping, or seedling, and is growing in a container.

"Licensed" means the holding of a valid license granted by the Authority.

"Licensee" means any person to whom a license has been granted by the Authority.

"Manufacturing" or "manufacture" means the production of marijuana products or the blending, infusing, compounding, or other preparation of marijuana and marijuana products, including marijuana extraction or preparation by means of chemical synthesis. "Manufacturing" or "manufacture" does not include cultivation or testing.

"Marijuana" means *(i)* any part of a plant of the genus Cannabis, whether growing or not, its seeds or resin; and every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds, its resin, or any extract containing one or more cannabinoids *or (ii) any substance containing (a) a total tetrahydrocannabinol concentration that exceeds 0.3 percent or (b) more than one milligram of tetrahydrocannabinol per 100 grams of total product weight, including a hemp product, as defined in § 3.2-4112, or an industrial hemp extract, as defined in § 3.2-5145.1.* "Marijuana" does not include *(1)* the mature stalks of such plant, fiber produced from such stalk, or oil or cake made from the seed of such plant, unless such stalks, fiber, oil, or cake is combined with other parts of plants of the genus Cannabis~~. "Marijuana" does not include (i)~~*; (2)* industrial hemp, as defined in § 3.2-4112, that is possessed by a person registered pursuant to subsection A of § 3.2-4115 or his agent ~~or (ii)~~*; (3) industrial hemp, as defined in § 3.2-4112, that is possessed by a person who holds a hemp producer license issued by the U.S. Department of Agriculture pursuant to 7 C.F.R. Part 990 or his agent; (4)* a hemp product, as defined in § 3.2-4112, ~~containing~~ *that (a) is not intended for human consumption, orally or by inhalation, (b) contains* a tetrahydrocannabinol concentration of no greater than 0.3 percent ~~that~~*, and (c)* is derived from industrial hemp, as defined in § 3.2-4112, that is grown, dealt, or processed in compliance with state or federal law*; (5) an industrial hemp extract, as defined in § 3.2-5145.1, that (a) is derived from industrial hemp, as defined in § 3.2-4112, grown, dealt, or processed in compliance with state or federal law and (b) contains a tetrahydrocannabinol concentration of no greater than 0.3 percent and no more than one milligram of tetrahydrocannabinol per 100 grams of total product weight*

at the time such industrial hemp extract is offered for sale at retail; or (6) any drug product containing tetrahydrocannabinol that is approved for marketing by the U.S. Food and Drug Administration and scheduled in the Drug Control Act (§ 54.1-3400 et seq.) by the Board of Pharmacy pursuant to § 54.1-3443.

"Marijuana concentrate" means marijuana that has undergone a process to concentrate one or more active cannabinoids, thereby increasing the product's potency. Resin from granular trichomes from a marijuana plant is a concentrate for purposes of this subtitle.

"Marijuana cultivation facility" means a facility licensed under this subtitle to cultivate, label, and package retail marijuana; to purchase or take possession of marijuana plants and seeds from other marijuana cultivation facilities; to transfer possession of and sell retail marijuana, immature marijuana plants, and marijuana seeds to marijuana wholesalers and retail marijuana stores; to transfer possession of and sell retail marijuana, marijuana plants, and marijuana seeds to other marijuana cultivation facilities; to transfer possession of and sell retail marijuana to marijuana manufacturing facilities; and to sell immature marijuana plants and marijuana seeds to consumers for the purpose of cultivating marijuana at home for personal use.

"Marijuana establishment" means a marijuana cultivation facility, a marijuana testing facility, a marijuana manufacturing facility, a marijuana wholesaler, or a retail marijuana store.

"Marijuana manufacturing facility" means a facility licensed under this subtitle to manufacture, label, and package retail marijuana and retail marijuana products; to purchase or take possession of retail marijuana from a marijuana cultivation facility or another marijuana manufacturing facility; and to transfer possession of and sell retail marijuana and retail marijuana products to marijuana wholesalers, retail marijuana stores, or other marijuana manufacturing facilities.

"Marijuana paraphernalia" means all equipment, products, and materials of any kind that are either designed for use or are intended for use in planting, propagating, cultivating, growing, harvesting, manufacturing, compounding, converting, producing, processing, preparing, strength testing, analyzing, packaging, repackaging, storing, containing, concealing, ingesting, inhaling, or otherwise introducing into the human body marijuana.

"Marijuana products" means (i) products that are composed of marijuana and other ingredients and are intended for use or consumption, ointments, and tinctures or (ii) marijuana concentrate.

"Marijuana testing facility" means a facility licensed under this subtitle to develop, research, or test marijuana, marijuana products, and other substances.

"Marijuana wholesaler" means a facility licensed under this subtitle to purchase or take possession of retail marijuana, retail marijuana products, immature marijuana plants, and marijuana seeds from a marijuana cultivation facility, a marijuana manufacturing facility, or another marijuana wholesaler and to transfer possession and sell or resell retail marijuana, retail marijuana products, immature marijuana plants, and marijuana seeds to a marijuana cultivation facility, marijuana manufacturing facility, retail marijuana store, or another marijuana wholesaler.

"Non-retail marijuana" means marijuana that is not cultivated, manufactured, or sold by a licensed marijuana establishment.

"Non-retail marijuana products" means marijuana products that are not manufactured and sold by a licensed marijuana establishment.

"Place or premises" means the real estate, together with any buildings or other improvements thereon, designated in the application for a license as the place at which the cultivation, manufacture, sale, or testing of retail marijuana or retail marijuana products shall be performed, except that portion of any such building or other improvement actually and exclusively used as a private residence.

"Public place" means any place, building, or conveyance to which the public has, or is permitted to have, access, including restaurants, soda fountains, hotel dining areas, lobbies and corridors of hotels, and any park, place of public resort or amusement, highway, street, lane, or sidewalk adjoining any highway, street, or lane.

"Residence" means any building or part of a building or structure where a person resides, but does not include any part of a building that is not actually and exclusively used as a private residence, nor any part of a hotel or club other than a private guest room thereof.

"Retail marijuana" means marijuana that is cultivated, manufactured, or sold by a licensed marijuana establishment.

"Retail marijuana products" means marijuana products that are manufactured and sold by a licensed marijuana establishment.

"Retail marijuana store" means a facility licensed under this subtitle to purchase or take possession of retail marijuana, retail marijuana products, immature marijuana plants, or marijuana seeds from a marijuana cultivation facility, marijuana manufacturing facility, or marijuana wholesaler and to sell retail marijuana, retail marijuana products, immature marijuana plants, or marijuana seeds to consumers.

"Sale" and "sell" includes soliciting or receiving an order for; keeping, offering, or exposing for sale;

peddling, exchanging, or bartering; or delivering otherwise than gratuitously, by any means, retail marijuana or retail marijuana products.

"Special agent" means an employee of the Virginia Cannabis Control Authority whom the Board has designated as a law-enforcement officer pursuant to this subtitle.

"Testing" or "test" means the research and analysis of marijuana, marijuana products, or other substances for contaminants, safety, or potency. "Testing" or "test" does not include cultivation or manufacturing.

"Tetrahydrocannabinol" or "THC" means any naturally occurring or synthetic tetrahydrocannabinol, including its salts, isomers, or salts of isomers.

"Total tetrahydrocannabinol concentration" means the total available tetrahydrocannabinol derived from the molar sum of tetrahydrocannabinol and tetrahydrocannabinolic acid.

§ 9.1-1101. Powers and duties of the Department.

A. It shall be the responsibility of the Department to provide forensic laboratory services upon request of the Superintendent of State Police; the Chief Medical Examiner, the Assistant Chief Medical Examiners, and local medical examiners; any attorney for the Commonwealth; any chief of police, sheriff, or sergeant responsible for law enforcement in the jurisdiction served by him; any local fire department; the head of any private police department that has been designated as a criminal justice agency by the Department of Criminal Justice Services as defined by § 9.1-101; or any state agency in any criminal matter. The Department shall provide such services to any federal investigatory agency within available resources.

B. The Department shall:

1. Provide forensic laboratory services to all law-enforcement agencies throughout the Commonwealth and provide laboratory services, research, and scientific investigations for agencies of the Commonwealth as needed;

2. Establish and maintain a DNA testing program in accordance with Article 1.1 (§ 19.2-310.2 et seq.) of Chapter 18 of Title 19.2 to determine identification characteristics specific to an individual; ~~and~~

3. Test the accuracy of equipment used to test the blood alcohol content of breath at least once every six months. Only equipment found to be accurate shall be used to test the blood alcohol content of breath*; and*

4. Determine the proper methods for detecting the concentration of tetrahydrocannabinol (THC) in substances for the purposes of Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1, Chapter 7 (§ 18.2-247 et seq.) of Title 18.2, and §§ 54.1-3401 and 54.1-3446. The testing methodology shall use post-decarboxylation testing or another equivalent method and shall consider the potential conversion of tetrahydrocannabinolic acid (THC-A) into THC.

C. The Department shall have the power and duty to:

1. Receive, administer, and expend all funds and other assistance available for carrying out the purposes of this chapter;

2. Make and enter into all contracts and agreements necessary or incidental to the performance of its duties and execution of its powers under this chapter including, but not limited to, contracts with the United States, units of general local government or combinations thereof in Virginia or other states, and with agencies and departments of the Commonwealth; and

3. Perform such other acts as may be necessary or convenient for the effective performance of its duties.

D. The Director may appoint and employ a deputy director and such other personnel as are needed to carry out the duties and responsibilities conferred by this chapter.

§ 18.2-247. Use of terms "controlled substances," "counterfeit controlled substance," "imitation controlled substance," "marijuana," and "Schedules I, II, III, IV, V, and VI" in Title 18.2.

A. ~~Wherever the terms~~ *As used in Title 18.2,*"controlled substances" and "Schedules I, II, III, IV, V, and VI" ~~are used in Title 18.2, such terms refer to~~ *mean the same as* those terms ~~as they~~ are used or defined in the Drug Control Act (§ 54.1-3400 et seq.).

B. ~~The term~~ *As used in this article, unless the context requires a different meaning,* "imitation controlled substance" ~~when used in this article~~ means (i) a counterfeit controlled substance or (ii) a pill, capsule, tablet, or substance in any form whatsoever ~~which~~ *that* is not a controlled substance subject to abuse~~,~~ and:

1. ~~Which~~ *That* by overall dosage unit appearance, including color, shape, size, marking, and packaging or by representations made, would cause the likelihood that such a pill, capsule, tablet, or substance in any other form whatsoever will be mistaken for a controlled substance unless such substance was introduced into commerce prior to the initial introduction into commerce of the controlled substance ~~which~~ *that* it is alleged to imitate; or

2. ~~Which~~ *That* by express or implied representations purports to act like a controlled substance as a stimulant or depressant of the central nervous system and ~~which~~ *that* is not commonly used or recognized for use in that particular formulation for any purpose other than for such stimulant or

depressant effect, unless marketed, promoted, or sold as permitted by the U.S. Food and Drug Administration.

C. In determining whether a pill, capsule, tablet, or substance in any other form whatsoever~~,~~ is an "imitation controlled substance," there shall be considered, in addition to all other relevant factors, comparisons with accepted methods of marketing for legitimate nonprescription drugs for medicinal purposes rather than for drug abuse or any similar nonmedicinal use, including consideration of the packaging of the drug and its appearance in overall finished dosage form, promotional materials, or representations, oral or written, concerning the drug, and the methods of distribution of the drug and where and how it is sold to the public.

D. ~~The term "marijuana" when~~ *As* used in this article, *unless the context requires a different meaning:*

"Counterfeit controlled substance" means a controlled substance that, without authorization, bears, is packaged in a container or wrapper that bears, or is otherwise labeled to bear the trademark, trade name, or other identifying mark, imprint, or device or any likeness thereof of a drug manufacturer, processor, packer, or distributor other than the manufacturer, processor, packer, or distributor that did in fact so manufacture, process, pack, or distribute such drug.

"Marijuana" means *(i)* any part of a plant of the genus Cannabis, whether growing or not, its seeds or resin; and every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds, its resin, or any extract containing one or more cannabinoids *or (ii) any substance containing (a) a total tetrahydrocannabinol concentration that exceeds 0.3 percent or (b) more than one milligram of tetrahydrocannabinol per 100 grams of total product weight, including a hemp product, as defined in § 3.2-4112, or an industrial hemp extract, as defined in § 3.2-5145.1*. Marijuana does not include *(1)* the mature stalks of such plant, fiber produced from such stalk, oil or cake made from the seed of such plant, unless such stalks, fiber, oil or cake is combined with other parts of plants of the genus Cannabis~~.~~ ~~Marijuana does not include (i)~~*; (2)* industrial hemp, as defined in § 3.2-4112, that is possessed by a person registered pursuant to subsection A of § 3.2-4115 or his agent; ~~(ii)~~ *(3)* industrial hemp, as defined in § 3.2-4112, that is possessed by a person who holds a hemp producer license issued by the U.S. Department of Agriculture pursuant to 7 C.F.R. Part 990 *or his agent*; ~~or (iii)~~ *(4)* a hemp product, as defined in § 3.2-4112, ~~containing~~ *that (a) is not intended for human consumption, orally or by inhalation, (b) contains* a tetrahydrocannabinol concentration of no greater than 0.3 percent ~~that~~*, and (c)* is derived from industrial hemp, as defined in § 3.2-4112, that is grown, dealt, or processed in compliance with state or federal law*; (5) an industrial hemp extract, as defined in § 3.2-5145.1, that (a) is derived from industrial hemp, as defined in § 3.2-4112, grown, dealt, or processed in compliance with state or federal law and (b) contains a tetrahydrocannabinol concentration of no greater than 0.3 percent and no more than one milligram of tetrahydrocannabinol per 100 grams of total product weight at the time such industrial hemp extract is offered for sale at retail; or (6) any drug product containing tetrahydrocannabinol that is approved for marketing by the U.S. Food and Drug Administration and scheduled in the Drug Control Act (§ 54.1-3400 et seq.) by the Board of Pharmacy pursuant to § 54.1-3443.*

"Tetrahydrocannabinol" or "THC" means any naturally occurring or synthetic tetrahydrocannabinol, including its salts, isomers, or salts of isomers.

"Total tetrahydrocannabinol concentration" means the total available tetrahydrocannabinol derived from the molar sum of tetrahydrocannabinol and tetrahydrocannabinolic acid.

E. ~~The term "counterfeit controlled substance" means a controlled substance that, without authorization, bears, is packaged in a container or wrapper that bears, or is otherwise labeled to bear, the trademark, trade name, or other identifying mark, imprint or device or any likeness thereof, of a drug manufacturer, processor, packer, or distributor other than the manufacturer, processor, packer, or distributor who did in fact so manufacture, process, pack or distribute such drug.~~

~~F.~~ The Department of Forensic Science shall determine the proper methods for detecting the concentration of ~~delta-9-tetrahydrocannabinol~~ *tetrahydrocannabinol* (THC) in substances for the purposes of this title, *Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1*, and §§ 54.1-3401 and 54.1-3446. The testing methodology shall use post-decarboxylation testing or other equivalent method and shall consider the potential conversion of ~~delta-9-tetrahydrocannibinol~~ *tetrahydrocannabinolic* acid (THC-A) into THC. ~~The test result shall include the total available THC derived from the sum of the THC and THC-A content.~~

§ 19.2-188.1. Testimony regarding identification of controlled substances.

A. In any preliminary hearing on a violation of Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1, Article 1 (§ 18.2-247 et seq.) of Chapter 7 of Title 18.2, or subdivision 6 of § 53.1-203, any law-enforcement officer shall be permitted to testify as to the results of field tests that have been approved by the Department of Forensic Science pursuant to regulations adopted in accordance with the Administrative Process Act (§ 2.2-4000 et seq.), regarding whether or not any substance the identity of which is at issue in such hearing is a controlled substance, imitation controlled substance, or marijuana, as defined in §

4.1-600 or 18.2-247.

B. In any trial for a violation of § 4.1-1105.1, any law-enforcement officer shall be permitted to testify as to the results of any marijuana field test approved as accurate and reliable by the Department of Forensic Science pursuant to regulations adopted in accordance with the Administrative Process Act (§ 2.2-4000 et seq.), regarding whether or not any plant material, the identity of which is at issue, is marijuana provided the defendant has been given written notice of his right to request a full chemical analysis. Such notice shall be on a form approved by the Supreme Court and shall be provided to the defendant prior to trial.

In any case in which the person accused of a violation of § 4.1-1105.1, or the attorney of record for the accused, desires a full chemical analysis of the alleged plant material, he may, by motion prior to trial before the court in which the charge is pending, request such a chemical analysis. Upon such motion, the court shall order that the analysis be performed by the Department of Forensic Science in accordance with the provisions of § 18.2-247 and shall prescribe in its order the method of custody, transfer, and return of evidence submitted for chemical analysis.

§ 54.1-3401. Definitions.

As used in this chapter, unless the context requires a different meaning:

"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 (i) a practitioner or by his authorized agent and under his direction or (ii) the patient or research subject at the direction and in the presence of the practitioner.

"Advertisement" means all representations disseminated in any manner or by any means, other than by labeling, for the purpose of inducing, or which are likely to induce, directly or indirectly, the purchase of drugs or devices.

"Agent" means an authorized person who acts on behalf of or at the direction of a manufacturer, distributor, or dispenser. It does not include a common or contract carrier, public warehouseman, or employee of the carrier or warehouseman.

"Anabolic steroid" means any drug or hormonal substance, chemically and pharmacologically related to testosterone, other than estrogens, progestins, corticosteroids, and dehydroepiandrosterone.

"Animal" means any nonhuman animate being endowed with the power of voluntary action.

"Automated drug dispensing system" means a mechanical or electronic system that performs operations or activities, other than compounding or administration, relating to pharmacy services, including the storage, dispensing, or distribution of drugs and the collection, control, and maintenance of all transaction information, to provide security and accountability for such drugs.

"Biological product" means a virus, therapeutic serum, toxin, antitoxin, vaccine, blood, blood component or derivative, allergenic product, protein other than a chemically synthesized polypeptide, or analogous product, or arsphenamine or any derivative of arsphenamine or any other trivalent organic arsenic compound, applicable to the prevention, treatment, or cure of a disease or condition of human beings.

"Biosimilar" means a biological product that is highly similar to a specific reference biological product, notwithstanding minor differences in clinically inactive compounds, such that there are no clinically meaningful differences between the reference biological product and the biological product that has been licensed as a biosimilar pursuant to 42 U.S.C. § 262(k) in terms of safety, purity, and potency of the product.

"Board" means the Board of Pharmacy.

"Bulk drug substance" means any substance that is represented for use, and that, when used in the compounding, manufacturing, processing, or packaging of a drug, becomes an active ingredient or a finished dosage form of the drug; however, "bulk drug substance" shall not include intermediates that are used in the synthesis of such substances.

"Change of ownership" of an existing entity permitted, registered, or licensed by the Board means (i) the sale or transfer of all or substantially all of the assets of the entity or of any corporation that owns or controls the entity; (ii) the creation of a partnership by a sole proprietor, the dissolution of a partnership, or change in partnership composition; (iii) the acquisition or disposal of 50 percent or more of the outstanding shares of voting stock of a corporation owning the entity or of the parent corporation of a wholly owned subsidiary owning the entity, except that this shall not apply to any corporation the voting stock of which is actively traded on any securities exchange or in any over-the-counter market; (iv) the merger of a corporation owning the entity or of the parent corporation of a wholly-owned subsidiary owning the entity with another business or corporation; or (v) the expiration or forfeiture of a corporation's charter.

"Co-licensed partner" means a person who, with at least one other person, has the right to engage in the manufacturing or marketing of a prescription drug, consistent with state and federal law.

"Compounding" means the combining of two or more ingredients to fabricate such ingredients into a single preparation and includes the mixing, assembling, packaging, or labeling of a drug or device (i) by

a pharmacist, or within a permitted pharmacy, pursuant to a valid prescription issued for a medicinal or therapeutic purpose in the context of a bona fide practitioner-patient-pharmacist relationship, or in expectation of receiving a valid prescription based on observed historical patterns of prescribing and dispensing; (ii) by a practitioner of medicine, osteopathy, podiatry, dentistry, or veterinary medicine as an incident to his administering or dispensing, if authorized to dispense, a controlled substance in the course of his professional practice; or (iii) for the purpose of, or as incident to, research, teaching, or chemical analysis and not for sale or for dispensing. The mixing, diluting, or reconstituting of a manufacturer's product drugs for the purpose of administration to a patient, when performed by a practitioner of medicine or osteopathy licensed under Chapter 29 (§ 54.1-2900 et seq.), a person supervised by such practitioner pursuant to subdivision A 6 or 19 of § 54.1-2901, or a person supervised by such practitioner or a licensed nurse practitioner or physician assistant pursuant to subdivision A 4 of § 54.1-2901 shall not be considered compounding.

"Controlled substance" means a drug, substance, or immediate precursor in Schedules I through VI of this chapter. The term shall not include distilled spirits, wine, malt beverages, or tobacco as those terms are defined or used in Title 3.2 or Title 4.1. The term "controlled substance" includes a controlled substance analog that has been placed into Schedule I or II by the Board pursuant to the regulatory authority in subsection D of § 54.1-3443.

"Controlled substance analog" means a substance the chemical structure of which is substantially similar to the chemical structure of a controlled substance in Schedule I or II and either (i) which has a stimulant, depressant, or hallucinogenic effect on the central nervous system that is substantially similar to or greater than the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance in Schedule I or II or (ii) with respect to a particular person, which such person represents or intends to have a stimulant, depressant, or hallucinogenic effect on the central nervous system that is substantially similar to or greater than the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance in Schedule I or II. "Controlled substance analog" does not include (a) any substance for which there is an approved new drug application as defined under § 505 of the federal Food, Drug, and Cosmetic Act (21 U.S.C. § 355) or that is generally recognized as safe and effective pursuant to §§ 501, 502, and 503 of the federal Food, Drug, and Cosmetic Act (21 U.S.C. §§ 351, 352, and 353) and 21 C.F.R. Part 330; (b) with respect to a particular person, any substance for which an exemption is in effect for investigational use for that person under § 505 of the federal Food, Drug, and Cosmetic Act to the extent that the conduct with respect to that substance is pursuant to such exemption; or (c) any substance to the extent not intended for human consumption before such an exemption takes effect with respect to that substance.

"DEA" means the Drug Enforcement Administration, U.S. Department of Justice, or its successor agency.

"Deliver" or "delivery" means the actual, constructive, or attempted transfer of any item regulated by this chapter, whether or not there exists an agency relationship, including delivery of a Schedule VI prescription device to an ultimate user or consumer on behalf of a medical equipment supplier by a manufacturer, nonresident manufacturer, wholesale distributor, nonresident wholesale distributor, warehouser, nonresident warehouser, third-party logistics provider, or nonresident third-party logistics provider at the direction of a medical equipment supplier in accordance with § 54.1-3415.1.

"Device" means instruments, apparatus, and contrivances, including their components, parts, and accessories, intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in man or animals or to affect the structure or any function of the body of man or animals.

"Dialysis care technician" or "dialysis patient care technician" means an individual who is certified by an organization approved by the Board of Health Professions pursuant to Chapter 27.01 (§ 54.1-2729.1 et seq.) and who, under the supervision of a licensed physician, nurse practitioner, physician assistant, or a registered nurse, assists in the care of patients undergoing renal dialysis treatments in a Medicare-certified renal dialysis facility.

"Dialysis solution" means either the commercially available, unopened, sterile solutions whose purpose is to be instilled into the peritoneal cavity during the medical procedure known as peritoneal dialysis, or commercially available solutions whose purpose is to be used in the performance of hemodialysis not to include any solutions administered to the patient intravenously.

"Dispense" means to deliver a drug to an ultimate user or research subject by or pursuant to the lawful order of a practitioner, including the prescribing and administering, packaging, labeling, or compounding necessary to prepare the substance for that delivery. However, dispensing shall not include the transportation of drugs mixed, diluted, or reconstituted in accordance with this chapter to other sites operated by such practitioner or that practitioner's medical practice for the purpose of administration of such drugs to patients of the practitioner or that practitioner's medical practice at such other sites. For practitioners of medicine or osteopathy, "dispense" shall only include the provision of drugs by a practitioner to patients to take with them away from the practitioner's place of practice.

"Dispenser" means a practitioner who dispenses.

"Distribute" means to deliver other than by administering or dispensing a controlled substance.

"Distributor" means a person who distributes.

"Drug" means (i) articles or substances recognized in the official United States Pharmacopoeia National Formulary or official Homeopathic Pharmacopoeia of the United States, or any supplement to any of them; (ii) articles or substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in man or animals; (iii) articles or substances, other than food, intended to affect the structure or any function of the body of man or animals; (iv) articles or substances intended for use as a component of any article specified in clause (i), (ii), or (iii); or (v) a biological product. "Drug" does not include devices or their components, parts, or accessories.

"Drug product" means a specific drug in dosage form from a known source of manufacture, whether by brand or therapeutically equivalent drug product name.

"Electronic prescription" means a written prescription that is generated on an electronic application and is transmitted to a pharmacy as an electronic data file; Schedule II through V prescriptions shall be transmitted in accordance with 21 C.F.R. Part 1300.

"Facsimile (FAX) prescription" means a written prescription or order that is transmitted by an electronic device over telephone lines that sends the exact image to the receiving pharmacy in hard copy form.

"FDA" means the U.S. Food and Drug Administration.

"Immediate precursor" means a substance which the Board of Pharmacy 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.

"Interchangeable" means a biosimilar that meets safety standards for determining interchangeability pursuant to 42 U.S.C. § 262(k)(4).

"Label" means a display of written, printed, or graphic matter upon the immediate container of any article. A requirement made by or under authority of this chapter that any word, statement, or other information appear on the label shall not be considered to be complied with unless such word, statement, or other information also appears on the outside container or wrapper, if any, of the retail package of such article or is easily legible through the outside container or wrapper.

"Labeling" means all labels and other written, printed, or graphic matter on an article or any of its containers or wrappers, or accompanying such article.

"Manufacture" means the production, preparation, propagation, conversion, or processing of any item regulated by this chapter, either directly or indirectly by extraction from 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. This term does not include compounding.

"Manufacturer" means every person who manufactures, a manufacturer's co-licensed partner, or a repackager.

"Marijuana" means *(i)* any part of a plant of the genus Cannabis whether growing or not, its seeds, or its resin; and every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds, its resin, or any extract containing one or more cannabinoids *or (ii) any substance containing (a) a total tetrahydrocannabinol concentration that exceeds 0.3 percent or (b) more than one milligram of tetrahydrocannabinol per 100 grams of total product weight, including a hemp product, as defined in § 3.2-4112, or an industrial hemp extract, as defined in § 3.2-5145.1*. Marijuana does not include *(1)* the mature stalks of such plant, fiber produced from such stalk, or oil or cake made from the seeds of such plant, unless such stalks, fiber, oil, or cake is combined with other parts of plants of the genus Cannabis. ~~Marijuana does not include (i)~~*; (2)* industrial hemp, as defined in § 3.2-4112, that is possessed by a person registered pursuant to subsection A of § 3.2-4115 or his agent~~, (ii)~~*; (3)* industrial hemp, as defined in § 3.2-4112, that is possessed by a person who holds a hemp producer license issued by the U.S. Department of Agriculture pursuant to 7 C.F.R. Part 990~~, or (iii)~~ *or his agent; (4)* a hemp product, as defined in § 3.2-4112, ~~containing~~ *that (a) is not intended for human consumption, orally or by inhalation, (b) contains* a tetrahydrocannabinol concentration of no greater than 0.3 percent ~~that~~*, and (c)* is derived from industrial hemp, as defined in § 3.2-4112, that is grown, dealt, or processed in compliance with state or federal law*; (5) an industrial hemp extract, as defined in § 3.2-5145.1, that (a) is derived from industrial hemp, as defined in § 3.2-4112, grown, dealt, or processed in compliance with state or federal law and (b) contains a tetrahydrocannabinol concentration of no greater than 0.3 percent and no more than one milligram of tetrahydrocannabinol per 100 grams of total product weight at the time such industrial hemp extract is offered for sale at retail; or (6) any drug product containing tetrahydrocannabinol that is approved for marketing by the U.S. Food and Drug Administration and scheduled in the Drug Control Act (§ 54.1-3400 et seq.) by the Board of Pharmacy pursuant to § 54.1-3443*.

"Medical equipment supplier" means any person, as defined in § 1-230, engaged in the delivery to the ultimate consumer, pursuant to the lawful order of a practitioner, of hypodermic syringes and needles, medicinal oxygen, Schedule VI controlled devices, those Schedule VI controlled substances with no medicinal properties that are used for the operation and cleaning of medical equipment, solutions for peritoneal dialysis, and sterile water or saline for irrigation.

"Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis: (i) opium, opiates, and any salt, compound, derivative, or preparation of opium or opiates; (ii) any salt, compound, isomer, derivative, or preparation thereof which is chemically equivalent or identical with any of the substances referred to in clause (i), but not including the isoquinoline alkaloids of opium; (iii) opium poppy and poppy straw; (iv) coca leaves and any salt, compound, derivative, or preparation of coca leaves, and any salt, compound, isomer, derivative, or preparation thereof which is chemically equivalent or identical with any of these substances, but not including decocainized coca leaves or extraction of coca leaves which do not contain cocaine or ecgonine.

"New drug" means (i) any drug, except a new animal drug or an animal feed bearing or containing a new animal drug, the composition of which is such that such drug is not generally recognized, among experts qualified by scientific training and experience to evaluate the safety and effectiveness of drugs, as safe and effective for use under the conditions prescribed, recommended, or suggested in the labeling, except that such a drug not so recognized shall not be deemed to be a "new drug" if at any time prior to the enactment of this chapter it was subject to the Food and Drugs Act of June 30, 1906, as amended, and if at such time its labeling contained the same representations concerning the conditions of its use, or (ii) any drug, except a new animal drug or an animal feed bearing or containing a new animal drug, the composition of which is such that such drug, as a result of investigations to determine its safety and effectiveness for use under such conditions, has become so recognized, but which has not, otherwise than in such investigations, been used to a material extent or for a material time under such conditions.

"Nuclear medicine technologist" means an individual who holds a current certification with the American Registry of Radiological Technologists or the Nuclear Medicine Technology Certification Board.

"Official compendium" means the official United States Pharmacopoeia National Formulary, official Homeopathic Pharmacopoeia of the United States, or any supplement to any of them.

"Official written order" means an order written on a form provided for that purpose by the U.S. Drug Enforcement Administration, under any laws of the United States making provision therefor, if such order forms are authorized and required by federal law, and if no such order form is provided then on an official form provided for that purpose by the Board of Pharmacy.

"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 addiction-sustaining liability. It does not include, unless specifically designated as controlled under Article 4 (§ 54.1-3437 et seq.), the dextrorotatory isomer of 3-methoxy-n-methylmorphinan and its salts (dextromethorphan). It does include its racemic and levorotatory forms.

"Opium poppy" means the plant of the species Papaver somniferum L., except the seeds thereof.

"Original package" means the unbroken container or wrapping in which any drug or medicine is enclosed together with label and labeling, put up by or for the manufacturer, wholesaler, or distributor for use in the delivery or display of such article.

"Outsourcing facility" means a facility that is engaged in the compounding of sterile drugs and is currently registered as an outsourcing facility with the U.S. Secretary of Health and Human Services and that complies with all applicable requirements of federal and state law, including the Federal Food, Drug, and Cosmetic Act.

"Person" means both the plural and singular, as the case demands, and includes an individual, partnership, corporation, association, governmental agency, trust, or other institution or entity.

"Pharmacist-in-charge" means the person who, being licensed as a pharmacist, signs the application for a pharmacy permit and assumes full legal responsibility for the operation of the relevant pharmacy in a manner complying with the laws and regulations for the practice of pharmacy and the sale and dispensing of controlled substances; the "pharmacist-in-charge" shall personally supervise the pharmacy and the pharmacy's personnel as required by § 54.1-3432.

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

"Practitioner" means a physician, dentist, licensed nurse practitioner pursuant to § 54.1-2957.01, licensed physician assistant pursuant to § 54.1-2952.1, pharmacist pursuant to § 54.1-3300, TPA-certified optometrist pursuant to Article 5 (§ 54.1-3222 et seq.) of Chapter 32, veterinarian, scientific investigator, or other person licensed, registered, or otherwise permitted to distribute, dispense, prescribe and

administer, or conduct research with respect to a controlled substance in the course of professional practice or research in the Commonwealth.

"Prescriber" means a practitioner who is authorized pursuant to §§ 54.1-3303 and 54.1-3408 to issue a prescription.

"Prescription" means an order for drugs or medical supplies, written or signed or transmitted by word of mouth, telephone, telegraph, or other means of communication to a pharmacist by a duly licensed physician, dentist, veterinarian, or other practitioner authorized by law to prescribe and administer such drugs or medical supplies.

"Prescription drug" means any drug required by federal law or regulation to be dispensed only pursuant to a prescription, including finished dosage forms and active ingredients subject to § 503(b) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. § 353(b)).

"Production" or "produce" includes the manufacture, planting, cultivation, growing, or harvesting of a controlled substance or marijuana.

"Proprietary medicine" means a completely compounded nonprescription drug in its unbroken, original package which does not contain any controlled substance or marijuana as defined in this chapter and is not in itself poisonous, and which is sold, offered, promoted, or advertised directly to the general public by or under the authority of the manufacturer or primary distributor, under a trademark, trade name, or other trade symbol privately owned, and the labeling of which conforms to the requirements of this chapter and applicable federal law. However, this definition shall not include a drug that is only advertised or promoted professionally to licensed practitioners, a narcotic or drug containing a narcotic, a drug that may be dispensed only upon prescription or the label of which bears substantially the statement "Warning — may be habit-forming," or a drug intended for injection.

"Radiopharmaceutical" means any drug that exhibits spontaneous disintegration of unstable nuclei with the emission of nuclear particles or photons and includes any non-radioactive reagent kit or radionuclide generator that is intended to be used in the preparation of any such substance, but does not include drugs such as carbon-containing compounds or potassium-containing salts that include trace quantities of naturally occurring radionuclides. The term also includes any biological product that is labeled with a radionuclide or intended solely to be labeled with a radionuclide.

"Reference biological product" means the single biological product licensed pursuant to 42 U.S.C. § 262(a) against which a biological product is evaluated in an application submitted to the U.S. Food and Drug Administration for licensure of biological products as biosimilar or interchangeable pursuant to 42 U.S.C. § 262(k).

"Sale" includes barter, exchange, or gift, or offer therefor, and each such transaction made by any person, whether as an individual, proprietor, agent, servant, or employee.

"Tetrahydrocannabinol" or "THC" means any naturally occurring or synthetic tetrahydrocannabinol, including its salts, isomers, or salts of isomers.

"Therapeutically equivalent drug products" means drug products that contain the same active ingredients and are identical in strength or concentration, dosage form, and route of administration and that are classified as being therapeutically equivalent by the U.S. Food and Drug Administration pursuant to the definition of "therapeutically equivalent drug products" set forth in the most recent edition of the Approved Drug Products with Therapeutic Equivalence Evaluations, otherwise known as the "Orange Book."

"Third-party logistics provider" means a person that provides or coordinates warehousing of or other logistics services for a drug or device in interstate commerce on behalf of a manufacturer, wholesale distributor, or dispenser of the drug or device but does not take ownership of the product or have responsibility for directing the sale or disposition of the product.

"Total tetrahydrocannabinol concentration" means the total available tetrahydrocannabinol derived from the molar sum of tetrahydrocannabinol and tetrahydrocannabinolic acid.

"USP-NF" means the current edition of the United States Pharmacopeia-National Formulary.

"Warehouser" means any person, other than a wholesale distributor, manufacturer, or third-party logistics provider, engaged in the business of (i) selling or otherwise distributing prescription drugs or devices to any person who is not the ultimate user or consumer and (ii) delivering Schedule VI prescription devices to the ultimate user or consumer pursuant to § 54.1-3415.1. No person shall be subject to any state or local tax by reason of this definition.

"Wholesale distribution" means (i) distribution of prescription drugs to persons other than consumers or patients and (ii) delivery of Schedule VI prescription devices to the ultimate user or consumer pursuant to § 54.1-3415.1, subject to the exemptions set forth in the federal Drug Supply Chain Security Act.

"Wholesale distributor" means any person other than a manufacturer, a manufacturer's co-licensed partner, a third-party logistics provider, or a repackager that engages in wholesale distribution.

The words "drugs" and "devices" as used in Chapter 33 (§ 54.1-3300 et seq.) and in this chapter shall not include surgical or dental instruments, physical therapy equipment, X-ray apparatus, or glasses

or lenses for the eyes.

The terms "pharmacist," "pharmacy," and "practice of pharmacy" as used in this chapter shall be defined as provided in Chapter 33 (§ 54.1-3300 et seq.) unless the context requires a different meaning.

§ 54.1-3408.3. Certification for use of cannabis oil for treatment.

A. As used in this section:

"Botanical cannabis" means cannabis that is composed wholly of usable cannabis from the same parts of the same chemovar of cannabis plant.

"Cannabis oil" means any formulation of processed Cannabis plant extract, which may include industrial hemp extracts, including isolates and distillates, acquired by a pharmaceutical processor pursuant to § 54.1-3442.6, or a dilution of the resin of the Cannabis plant that contains no more than 10 milligrams of ~~delta-9-tetrahydrocannabinol~~ *tetrahydrocannabinol* per dose. "Cannabis oil" does not include industrial hemp, as defined in § 3.2-4112, that is grown, dealt, or processed in compliance with state or federal law, unless it has been grown and processed in the Commonwealth by a registered industrial hemp processor and acquired and formulated by a pharmaceutical processor.

"Cannabis product" means a product that is (i) produced by a pharmaceutical processor, registered with the Board, and compliant with testing requirements and (ii) composed of cannabis oil or botanical cannabis.

"Designated caregiver facility" means any hospice or hospice facility licensed pursuant to § 32.1-162.3, or home care organization as defined in § 32.1-162.7 that provides pharmaceutical services or home health services, private provider licensed by the Department of Behavioral Health and Developmental Services pursuant to Article 2 (§ 37.2-403 et seq.) of Chapter 4 of Title 37.2, assisted living facility licensed pursuant to § 63.2-1701, or adult day care center licensed pursuant to § 63.2-1701.

"Practitioner" means a practitioner of medicine or osteopathy licensed by the Board of Medicine, a physician assistant licensed by the Board of Medicine, or a nurse practitioner jointly licensed by the Board of Medicine and the Board of Nursing.

"Registered agent" means an individual designated by a patient who has been issued a written certification, or, if such patient is a minor or a vulnerable adult as defined in § 18.2-369, designated by such patient's parent or legal guardian, and registered with the Board pursuant to subsection G.

"Usable cannabis" means any cannabis plant material, including seeds, but not (i) resin that has been extracted from any part of the cannabis plant, its seeds, or its resin; (ii) the mature stalks, fiber produced from the stalks, or any other compound, manufacture, salt, or derivative, mixture, or preparation of the mature stalks; or (iii) oil or cake made from the seeds of the plant.

B. A practitioner in the course of his professional practice may issue a written certification for the use of cannabis products for treatment or to alleviate the symptoms of any diagnosed condition or disease determined by the practitioner to benefit from such use. The practitioner shall use his professional judgment to determine the manner and frequency of patient care and evaluation and may employ the use of telemedicine, provided that the use of telemedicine includes the delivery of patient care through real-time interactive audio-visual technology. If a practitioner determines it is consistent with the standard of care to dispense botanical cannabis to a minor, the written certification shall specifically authorize such dispensing. If not specifically included on the initial written certification, authorization for botanical cannabis may be communicated verbally or in writing to the pharmacist at the time of dispensing.

C. The written certification shall be on a form provided by the Board of Pharmacy. Such written certification shall contain the name, address, and telephone number of the practitioner; the name and address of the patient issued the written certification; the date on which the written certification was made; and the signature or authentic electronic signature of the practitioner. Such written certification issued pursuant to subsection B shall expire no later than one year after its issuance unless the practitioner provides in such written certification an earlier expiration. A written certification shall not be issued to a patient by more than one practitioner during any given time period.

D. No practitioner shall be prosecuted under § 18.2-248 or 18.2-248.1 for the issuance of a certification for the use of cannabis products for the treatment or to alleviate the symptoms of a patient's diagnosed condition or disease pursuant to a written certification issued pursuant to subsection B. Nothing in this section shall preclude the Board of Medicine from sanctioning a practitioner for failing to properly evaluate or treat a patient's medical condition or otherwise violating the applicable standard of care for evaluating or treating medical conditions.

E. A practitioner who issues a written certification to a patient pursuant to this section shall register with the Board and shall hold sufficient education and training to exercise appropriate professional judgment in the certification of patients. The Board shall not limit the number of patients to whom a practitioner may issue a written certification. The Board may report information to the applicable licensing board on unusual patterns of certifications issued by a practitioner.

F. No patient shall be required to physically present the written certification after the initial dispensing by any pharmaceutical processor or cannabis dispensing facility under each written certification, provided that the pharmaceutical processor or cannabis dispensing facility maintains an electronic copy of the written certification. Pharmaceutical processors and cannabis dispensing facilities shall electronically transmit, on a monthly basis, all new written certifications received by the pharmaceutical processor or cannabis dispensing facility to the Board.

G. A patient, or, if such patient is a minor or a vulnerable adult as defined in § 18.2-369, such patient's parent or legal guardian, may designate an individual to act as his registered agent for the purposes of receiving cannabis products pursuant to a valid written certification. Such designated individual shall register with the Board. The Board may set a limit on the number of patients for whom any individual is authorized to act as a registered agent.

H. Upon delivery of a cannabis product by a pharmaceutical processor or cannabis dispensing facility to a designated caregiver facility, any employee or contractor of a designated caregiver facility, who is licensed or registered by a health regulatory board and who is authorized to possess, distribute, or administer medications, may accept delivery of the cannabis product on behalf of a patient or resident for subsequent delivery to the patient or resident and may assist in the administration of the cannabis product to the patient or resident as necessary.

I. Information obtained under the registration process shall be confidential and shall not be subject to the disclosure provisions of the Virginia Freedom of Information Act (§ 2.2-3700 et seq.). However, reasonable access to registry information shall be provided to (i) the Chairmen of the House Committee for Courts of Justice and the Senate Committee on the Judiciary, (ii) state and federal agencies or local law enforcement for the purpose of investigating or prosecuting a specific individual for a specific violation of law, (iii) licensed practitioners or pharmacists, or their agents, for the purpose of providing patient care and drug therapy management and monitoring of drugs obtained by a patient, (iv) a pharmaceutical processor or cannabis dispensing facility involved in the treatment of a patient, or (v) a registered agent, but only with respect to information related to such patient.

§ 54.1-3442.6. Permit to operate pharmaceutical processor or cannabis dispensing facility.

A. No person shall operate a pharmaceutical processor or a cannabis dispensing facility without first obtaining a permit from the Board. The application for such permit shall be made on a form provided by the Board and signed by a pharmacist who will be in full and actual charge of the pharmaceutical processor's dispensing area or cannabis dispensing facility. The Board shall establish an application fee and other general requirements for such application.

B. Each permit shall expire annually on a date determined by the Board in regulation. The number of permits that the Board may issue or renew in any year is limited to one pharmaceutical processor and up to five cannabis dispensing facilities for each health service area established by the Board of Health. Permits shall be displayed in a conspicuous place on the premises of the pharmaceutical processor and cannabis dispensing facility.

C. The Board shall adopt regulations establishing health, safety, and security requirements for pharmaceutical processors and cannabis dispensing facilities. Such regulations shall include requirements for (i) physical standards; (ii) location restrictions; (iii) security systems and controls; (iv) minimum equipment and resources; (v) recordkeeping; (vi) labeling, including the potency of each botanical cannabis product and the amounts recommended by the practitioner or dispensing pharmacist, and packaging; (vii) routine inspections no more frequently than once annually; (viii) processes for safely and securely dispensing and delivering in person cannabis products to a patient, his registered agent, or, if such patient is a minor or a vulnerable adult as defined in § 18.2-369, such patient's parent or legal guardian; (ix) dosage limitations for cannabis oil that provide that each dispensed dose of cannabis oil not exceed 10 milligrams of ~~delta-9-tetrahydrocannabinol~~ *tetrahydrocannabinol*; (x) a process for the wholesale distribution of and the transfer of usable cannabis, botanical cannabis, cannabis oil, and cannabis products between pharmaceutical processors, between a pharmaceutical processors and a cannabis dispensing facility, and between cannabis dispensing facilities; (xi) an allowance for the sale of devices for administration of dispensed cannabis products and hemp-based CBD products that meet the applicable standards set forth in state and federal law, including the laboratory testing standards set forth in subsection M; (xii) an allowance for the use and distribution of inert product samples containing no cannabinoids for patient demonstration exclusively at the pharmaceutical processor or cannabis dispensing facility, and not for further distribution or sale, without the need for a written certification; (xiii) a process for acquiring industrial hemp extracts and formulating such extracts into cannabis products; and (xiv) an allowance for the advertising and promotion of the pharmaceutical processor's products and operations, which shall not limit the pharmaceutical processor from the provision of educational material to practitioners who issue written certifications and patients. The Board shall also adopt regulations for pharmaceutical processors that include requirements for (a) processes for safely and securely cultivating Cannabis plants intended for producing cannabis products, (b) the secure disposal of agricultural waste, and (c) a process for registering cannabis oil products.

D. The Board shall require that, after processing and before dispensing any cannabis products, a pharmaceutical processor shall make a sample available from each batch of cannabis product for testing by an independent laboratory located in Virginia meeting Board requirements. A valid sample size for testing shall be determined by each laboratory and may vary due to sample matrix, analytical method, and laboratory-specific procedures. A minimum sample size of 0.5 percent of individual units for dispensing or distribution from each homogenized batch of cannabis oil is required to achieve a representative cannabis oil sample for analysis. A minimum sample size, to be determined by the certified testing laboratory, from each batch of botanical cannabis is required to achieve a representative botanical cannabis sample for analysis. Botanical cannabis products shall only be tested for the following: total cannabidiol (CBD); total tetrahydrocannabinol (THC); terpenes; pesticide chemical residue; heavy metals; mycotoxins; moisture; and microbiological contaminants. Testing thresholds shall be consistent with generally accepted cannabis industry thresholds. The pharmaceutical processor may remediate botanical cannabis or cannabis oil that fails any quality testing standard except pesticides. Following remediation, all remediated botanical cannabis or cannabis oil shall be subject to laboratory testing and approved upon satisfaction of applicable testing standards, which shall not be more stringent than initial testing prior to remediation. If a batch of botanical cannabis fails retesting after remediation, it shall be considered usable cannabis and may be processed into cannabis oil. Stability testing shall not be required for any cannabis product with an expiration date assigned by the pharmaceutical processor of six months or less from the date of the cannabis product registration approval. Stability testing required for assignment of an expiration date longer than six months shall be limited to microbial testing, on a pass/fail basis, and potency testing, on a 10 percent deviation basis, of active ingredients.

E. A laboratory testing samples for a pharmaceutical processor shall obtain a controlled substances registration certificate pursuant to § 54.1-3423 and shall comply with quality standards established by the Board in regulation.

F. Every pharmaceutical processor's dispensing area or cannabis dispensing facility shall be under the personal supervision of a licensed pharmacist on the premises of the pharmaceutical processor or cannabis dispensing facility. The pharmaceutical processor shall ensure that security measures are adequate to protect the cannabis from diversion at all times, and the pharmacist-in-charge shall have concurrent responsibility for preventing diversion from the dispensing area.

Every pharmaceutical processor shall designate a person who shall have oversight of the cultivation and production areas of the pharmaceutical processor and shall provide such information to the Board. The Board shall direct all communications related to enforcement of requirements related to cultivation and production of cannabis oil products by the pharmaceutical processor to such designated person.

G. The Board shall require the material owners of an applicant for a pharmaceutical processor or cannabis dispensing facility permit to submit to fingerprinting and provide personal descriptive information to be forwarded along with his fingerprints through the Central Criminal Records Exchange to the Federal Bureau of Investigation for the purpose of obtaining criminal history record information regarding the applicant's material owners. The cost of fingerprinting and the criminal history record search shall be paid by the applicant. The Central Criminal Records Exchange shall forward the results of the criminal history background check to the Board or its designee, which shall be a governmental entity. A pharmaceutical processor shall maintain evidence of criminal background checks for all employees and delivery agents of the pharmaceutical processor. Criminal background checks of employees and delivery agents may be conducted by any service sufficient to disclose any federal and state criminal convictions.

H. In addition to other employees authorized by the Board, a pharmaceutical processor may employ individuals who may have less than two years of experience (i) to perform cultivation-related duties under the supervision of an individual who has received a degree in a field related to the cultivation of plants or a certification recognized by the Board or who has at least two years of experience cultivating plants, (ii) to perform extraction-related duties under the supervision of an individual who has a degree in chemistry or pharmacology or at least two years of experience extracting chemicals from plants, and (iii) to perform duties at the pharmaceutical processor and cannabis dispensing facility upon certification as a pharmacy technician.

I. A pharmaceutical processor to whom a permit has been issued by the Board may establish up to five cannabis dispensing facilities for the dispensing of cannabis products that have been cultivated and produced on the premises of a pharmaceutical processor permitted by the Board. Each cannabis dispensing facility shall be located within the same health service area as the pharmaceutical processor.

J. No person who has been convicted of a felony under the laws of the Commonwealth or another jurisdiction within the last five years shall be employed by or act as an agent of a pharmaceutical processor or cannabis dispensing facility.

K. Every pharmaceutical processor or cannabis dispensing facility shall adopt policies for pre-employment drug screening and regular, ongoing, random drug screening of employees.

L. A pharmacist at the pharmaceutical processor's dispensing area and the cannabis dispensing facility shall determine the number of pharmacy interns, pharmacy technicians, and pharmacy technician trainees who can be safely and competently supervised at one time; however, no pharmacist shall supervise more than six persons performing the duties of a pharmacy technician at one time in the pharmaceutical processor's dispensing area or cannabis dispensing facility.

M. A pharmaceutical processor may acquire industrial hemp extracts grown and processed in Virginia, and in compliance with state or federal law, from a registered industrial hemp dealer or processor. A pharmaceutical processor may process and formulate such extracts into an allowable dosage of cannabis product. Industrial hemp extracts acquired and formulated by a pharmaceutical processor are subject to the same third-party testing requirements that may apply to cannabis plant extract. Testing shall be performed by a laboratory located in Virginia and in compliance with state law governing the testing of cannabis products. The industrial hemp dealer or processor shall provide such third-party testing results to the pharmaceutical processor before industrial hemp extracts may be acquired.

N. With the exception of § 2.2-4031, neither the provisions of the Administrative Process Act (§ 2.2-4000 et seq.) nor public participation guidelines adopted pursuant thereto shall apply to the adoption of any regulation pursuant to this section. Prior to adopting any regulation pursuant to this section, the Board of Pharmacy shall publish a notice of opportunity to comment in the Virginia Register of Regulations and post the action on the Virginia Regulatory Town Hall. Such notice of opportunity to comment shall contain (i) a summary of the proposed regulation; (ii) the text of the proposed regulation; and (iii) the name, address, and telephone number of the agency contact person responsible for receiving public comments. Such notice shall be made at least 60 days in advance of the last date prescribed in such notice for submittals of public comment. The legislative review provisions of subsections A and B of § 2.2-4014 shall apply to the promulgation or final adoption process for regulations pursuant to this section. The Board of Pharmacy shall consider and keep on file all public comments received for any regulation adopted pursuant to this section.

O. The Board shall register all cannabis products that meet testing, labeling, and packaging standards.

§ 54.1-3442.7. Dispensing cannabis products; report.

A. A pharmaceutical processor or cannabis dispensing facility shall dispense or deliver cannabis products only in person to (i) a patient who is a Virginia resident or temporarily resides in Virginia and has been issued a valid written certification; (ii) such patient's registered agent; or (iii) if such patient is a minor or a vulnerable adult as defined in § 18.2-369, such patient's parent or legal guardian who is a Virginia resident or temporarily resides in Virginia. A companion may accompany a patient into a pharmaceutical processor's dispensing area or cannabis dispensing facility. Prior to the initial dispensing of cannabis products pursuant to each written certification, a pharmacist or pharmacy technician employed by the pharmaceutical processor or cannabis dispensing facility shall make and maintain, on site or remotely by electronic means, for two years a paper or electronic copy of the written certification that provides an exact image of the document that is clearly legible; shall view, in person or by audiovisual means, a current photo identification of the patient, registered agent, parent, or legal guardian; and shall verify current board registration of the practitioner and the corresponding registered agent if applicable. Thereafter, an initial dispensing may be delivered to the patient, registered agent, parent, legal guardian, or designated caregiver facility. Prior to any subsequent dispensing of cannabis products pursuant to each written certification, an employee or delivery agent shall view a current photo identification of the patient, registered agent, parent, or legal guardian and the current board registration issued to the registered agent if applicable. No pharmaceutical processor or cannabis dispensing facility shall dispense more than a 90-day supply, as determined by the dispensing pharmacist or certifying practitioner, for any patient during any 90-day period. A pharmaceutical processor or cannabis dispensing facility may dispense less than a 90-day supply of a cannabis product for any patient during any 90-day period; however, a pharmaceutical processor or cannabis dispensing facility may dispense more than one cannabis product to a patient at one time. No more than four ounces of botanical cannabis shall be dispensed for each 30-day period for which botanical cannabis is dispensed. The Board shall establish in regulation an amount of cannabis oil that constitutes a 90-day supply to treat or alleviate the symptoms of a patient's diagnosed condition or disease. In determining the appropriate amount of a cannabis product to be dispensed to a patient, a pharmaceutical processor or cannabis dispensing facility shall consider all cannabis products dispensed to the patient and adjust the amount dispensed accordingly.

B. A pharmaceutical processor or cannabis dispensing facility shall dispense only cannabis products produced on the premises of a pharmaceutical processor permitted by the Board or cannabis products that have been formulated with extracts from industrial hemp acquired by a pharmaceutical processor from a registered industrial hemp dealer or processor pursuant to § 54.1-3442.6. A pharmaceutical processor may begin cultivation upon being issued a permit by the Board.

C. The Board shall report annually by December 1 to the Chairmen of the House Committee for Health, Welfare and Institutions and the Senate Committee on Education and Health on the operation of

pharmaceutical processors and cannabis dispensing facilities issued a permit by the Board.

D. The concentration of ~~delta-9-tetrahydrocannabinol~~ *tetrahydrocannabinol* in any cannabis product on site may be up to 10 percent greater than or less than the level of ~~delta-9-tetrahydrocannabinol~~ *tetrahydrocannabinol* measured for labeling. A pharmaceutical processor and cannabis dispensing facility shall ensure that such concentration in any cannabis product on site is within such range. A pharmaceutical processor producing cannabis products shall establish a stability testing schedule of cannabis products.

§ 59.1-200. Prohibited practices.

A. The following fraudulent acts or practices committed by a supplier in connection with a consumer transaction are hereby declared unlawful:

1. Misrepresenting goods or services as those of another;

2. Misrepresenting the source, sponsorship, approval, or certification of goods or services;

3. Misrepresenting the affiliation, connection, or association of the supplier, or of the goods or services, with another;

4. Misrepresenting geographic origin in connection with goods or services;

5. Misrepresenting that goods or services have certain quantities, characteristics, ingredients, uses, or benefits;

6. Misrepresenting that goods or services are of a particular standard, quality, grade, style, or model;

7. Advertising or offering for sale goods that are used, secondhand, repossessed, defective, blemished, deteriorated, or reconditioned, or that are "seconds," irregulars, imperfects, or "not first class," without clearly and unequivocally indicating in the advertisement or offer for sale that the goods are used, secondhand, repossessed, defective, blemished, deteriorated, reconditioned, or are "seconds," irregulars, imperfects or "not first class";

8. Advertising goods or services with intent not to sell them as advertised, or with intent not to sell at the price or upon the terms advertised.

In any action brought under this subdivision, the refusal by any person, or any employee, agent, or servant thereof, to sell any goods or services advertised or offered for sale at the price or upon the terms advertised or offered, shall be prima facie evidence of a violation of this subdivision. This paragraph shall not apply when it is clearly and conspicuously stated in the advertisement or offer by which such goods or services are advertised or offered for sale, that the supplier or offeror has a limited quantity or amount of such goods or services for sale, and the supplier or offeror at the time of such advertisement or offer did in fact have or reasonably expected to have at least such quantity or amount for sale;

9. Making false or misleading statements of fact concerning the reasons for, existence of, or amounts of price reductions;

10. Misrepresenting that repairs, alterations, modifications, or services have been performed or parts installed;

11. Misrepresenting by the use of any written or documentary material that appears to be an invoice or bill for merchandise or services previously ordered;

12. Notwithstanding any other provision of law, using in any manner the words "wholesale," "wholesaler," "factory," or "manufacturer" in the supplier's name, or to describe the nature of the supplier's business, unless the supplier is actually engaged primarily in selling at wholesale or in manufacturing the goods or services advertised or offered for sale;

13. Using in any contract or lease any liquidated damage clause, penalty clause, or waiver of defense, or attempting to collect any liquidated damages or penalties under any clause, waiver, damages, or penalties that are void or unenforceable under any otherwise applicable laws of the Commonwealth, or under federal statutes or regulations;

13a. Failing to provide to a consumer, or failing to use or include in any written document or material provided to or executed by a consumer, in connection with a consumer transaction any statement, disclosure, notice, or other information however characterized when the supplier is required by 16 C.F.R. Part 433 to so provide, use, or include the statement, disclosure, notice, or other information in connection with the consumer transaction;

14. Using any other deception, fraud, false pretense, false promise, or misrepresentation in connection with a consumer transaction;

15. Violating any provision of § 3.2-6509, 3.2-6512, 3.2-6513, 3.2-6513.1, 3.2-6514, 3.2-6515, 3.2-6516, or 3.2-6519 is a violation of this chapter;

16. Failing to disclose all conditions, charges, or fees relating to:

a. The return of goods for refund, exchange, or credit. Such disclosure shall be by means of a sign attached to the goods, or placed in a conspicuous public area of the premises of the supplier, so as to be readily noticeable and readable by the person obtaining the goods from the supplier. If the supplier does not permit a refund, exchange, or credit for return, he shall so state on a similar sign. The provisions of this subdivision shall not apply to any retail merchant who has a policy of providing, for a period of not

less than 20 days after date of purchase, a cash refund or credit to the purchaser's credit card account for the return of defective, unused, or undamaged merchandise upon presentation of proof of purchase. In the case of merchandise paid for by check, the purchase shall be treated as a cash purchase and any refund may be delayed for a period of 10 banking days to allow for the check to clear. This subdivision does not apply to sale merchandise that is obviously distressed, out of date, post season, or otherwise reduced for clearance; nor does this subdivision apply to special order purchases where the purchaser has requested the supplier to order merchandise of a specific or unusual size, color, or brand not ordinarily carried in the store or the store's catalog; nor shall this subdivision apply in connection with a transaction for the sale or lease of motor vehicles, farm tractors, or motorcycles as defined in § 46.2-100;

b. A layaway agreement. Such disclosure shall be furnished to the consumer (i) in writing at the time of the layaway agreement, or (ii) by means of a sign placed in a conspicuous public area of the premises of the supplier, so as to be readily noticeable and readable by the consumer, or (iii) on the bill of sale. Disclosure shall include the conditions, charges, or fees in the event that a consumer breaches the agreement;

16a. Failing to provide written notice to a consumer of an existing open-end credit balance in excess of $5 (i) on an account maintained by the supplier and (ii) resulting from such consumer's overpayment on such account. Suppliers shall give consumers written notice of such credit balances within 60 days of receiving overpayments. If the credit balance information is incorporated into statements of account furnished consumers by suppliers within such 60-day period, no separate or additional notice is required;

17. If a supplier enters into a written agreement with a consumer to resolve a dispute that arises in connection with a consumer transaction, failing to adhere to the terms and conditions of such an agreement;

18. Violating any provision of the Virginia Health Club Act, Chapter 24 (§ 59.1-294 et seq.);

19. Violating any provision of the Virginia Home Solicitation Sales Act, Chapter 2.1 (§ 59.1-21.1 et seq.);

20. Violating any provision of the Automobile Repair Facilities Act, Chapter 17.1 (§ 59.1-207.1 et seq.);

21. Violating any provision of the Virginia Lease-Purchase Agreement Act, Chapter 17.4 (§ 59.1-207.17 et seq.);

22. Violating any provision of the Prizes and Gifts Act, Chapter 31 (§ 59.1-415 et seq.);

23. Violating any provision of the Virginia Public Telephone Information Act, Chapter 32 (§ 59.1-424 et seq.);

24. Violating any provision of § 54.1-1505;

25. Violating any provision of the Motor Vehicle Manufacturers' Warranty Adjustment Act, Chapter 17.6 (§ 59.1-207.34 et seq.);

26. Violating any provision of § 3.2-5627, relating to the pricing of merchandise;

27. Violating any provision of the Pay-Per-Call Services Act, Chapter 33 (§ 59.1-429 et seq.);

28. Violating any provision of the Extended Service Contract Act, Chapter 34 (§ 59.1-435 et seq.);

29. Violating any provision of the Virginia Membership Camping Act, Chapter 25 (§ 59.1-311 et seq.);

30. Violating any provision of the Comparison Price Advertising Act, Chapter 17.7 (§ 59.1-207.40 et seq.);

31. Violating any provision of the Virginia Travel Club Act, Chapter 36 (§ 59.1-445 et seq.);

32. Violating any provision of §§ 46.2-1231 and 46.2-1233.1;

33. Violating any provision of Chapter 40 (§ 54.1-4000 et seq.) of Title 54.1;

34. Violating any provision of Chapter 10.1 (§ 58.1-1031 et seq.) of Title 58.1;

35. Using the consumer's social security number as the consumer's account number with the supplier, if the consumer has requested in writing that the supplier use an alternate number not associated with the consumer's social security number;

36. Violating any provision of Chapter 18 (§ 6.2-1800 et seq.) of Title 6.2;

37. Violating any provision of § 8.01-40.2;

38. Violating any provision of Article 7 (§ 32.1-212 et seq.) of Chapter 6 of Title 32.1;

39. Violating any provision of Chapter 34.1 (§ 59.1-441.1 et seq.);

40. Violating any provision of Chapter 20 (§ 6.2-2000 et seq.) of Title 6.2;

41. Violating any provision of the Virginia Post-Disaster Anti-Price Gouging Act, Chapter 46 (§ 59.1-525 et seq.);

42. Violating any provision of Chapter 47 (§ 59.1-530 et seq.);

43. Violating any provision of § 59.1-443.2;

44. Violating any provision of Chapter 48 (§ 59.1-533 et seq.);

45. Violating any provision of Chapter 25 (§ 6.2-2500 et seq.) of Title 6.2;

46. Violating the provisions of clause (i) of subsection B of § 54.1-1115;

47. Violating any provision of § 18.2-239;

48. Violating any provision of Chapter 26 (§ 59.1-336 et seq.);

49. Selling, offering for sale, or manufacturing for sale a children's product the supplier knows or has reason to know was recalled by the U.S. Consumer Product Safety Commission. There is a rebuttable presumption that a supplier has reason to know a children's product was recalled if notice of the recall has been posted continuously at least 30 days before the sale, offer for sale, or manufacturing for sale on the website of the U.S. Consumer Product Safety Commission. This prohibition does not apply to children's products that are used, secondhand or "seconds";

50. Violating any provision of Chapter 44.1 (§ 59.1-518.1 et seq.);

51. Violating any provision of Chapter 22 (§ 6.2-2200 et seq.) of Title 6.2;

52. Violating any provision of § 8.2-317.1;

53. Violating subsection A of § 9.1-149.1;

54. Selling, offering for sale, or using in the construction, remodeling, or repair of any residential dwelling in the Commonwealth, any drywall that the supplier knows or has reason to know is defective drywall. This subdivision shall not apply to the sale or offering for sale of any building or structure in which defective drywall has been permanently installed or affixed;

55. Engaging in fraudulent or improper or dishonest conduct as defined in § 54.1-1118 while engaged in a transaction that was initiated (i) during a declared state of emergency as defined in § 44-146.16 or (ii) to repair damage resulting from the event that prompted the declaration of a state of emergency, regardless of whether the supplier is licensed as a contractor in the Commonwealth pursuant to Chapter 11 (§ 54.1-1100 et seq.) of Title 54.1;

56. Violating any provision of Chapter 33.1 (§ 59.1-434.1 et seq.);

57. Violating any provision of § 18.2-178, 18.2-178.1, or 18.2-200.1;

58. Violating any provision of Chapter 17.8 (§ 59.1-207.45 et seq.);

59. Violating any provision of subsection E of § 32.1-126;

60. Violating any provision of § 54.1-111 relating to the unlicensed practice of a profession licensed under Chapter 11 (§ 54.1-1100 et seq.) or Chapter 21 (§ 54.1-2100 et seq.) of Title 54.1;

61. Violating any provision of § 2.2-2001.5;

62. Violating any provision of Chapter 5.2 (§ 54.1-526 et seq.) of Title 54.1;

63. Violating any provision of § 6.2-312;

64. Violating any provision of Chapter 20.1 (§ 6.2-2026 et seq.) of Title 6.2;

65. Violating any provision of Chapter 26 (§ 6.2-2600 et seq.) of Title 6.2;

66. Violating any provision of Chapter 54 (§ 59.1-586 et seq.);

67. Knowingly violating any provision of § 8.01-27.5;

68. Failing to make available a conspicuous online option to cancel a recurring purchase of a good or service as required by § 59.1-207.46;

69. Selling or offering for sale to a person younger than 21 years of age any substance intended for human consumption, orally or by inhalation, that contains tetrahydrocannabinol. This subdivision shall not (i) apply to products that are approved for marketing by the U.S. Food and Drug Administration and scheduled in the Drug Control Act (§ 54.1-3400 et seq.) or (ii) be construed to prohibit any conduct permitted under Article 4.2 *(§ 54.1-3442.5 et seq.)* of Chapter 34 of Title 54.1 of the Code of Virginia;

70. Selling or offering for sale any substance intended for human consumption, orally or by inhalation, that contains tetrahydrocannabinol, unless such substance is (i) contained in child-resistant packaging, as defined in § 4.1-600; (ii) equipped with a label that states, in English and in a font no less than 1/16 of an inch, (a) that the substance contains tetrahydrocannabinol and may not be sold to persons younger than 21 years of age, (b) all ingredients contained in the substance, (c) the amount of such substance that constitutes a single serving, and (d) the total percentage and milligrams of tetrahydrocannabinol included in the substance and the number of milligrams of tetrahydrocannabinol that are contained in each serving; and (iii) accompanied by a certificate of analysis, produced by an independent laboratory that is *registered with the Virginia Cannabis Control Authority and* accredited pursuant to standard ISO/IEC 17025 of the International Organization of Standardization by a third-party accrediting body, that states the tetrahydrocannabinol concentration of the substance or the tetrahydrocannabinol concentration of the batch from which the substance originates. This subdivision shall not (i) apply to products that are approved for marketing by the U.S. Food and Drug Administration and scheduled in the Drug Control Act (§ 54.1-3400 et seq.) or (ii) be construed to prohibit any conduct permitted under Article 4.2 *(§ 54.1-3442.5 et seq.)* of Chapter 34 of Title 54.1 of the Code of Virginia;

71. Manufacturing, offering for sale at retail, or selling at retail an industrial hemp extract, as defined in § 3.2-5145.1, a food containing an industrial hemp extract, or a substance containing tetrahydrocannabinol that depicts or is in the shape of a human, animal, vehicle, or fruit; and

72. Selling or offering for sale any substance intended for human consumption, orally or by

inhalation, that contains tetrahydrocannabinol and, without authorization, bears, is packaged in a container or wrapper that bears, or is otherwise labeled to bear the trademark, trade name, famous mark as defined in 15 U.S.C. § 1125, or other identifying mark, imprint, or device, or any likeness thereof, of a manufacturer, processor, packer, or distributor of a product intended for human consumption other than the manufacturer, processor, packer, or distributor that did in fact so manufacture, process, pack, or distribute such substance.

B. Nothing in this section shall be construed to invalidate or make unenforceable any contract or lease solely by reason of the failure of such contract or lease to comply with any other law of the Commonwealth or any federal statute or regulation, to the extent such other law, statute, or regulation provides that a violation of such law, statute, or regulation shall not invalidate or make unenforceable such contract or lease.

§ 59.1-206. Civil penalties; attorney fees.

A. In any action brought under this chapter, if the court finds that a person has willfully engaged in an act or practice in violation of § 59.1-200 or 59.1-200.1, the Attorney General, the attorney for the Commonwealth, or the attorney for the county, city, or town may recover for the Literary Fund, upon petition to the court, a civil penalty of not more than $2,500 per violation. *If the court finds that a person has willfully committed a second or subsequent violation of subdivision A 69, 70, 71, or 72 of § 59.1-200, the Attorney General, the attorney for the Commonwealth, or the attorney for the county, city, or town may recover for the Literary Fund, upon petition to the court, a civil penalty of not more than $5,000 per violation.*

B. For purposes of this section, prima facie evidence of a willful violation may be shown when the Attorney General, the attorney for the Commonwealth, or the attorney for the county, city, or town notifies the alleged violator by certified mail that an act or practice is a violation of § 59.1-200 or 59.1-200.1, and the alleged violator, after receipt of said notice, continues to engage in the act or practice.

~~B.~~ *C.* Any person who willfully violates the terms of an assurance of voluntary compliance or an injunction issued under § 59.1-203 shall forfeit and pay to the Literary Fund a civil penalty of not more than $5,000 per violation. For purposes of this section, the circuit court issuing an injunction shall retain jurisdiction, and the cause shall be continued, and in such cases the Attorney General, the attorney for the Commonwealth, or the attorney for the county, city, or town may petition for recovery of civil penalties.

~~C.~~ *D.* In any action pursuant to subsection A or ~~B~~ *C* and in addition to any other amount awarded, the Attorney General, the attorney for the Commonwealth, or the attorney for the county, city, or town may recover any applicable civil penalty or penalties, costs, reasonable expenses incurred by the state or local agency in investigating and preparing the case not to exceed $1,000 per violation, and ~~attorney's~~ *attorney* fees. Such civil penalty or penalties, costs, reasonable expenses, and ~~attorney's~~ *attorney* fees shall be paid into the general fund of the Commonwealth or of the county, city, or town which such attorney represented.

~~D.~~ *E.* Nothing in this section shall be construed as limiting the power of the court to punish as contempt the violation of any order issued by the court, or as limiting the power of the court to enter other orders under § 59.1-203 or 59.1-205.

~~E.~~ *F.* The right of trial by jury as provided by law shall be preserved in actions brought under this section.

2. That the Board of Directors (the Board) of the Virginia Cannabis Control Authority shall promulgate regulations and establish a registration process to implement the provisions of subdivision A 70 of § 59.1-200 of the Code of Virginia, as amended by this act. The Board's initial adoption of such regulations shall be subject to the provisions of § 2.2-4031 of the Code of Virginia and exempt from all other provisions of the Administrative Process Act (§ 2.2-4000 et seq. of the Code of Virginia).

3. That the provisions of this act may result in a net increase in periods of imprisonment or commitment. Pursuant to § 30-19.1:4 of the Code of Virginia, the estimated amount of the necessary appropriation cannot be determined for periods of imprisonment in state adult correctional facilities; therefore, Chapter 2 of the Acts of Assembly of 2022, Special Session I, requires the Virginia Criminal Sentencing Commission to assign a minimum fiscal impact of $50,000. Pursuant to § 30-19.1:4 of the Code of Virginia, the estimated amount of the necessary appropriation cannot be determined for periods of commitment to the custody of the Department of Juvenile Justice.