§ 18.2-251.1. Possession or distribution of marijuana for medical purposes permitted

A. No person shall be prosecuted under § 18.2-250 or § 18.2-250.1 for the possession of marijuana or tetrahydrocannabinol when that possession occurs pursuant to a valid prescription issued by a medical doctor in the course of his professional practice for treatment of cancer or glaucoma.

B. No medical doctor shall be prosecuted under § 18.2-248 or § 18.2-248.1 for dispensing or distributing marijuana or tetrahydrocannabinol for medical purposes when such action occurs in the course of his professional practice for treatment of cancer or glaucoma.

C. No pharmacist shall be prosecuted under §§ 18.2-248 to 18.2-248.1 for dispensing or distributing marijuana or tetrahydrocannabinol to any person who holds a valid prescription of a medical doctor for such substance issued in the course of such doctor's professional practice for treatment of cancer or glaucoma.

1979, c. 435.

The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

Code of Virginia
Title 18.2. Crimes and Offenses Generally
Chapter 7. Crimes Involving Health and Safety
Article 1. Drugs

§ 18.2-248. Manufacturing, selling, giving, distributing, or possessing with intent to manufacture, sell, give, or distribute a controlled substance or an imitation controlled substance prohibited; penalties

A. Except as authorized in the Drug Control Act (§ 54.1-3400 et seq.), it shall be unlawful for any person to manufacture, sell, give, distribute, or possess with intent to manufacture, sell, give or distribute a controlled substance or an imitation controlled substance.

B. In determining whether any person intends to manufacture, sell, give or distribute an imitation controlled substance, the court may consider, in addition to all other relevant evidence, whether any distribution or attempted distribution of such pill, capsule, tablet or substance in any other form whatsoever included an exchange of or a demand for money or other property as consideration, and, if so, whether the amount of such consideration was substantially greater than the reasonable value of such pill, capsule, tablet or substance in any other form whatsoever, considering the actual chemical composition of such pill, capsule, tablet or substance in any other form whatsoever and, where applicable, the price at which over-the-counter substances of like chemical composition sell.

C. Except as provided in subsection C1, any person who violates this section with respect to a controlled substance classified in Schedule I or II shall upon conviction be imprisoned for not less than five nor more than 40 years and fined not more than $500,000. Upon a second conviction of such a violation, and it is alleged in the warrant, indictment, or information that the person has been before convicted of such an offense or of a substantially similar offense in any other jurisdiction, which offense would be a felony if committed in the Commonwealth, and such prior conviction occurred before the date of the offense alleged in the warrant, indictment, or information, any such person may, in the discretion of the court or jury imposing the sentence, be sentenced to imprisonment for life or for any period not less than five years, three years of which shall be a mandatory minimum term of imprisonment to be served consecutively with any other sentence, and he shall be fined not more than $500,000.

When a person is convicted of a third or subsequent offense under this subsection and it is alleged in the warrant, indictment or information that he has been before convicted of two or more such offenses or of substantially similar offenses in any other jurisdiction which offenses would be felonies if committed in the Commonwealth and such prior convictions occurred before the date of the offense alleged in the warrant, indictment, or information, he shall be sentenced to imprisonment for life or for a period of not less than 10 years, 10 years of which shall be a mandatory minimum term of imprisonment to be served consecutively with any other sentence, and he shall be fined not more than $500,000.

Any person who manufactures, sells, gives, distributes or possesses with the intent to manufacture, sell, give, or distribute the following is guilty of a felony punishable by a fine of not more than $1 million and imprisonment for five years to life, five years of which shall be a mandatory minimum term of imprisonment to be served consecutively with any other sentence:

1. 100 grams or more of a mixture or substance containing a detectable amount of heroin;

2. 500 grams or more of a mixture or substance containing a detectable amount of:

a. Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives of ecgonine or their salts have been removed;

b. Cocaine, its salts, optical and geometric isomers, and salts of isomers;

c. Ecgonine, its derivatives, their salts, isomers, and salts of isomers; or

d. Any compound, mixture, or preparation that contains any quantity of any of the substances referred to in subdivisions 2a through 2c;

3. 250 grams or more of a mixture or substance described in subdivisions 2a through 2d that contain cocaine base; or

4. 10 grams or more of methamphetamine, its salts, isomers, or salts of its isomers or 20 grams or more of a mixture or substance containing a detectable amount of methamphetamine, its salts, isomers, or salts of its isomers.

The mandatory minimum term of imprisonment to be imposed for a violation of this subsection shall not be applicable if the court finds that:

a. The person does not have a prior conviction for an offense listed in subsection C of § 17.1-805;

b. The person did not use violence or credible threats of violence or possess a firearm or other dangerous weapon in connection with the offense or induce another participant in the offense to do so;

c. The offense did not result in death or serious bodily injury to any person;

d. The person was not an organizer, leader, manager, or supervisor of others in the offense, and was not engaged in a continuing criminal enterprise as defined in subsection I; and

e. Not later than the time of the sentencing hearing, the person has truthfully provided to the Commonwealth all information and evidence the person has concerning the offense or offenses that were part of the same course of conduct or of a common scheme or plan, but the fact that the person has no relevant or useful other information to provide or that the Commonwealth already is aware of the information shall not preclude a determination by the court that the defendant has complied with this requirement.

C1. Any person who violates this section with respect to the manufacturing of methamphetamine, its salts, isomers, or salts of its isomers or less than 200 grams of a mixture or substance containing a detectable amount of methamphetamine, its salts, isomers, or salts of its isomers shall, upon conviction, be imprisoned for not less than 10 nor more than 40 years and fined not more than $500,000. Upon a second conviction of such a violation, any such person may, in the discretion of the court or jury imposing the sentence, be sentenced to imprisonment for life or for any period not less than 10 years, and be fined not more than $500,000. When a person is convicted of a third or subsequent offense under this subsection and it is alleged in the warrant, indictment, or information that he has been previously convicted of two or more such offenses or of substantially similar offenses in any other jurisdiction, which offenses would be felonies if committed in the Commonwealth and such prior convictions occurred before the date

of the offense alleged in the warrant, indictment, or information, he shall be sentenced to imprisonment for life or for a period not less than 10 years, three years of which shall be a mandatory minimum term of imprisonment to be served consecutively with any other sentence and he shall be fined not more than $500,000.

Upon conviction, in addition to any other punishment, a person found guilty of this offense shall be ordered by the court to make restitution, as the court deems appropriate, to any innocent property owner whose property is damaged, destroyed, or otherwise rendered unusable as a result of such methamphetamine production. This restitution shall include the person's or his estate's estimated or actual expenses associated with cleanup, removal, or repair of the affected property. If the property that is damaged, destroyed, or otherwise rendered unusable as a result of such methamphetamine production is property owned in whole or in part by the person convicted, the court shall order the person to pay to the Methamphetamine Cleanup Fund authorized in § 18.2-248.04 the reasonable estimated or actual expenses associated with cleanup, removal, or repair of the affected property or, if actual or estimated expenses cannot be determined, the sum of $10,000. The convicted person shall also pay the cost of certifying that any building that is cleaned up or repaired pursuant to this section is safe for human occupancy according to the guidelines established pursuant to § 32.1-11.7.

D. If such person proves that he gave, distributed or possessed with intent to give or distribute a controlled substance classified in Schedule I or II only as an accommodation to another individual who is not an inmate in a community correctional facility, local correctional facility or state correctional facility as defined in § 53.1-1 or in the custody of an employee thereof, and not with intent to profit thereby from any consideration received or expected nor to induce the recipient or intended recipient of the controlled substance to use or become addicted to or dependent upon such controlled substance, he shall be guilty of a Class 5 felony.

E. If the violation of the provisions of this article consists of the filling by a pharmacist of the prescription of a person authorized under this article to issue the same, which prescription has not been received in writing by the pharmacist prior to the filling thereof, and such written prescription is in fact received by the pharmacist within one week of the time of filling the same, or if such violation consists of a request by such authorized person for the filling by a pharmacist of a prescription which has not been received in writing by the pharmacist and such prescription is, in fact, written at the time of such request and delivered to the pharmacist within one week thereof, either such offense shall constitute a Class 4 misdemeanor.

E1. Any person who violates this section with respect to a controlled substance classified in Schedule III except for an anabolic steroid classified in Schedule III, constituting a violation of § 18.2-248.5, shall be guilty of a Class 5 felony.

E2. Any person who violates this section with respect to a controlled substance classified in Schedule IV shall be guilty of a Class 6 felony.

E3. Any person who proves that he gave, distributed or possessed with the intent to give or distribute a controlled substance classified in Schedule III or IV, except for an anabolic steroid classified in Schedule III, constituting a violation of § 18.2-248.5, only as an accommodation to another individual who is not an inmate in a community correctional facility, local correctional facility or state correctional facility as defined in § 53.1-1 or in the custody of an employee thereof, and not with the intent to profit thereby from any consideration received or expected nor to induce the recipient or intended recipient of the controlled substance to use or become

addicted to or dependent upon such controlled substance, is guilty of a Class 1 misdemeanor.

F. Any person who violates this section with respect to a controlled substance classified in Schedule V or Schedule VI or an imitation controlled substance which imitates a controlled substance classified in Schedule V or Schedule VI, shall be guilty of a Class 1 misdemeanor.

G. Any person who violates this section with respect to an imitation controlled substance which imitates a controlled substance classified in Schedule I, II, III, or IV shall be guilty of a Class 6 felony. In any prosecution brought under this subsection, it is not a defense to a violation of this subsection that the defendant believed the imitation controlled substance to actually be a controlled substance.

H. Any person who manufactures, sells, gives, distributes or possesses with the intent to manufacture, sell, give or distribute the following:

1. 1.0 kilograms or more of a mixture or substance containing a detectable amount of heroin;

2. 5.0 kilograms or more of a mixture or substance containing a detectable amount of:

a. Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives of ecgonine or their salts have been removed;

b. Cocaine, its salts, optical and geometric isomers, and salts of isomers;

c. Ecgonine, its derivatives, their salts, isomers, and salts of isomers; or

d. Any compound, mixture, or preparation which contains any quantity of any of the substances referred to in subdivisions a through c;

3. 2.5 kilograms or more of a mixture or substance described in subdivision 2 which contains cocaine base;

4. 100 kilograms or more of a mixture or substance containing a detectable amount of marijuana; or

5. 100 grams or more of methamphetamine, its salts, isomers, or salts of its isomers or 200 grams or more of a mixture or substance containing a detectable amount of methamphetamine, its salts, isomers, or salts of its isomers shall be guilty of a felony punishable by a fine of not more than $1 million and imprisonment for 20 years to life, 20 years of which shall be a mandatory minimum sentence. Such mandatory minimum sentence shall not be applicable if the court finds that (i) the person does not have a prior conviction for an offense listed in subsection C of § 17.1-805;(ii) the person did not use violence or credible threats of violence or possess a firearm or other dangerous weapon in connection with the offense or induce another participant in the offense to do so; (iii) the offense did not result in death or serious bodily injury to any person; (iv) the person was not an organizer, leader, manager, or supervisor of others in the offense, and was not engaged in a continuing criminal enterprise as defined in subsection I of this section; and (v) not later than the time of the sentencing hearing, the person has truthfully provided to the Commonwealth all information and evidence the person has concerning the offense or offenses that were part of the same course of conduct or of a common scheme or plan, but the fact that the person has no relevant or useful other information to provide or that the Commonwealth already is aware of the information shall not preclude a determination by the court that the defendant has complied with this requirement.

H1. Any person who was the principal or one of several principal administrators, organizers or leaders of a continuing criminal enterprise shall be guilty of a felony if (i) the enterprise received at least $100,000 but less than $250,000 in gross receipts during any 12-month period of its existence from the manufacture, importation, or distribution of heroin or cocaine or ecgonine or methamphetamine or the derivatives, salts, isomers, or salts of isomers thereof or marijuana or (ii) the person engaged in the enterprise to manufacture, sell, give, distribute or possess with the intent to manufacture, sell, give or distribute the following during any 12-month period of its existence:

1. At least 1.0 kilograms but less than 5.0 kilograms of a mixture or substance containing a detectable amount of heroin;

2. At least 5.0 kilograms but less than 10 kilograms of a mixture or substance containing a detectable amount of:

a. Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives of ecgonine or their salts have been removed;

b. Cocaine, its salts, optical and geometric isomers, and salts of isomers;

c. Ecgonine, its derivatives, their salts, isomers, and salts of isomers; or

d. Any compound, mixture, or preparation which contains any quantity of any of the substances referred to in subdivisions a through c;

3. At least 2.5 kilograms but less than 5.0 kilograms of a mixture or substance described in subdivision 2 which contains cocaine base;

4. At least 100 kilograms but less than 250 kilograms of a mixture or substance containing a detectable amount of marijuana; or

5. At least 100 grams but less than 250 grams of methamphetamine, its salts, isomers, or salts of its isomers or at least 200 grams but less than 1.0 kilograms of a mixture or substance containing a detectable amount of methamphetamine, its salts, isomers, or salts of its isomers.

A conviction under this section shall be punishable by a fine of not more than $1 million and imprisonment for 20 years to life, 20 years of which shall be a mandatory minimum sentence.

H2. Any person who was the principal or one of several principal administrators, organizers or leaders of a continuing criminal enterprise if (i) the enterprise received $250,000 or more in gross receipts during any 12-month period of its existence from the manufacture, importation, or distribution of heroin or cocaine or ecgonine or methamphetamine or the derivatives, salts, isomers, or salts of isomers thereof or marijuana or (ii) the person engaged in the enterprise to manufacture, sell, give, distribute or possess with the intent to manufacture, sell, give or distribute the following during any 12-month period of its existence:

1. At least 5.0 kilograms of a mixture or substance containing a detectable amount of heroin;

2. At least 10 kilograms of a mixture or substance containing a detectable amount of:

a. Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives of ecgonine or their salts have been removed;

b. Cocaine, its salts, optical and geometric isomers, and salts of isomers;

c. Ecgonine, its derivatives, their salts, isomers, and salts of isomers; or

d. Any compound, mixture, or preparation which contains any quantity of any of the substances referred to in subdivisions a through c;

3. At least 5.0 kilograms of a mixture or substance described in subdivision 2 which contains cocaine base;

4. At least 250 kilograms of a mixture or substance containing a detectable amount of marijuana; or

5. At least 250 grams of methamphetamine, its salts, isomers, or salts of its isomers or at least 1.0 kilograms of a mixture or substance containing a detectable amount of methamphetamine, its salts, isomers, or salts of its isomers shall be guilty of a felony punishable by a fine of not more than $1 million and imprisonment for life, which shall be served with no suspension in whole or in part. Such punishment shall be made to run consecutively with any other sentence. However, the court may impose a mandatory minimum sentence of 40 years if the court finds that the defendant substantially cooperated with law-enforcement authorities.

I. For purposes of this section, a person is engaged in a continuing criminal enterprise if (i) he violates any provision of this section, the punishment for which is a felony and either (ii) such violation is a part of a continuing series of violations of this section which are undertaken by such person in concert with five or more other persons with respect to whom such person occupies a position of organizer, a supervisory position, or any other position of management, and from which such person obtains substantial income or resources or (iii) such violation is committed, with respect to methamphetamine or other controlled substance classified in Schedule I or II, for the benefit of, at the direction of, or in association with any criminal street gang as defined in § 18.2-46.1.

J. Except as authorized in the Drug Control Act (§ 54.1-3400 et seq.), any person who possesses any two or more different substances listed below with the intent to manufacture methamphetamine, methcathinone, or amphetamine is guilty of a Class 6 felony: liquefied ammonia gas, ammonium nitrate, ether, hypophosphorus acid solutions, hypophosphite salts, hydrochloric acid, iodine crystals or tincture of iodine, phenylacetone, phenylacetic acid, red phosphorus, methylamine, methyl formamide, lithium, sodium metal, sulfuric acid, sodium hydroxide, potassium dichromate, sodium dichromate, potassium permanganate, chromium trioxide, methylbenzene, methamphetamine precursor drugs, trichloroethane, or 2-propanone.

K. The term "methamphetamine precursor drug," when used in this article, means a drug or product containing ephedrine, pseudoephedrine, or phenylpropanolamine or any of their salts, optical isomers, or salts of optical isomers.

Code 1950, § 54-524.101:1; 1972, c. 798; 1973, c. 479; 1974, c. 586; 1975, cc. 14, 15; 1976, c. 614; 1977, c. 409; 1978, cc. 177, 779; 1979, c. 435; 1982, cc. 276, 462; 1985, c. 569; 1986, c. 453; 1988, c. 355; 1990, c. 82; 1991, c. 13; 1992, cc. 685, 737, 756; 1995, c. 538;1999, c. 722;2000, cc. 1020, 1041;2004, c. 461;2005, cc. 174, 759, 796, 923, 941;2006, cc. 697, 759;2008, cc. 79, 618;2009, c. 750;2012, cc. 219, 710, 844;2013, c. 426;2014, c. 513.

The chapters of the acts of assembly referenced in the historical citation at the end of this

section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

§ 54.1-3408.3. (Effective until January 1, 2024) Certification for use of cannabis products 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 dispensing facility" means the same as that term is defined in § 54.1-3442.5.

"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, except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.), no more than 10 milligrams of tetrahydrocannabinol per dose. "Cannabis oil" does not include industrial hemp, as defined in § 3.2-4112, that is grown, handled, 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 (i) is formulated with cannabis oil or botanical cannabis; (ii) is produced by a pharmaceutical processor and sold by a pharmaceutical processor or cannabis dispensing facility; (iii) is registered with the Board; (iv) contains, except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.), no more than 10 milligrams of tetrahydrocannabinol per dose; and (v) is compliant with testing requirements.

"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.

"Pharmaceutical processor" means the same as that term is defined in § 54.1-3442.5.

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

"Registered agent" means an individual (i) 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 (ii) registered with the Board or listed on the patient's written certification 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. No practitioner may issue a written certification while such practitioner is on the premises of a pharmaceutical processor or cannabis dispensing facility. A pharmaceutical processor shall not endorse or promote any practitioner who issues certifications to patients. 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. A practitioner who issues written certifications shall not directly or indirectly accept, solicit, or receive anything of value from a pharmaceutical processor, cannabis dispensing facility, or any person associated with a pharmaceutical processor, cannabis dispensing facility, or provider of paraphernalia, excluding information on products or educational materials on the benefits and risks of cannabis products.

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 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 a practitioner's professional licensing board from sanctioning the 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 (i) shall hold sufficient education and training to exercise appropriate professional judgment in the certification of patients; (ii) shall not offer a discount or any other thing of value to a patient or a patient's parent, guardian, or registered agent that is contingent on or encourages the person's decision to use a particular pharmaceutical processor or cannabis product; (iii) shall not issue a certification to himself or his family members, employees, or coworkers; (iv) shall not provide product samples containing cannabis other than those approved by the U.S. Food and Drug Administration; and (v) shall not accept compensation from a pharmaceutical processor or cannabis dispensing facility. 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 unless the individual's name is listed on the patient's written certification. An individual may, on the basis of medical need and in the discretion of the patient's registered practitioner, be listed on the patient's written certification upon the patient's request. 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 patient certification or agent 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.

2015, cc. 7, 8;2017, c. 613;2018, cc. 246, 809;2019, cc. 653, 654, 681, 690;2020, cc. 730, 831, 928, 1278;2021 Sp. Sess. I, cc. 205, 227, 228;2022, cc. 259, 391, 392, 642;2023, cc. 183, 744, 760, 780, 794, 799.

This section has more than one version with varying effective dates. Scroll down to see all versions.
The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

§ 54.1-3408.3. (Effective January 1, 2024) Certification for use of cannabis for treatment

A. As used in this section, "botanical cannabis," "cannabis oil," "cannabis product," and "practitioner" mean the same as those terms are defined in § 4.1-1600.

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 in accordance with the provisions of § 4.1-1601.

2015, cc. 7, 8;2017, c. 613;2018, cc. 246, 809;2019, cc. 653, 654, 681, 690;2020, cc. 730, 831, 928, 1278;2021 Sp. Sess. I, cc. 205, 227, 228;2022, cc. 259, 391, 392, 642;2023, cc. 183, 740, 744, 760, 773, 780, 794, 799.

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Code of Virginia
Title 40.1. Labor and Employment
Chapter 3. Protection of Employees
Article 1. General Provisions

§ 40.1-27.4. (Effective until January 1, 2024) Discipline for employee's medicinal use of cannabis oil prohibited

A. As used in this section, "cannabis oil" means the same as that term is defined in § 54.1-3408.3.

B. No employer shall discharge, discipline, or discriminate against an employee for such employee's lawful use of cannabis oil pursuant to a valid written certification issued by a practitioner for the treatment or to eliminate the symptoms of the employee's diagnosed condition or disease pursuant to § 54.1-3408.3.

C. Notwithstanding the provisions of subsection B, nothing in this section shall (i) restrict an employer's ability to take any adverse employment action for any work impairment caused by the use of cannabis oil or to prohibit possession during work hours, (ii) require an employer to commit any act that would cause the employer to be in violation of federal law or that would result in the loss of a federal contract or federal funding, or (iii) require any defense industrial base sector employer or prospective employer, as defined by the U.S. Cybersecurity and Infrastructure Security Agency, to hire or retain any applicant or employee who tests positive for tetrahydrocannabinol (THC) in excess of 50 ng/ml for a urine test or 10 pg/mg for a hair test.

2021, Sp. Sess. I, c. 395.

This section has more than one version with varying effective dates. Scroll down to see all versions.
The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

§ 40.1-27.4. (Effective January 1, 2024) Discipline for employee's medicinal use of cannabis oil prohibited

A. As used in this section, "cannabis oil" means the same as that term is defined in § 4.1-1600.

B. No employer shall discharge, discipline, or discriminate against an employee for such employee's lawful use of cannabis oil pursuant to a valid written certification issued by a practitioner for the treatment or to eliminate the symptoms of the employee's diagnosed condition or disease pursuant to § 4.1-1601.

C. Notwithstanding the provisions of subsection B, nothing in this section shall (i) restrict an employer's ability to take any adverse employment action for any work impairment caused by the use of cannabis oil or to prohibit possession during work hours, (ii) require an employer to commit any act that would cause the employer to be in violation of federal law or that would result in the loss of a federal contract or federal funding, or (iii) require any defense industrial base sector employer or prospective employer, as defined by the U.S. Cybersecurity and Infrastructure Security Agency, to hire or retain any applicant or employee who tests positive for tetrahydrocannabinol (THC) in excess of 50 ng/ml for a urine test or 10 pg/mg for a hair test.

2021, Sp. Sess. I, c. 395;2023, cc. 740, 773.

This section has more than one version with varying effective dates. Scroll down to see all versions.
The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

Code of Virginia
Title 54.1. Professions and Occupations
Subtitle III. Professions and Occupations Regulated by Boards within the Department of Health Professions
Chapter 29. Medicine and Other Healing Arts
Article 1. General Provisions

§ 54.1-2903. (Effective until January 1, 2024) What constitutes practice; advertising in connection with medical practice

A. Any person shall be regarded as practicing the healing arts who actually engages in such practice as defined in this chapter, or who opens an office for such purpose, or who advertises or announces to the public in any manner a readiness to practice or who uses in connection with his name the words or letters "Doctor," "Dr.," "M.D.," "D.O.," "D.P.M.," "D.C.," "Healer," "N.P.," or any other title, word, letter or designation intending to designate or imply that he is a practitioner of the healing arts or that he is able to heal, cure or relieve those suffering from any injury, deformity or disease.

Signing a birth or death certificate, or signing any statement certifying that the person so signing has rendered professional service to the sick or injured, or signing or issuing a prescription for drugs or other remedial agents, shall be prima facie evidence that the person signing or issuing such writing is practicing the healing arts within the meaning of this chapter except where persons other than physicians are required to sign birth certificates.

B. No person regulated under this chapter shall use the title "Doctor" or the abbreviation "Dr." in writing or in advertising in connection with his practice unless he simultaneously uses words, initials, an abbreviation or designation, or other language that identifies the type of practice for which he is licensed. No person regulated under this chapter shall include in any advertisement a reference to marijuana, as defined in § 18.2-247, unless such advertisement is for the treatment of addiction or substance abuse. However, nothing in this subsection shall prevent a person from including in any advertisement that such person is registered with the Board of Pharmacy to issue written certifications for the use of cannabis products, as defined in § 54.1-3408.3.

Code 1950, § 54-275; 1958, c. 161; 1966, c. 657; 1973, c. 529; 1975, c. 508; 1988, c. 765; 1991, c. 102; 1996, cc. 937, 980;2000, c. 688;2018, c. 776;2019, c. 656;2021 Sp. Sess. I, cc. 227, 228.

This section has more than one version with varying effective dates. Scroll down to see all versions.
The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

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other title, word, letter or designation intending to designate or imply that he is a practitioner of the healing arts or that he is able to heal, cure or relieve those suffering from any injury, deformity or disease.

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Code 1950, § 54-275; 1958, c. 161; 1966, c. 657; 1973, c. 529; 1975, c. 508; 1988, c. 765; 1991, c. 102; 1996, cc. 937, 980;2000, c. 688;2018, c. 776;2019, c. 656;2021 Sp. Sess. I, cc. 227, 228;2023, cc. 740, 773.

The chapters of the acts of assembly referenced in the historical citation at the end of this section(s) may not constitute a comprehensive list of such chapters and may exclude chapters whose provisions have expired.

CHAPTER 773

An Act to amend and reenact §§ 4.1-604, 4.1-605, 4.1-627, 18.2-251.1:1, 18.2-251.1:2, 22.1-277, 32.1-127, 32.1-162.6:1, 40.1-27.4, 46.2-341.20:7, 54.1-2522.1, as it is currently effective and as it shall become effective, 54.1-2903, 54.1-3408.3, 59.1-200, and 63.2-1803.01 of the Code of Virginia; to amend the Code of Virginia by adding in Title 4.1 a chapter numbered 16, consisting of sections numbered 4.1-1600 through 4.1-1605; and to repeal Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia and the twenty-first enactment of Chapter 550 and the twenty-first enactment of Chapter 551 of the Acts of Assembly of 2021, Special Session I, relating to medical cannabis program; transition from Board of Pharmacy to Virginia Cannabis Control Authority.

[H 1598]

Approved April 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That §§ 4.1-604, 4.1-605, 4.1-627, 18.2-251.1:1, 18.2-251.1:2, 22.1-277, 32.1-127, 32.1-162.6:1, 40.1-27.4, 46.2-341.20:7, 54.1-2522.1, as it is currently effective and as it shall become effective, 54.1-2903, 54.1-3408.3, 59.1-200, and 63.2-1803.01 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding in Title 4.1 a chapter numbered 16, consisting of sections numbered 4.1-1600 through 4.1-1605, as follows:

§ 4.1-604. Powers and duties of the Board.

The Board shall have the following powers and duties:

1. Promulgate regulations in accordance with the Administrative Process Act (§ 2.2-4000 et seq.) and § 4.1-606;

2. Control the possession, sale, transportation, and delivery of marijuana and marijuana products;

3. Grant, suspend, ~~and~~ *restrict,* revoke ~~licenses for the cultivation, manufacture, distribution, sale, and testing of marijuana and marijuana products as provided by law~~, *or refuse to grant or renew any license or permit issued or authorized pursuant to this subtitle*;

4. Determine the nature, form, and capacity of all containers used for holding marijuana products to be kept or sold and prescribe the form and content of all labels and seals to be placed thereon;

5. Maintain actions to enjoin common nuisances as defined in § 4.1-1113;

6. Establish standards and implement an online course for employees of retail marijuana stores that trains employees on how to educate consumers on the potential risks of marijuana use;

7. Establish a plan to develop and disseminate to retail marijuana store licensees a pamphlet or similar document regarding the potential risks of marijuana use to be prominently displayed and made available to consumers;

8. Establish a position for a Cannabis Social Equity Liaison who shall lead the Cannabis Business Equity and Diversity Support Team and liaise with the Director of Diversity, Equity, and Inclusion on matters related to diversity, equity, and inclusion standards in the marijuana industry;

9. Establish a Cannabis Business Equity and Diversity Support Team, which shall (i) develop requirements for the creation and submission of diversity, equity, and inclusion plans by persons who wish to possess a license in more than one license category pursuant to subsection C of § 4.1-805, which may include a requirement that the licensee participate in social equity apprenticeship plan, and an approval process and requirements for implementation of such plans; (ii) be responsible for conducting an analysis of potential barriers to entry for small, women-owned, and minority-owned businesses and veteran-owned businesses interested in participating in the marijuana industry and recommending strategies to effectively mitigate such potential barriers; (iii) provide assistance with business planning for potential marijuana establishment licensees; (iv) spread awareness of business opportunities related to the marijuana marketplace in areas disproportionately impacted by marijuana prohibition and enforcement; (v) provide technical assistance in navigating the administrative process to potential marijuana establishment licensees; and (vi) conduct other outreach initiatives in areas disproportionately impacted by marijuana prohibition and enforcement as necessary;

10. Establish a position for an individual with professional experience in a health related field who shall staff the Cannabis Public Health Advisory Council, established pursuant to § 4.1-603, liaise with the Office of the Secretary of Health and Human Resources and relevant health and human services agencies and organizations, and perform other duties as needed.

11. Establish and implement a plan, in coordination with the Cannabis Social Equity Liaison and the Director of Diversity, Equity, and Inclusion to promote and encourage participation in the marijuana industry by people from communities that have been disproportionately impacted by marijuana prohibition and enforcement and to positively impact those communities;

12. Sue and be sued, implead and be impleaded, and complain and defend in all courts;

13. Adopt, use, and alter at will a common seal;

14. Fix, alter, charge, and collect rates, rentals, fees, and other charges for the use of property of, the sale of products of, or services rendered by the Authority at rates to be determined by the Authority for the purpose of providing for the payment of the expenses of the Authority;

15. Make and enter into all contracts and agreements necessary or incidental to the performance of its duties, the furtherance of its purposes, and the execution of its powers under this subtitle, including agreements with any person or federal agency;

16. Employ, at its discretion, consultants, researchers, architects, engineers, accountants, financial experts, investment bankers, superintendents, managers, and such other employees and special agents as may be necessary and fix their compensation to be payable from funds made available to the Authority. Legal services for the Authority shall be provided by the Attorney General in accordance with Chapter 5 (§ 2.2-500 et seq.) of Title 2.2;

17. Receive and accept from any federal or private agency, foundation, corporation, association, or person grants or other aid to be expended in accomplishing the objectives of the Authority, and receive and accept from the Commonwealth or any state and any municipality, county, or other political subdivision thereof or from any other source aid or contributions of either money, property, or other things of value, to be held, used, and applied only for the purposes for which such grants and contributions may be made. All federal moneys accepted under this section shall be accepted and expended by the Authority upon such terms and conditions as are prescribed by the United States and as are consistent with state law, and all state moneys accepted under this section shall be expended by the Authority upon such terms and conditions as are prescribed by the Commonwealth;

18. Adopt, alter, and repeal bylaws, rules, and regulations governing the manner in which its business shall be transacted and the manner in which the powers of the Authority shall be exercised and its duties performed. The Board may delegate or assign any duty or task to be performed by the Authority to any officer or employee of the Authority. The Board shall remain responsible for the performance of any such duties or tasks. Any delegation pursuant to this subdivision shall, where appropriate, be accompanied by written guidelines for the exercise of the duties or tasks delegated. Where appropriate, the guidelines shall require that the Board receive summaries of actions taken. Such delegation or assignment shall not relieve the Board of the responsibility to ensure faithful performance of the duties and tasks;

19. Conduct or engage in any lawful business, activity, effort, or project consistent with the Authority's purposes or necessary or convenient to exercise its powers;

20. Develop policies and procedures generally applicable to the procurement of goods, services, and construction, based upon competitive principles;

21. Develop policies and procedures consistent with Article 4 (§ 2.2-4347 et seq.) of Chapter 43 of Title 2.2;

22. Acquire, purchase, hold, use, lease, or otherwise dispose of any property, real, personal or mixed, tangible or intangible, or any interest therein necessary or desirable for carrying out the purposes of the Authority; lease as lessee any property, real, personal or mixed, tangible or intangible, or any interest therein, at such annual rental and on such terms and conditions as may be determined by the Board; lease as lessor to any person any property, real, personal or mixed, tangible or intangible, or any interest therein, at any time acquired by the Authority, whether wholly or partially completed, at such annual rental and on such terms and conditions as may be determined by the Board; sell, transfer, or convey any property, real, personal or mixed, tangible or intangible, or any interest therein, at any time acquired or held by the Authority on such terms and conditions as may be determined by the Board; and occupy and improve any land or building required for the purposes of this subtitle;

23. Purchase, lease, or acquire the use of, by any manner, any plant or equipment that may be considered necessary or useful in carrying into effect the purposes of this subtitle, including rectifying, blending, and processing plants;

24. Appoint every agent and employee required for its operations, require any or all of them to give bonds payable to the Commonwealth in such penalty as shall be fixed by the Board, and engage the services of experts and professionals;

25. Hold and conduct hearings, issue subpoenas requiring the attendance of witnesses and the production of records, memoranda, papers, and other documents before the Board or any agent of the Board, and administer oaths and take testimony thereunder. The Board may authorize any Board member or agent of the Board to hold and conduct hearings, issue subpoenas, administer oaths and take testimony thereunder, and decide cases, subject to final decision by the Board, on application of any party aggrieved. The Board may enter into consent agreements and may request and accept from any applicant ~~or~~, licensee, *or permittee* a consent agreement in lieu of proceedings on (i) objections to the issuance of a license *or permit* or (ii) disciplinary action. Any such consent agreement *(a)* shall include findings of fact *and provisions regarding whether the terms of the consent agreement are confidential* and *(b)* may include an admission or a finding of a violation. A consent agreement shall not be considered a case decision of the Board and shall not be subject to judicial review under the provisions

of the Administrative Process Act (§ 2.2-4000 et seq.), but may be considered by the Board in future disciplinary proceedings;

26. Make a reasonable charge for preparing and furnishing statistical information and compilations to persons other than (i) officials, including court and police officials, of the Commonwealth and of its subdivisions if the information requested is for official use and (ii) persons who have a personal or legal interest in obtaining the information requested if such information is not to be used for commercial or trade purposes;

27. ~~Assess~~ *Take appropriate disciplinary action and assess* and collect civil penalties and civil charges for violations of this subtitle and Board regulations;

28. Review and approve any proposed legislative or regulatory changes suggested by the Chief Executive Officer as the Board deems appropriate;

29. Report quarterly to the Secretary of Public Safety and Homeland Security on the law-enforcement activities undertaken to enforce the provisions of this subtitle;

30. Establish and collect fees for all permits set forth in this subtitle, including fees associated with applications for such permits;

31. Develop and make available on its website guidance documents regarding compliance and safe practices for persons who cultivate marijuana at home for personal use, which shall include information regarding cultivation practices that promote personal and public safety, including child protection, and discourage practices that create a nuisance;

32. Develop and make available on its website a resource that provides information regarding (i) responsible marijuana consumption; (ii) health risks and other dangers associated with marijuana consumption, including inability to operate a motor vehicle and other types of transportation and equipment; and (iii) ancillary effects of marijuana consumption, including ineligibility for certain employment opportunities. The Board shall require that the web address for such resource be included on the label of all retail marijuana and retail marijuana product as provided in § 4.1-1402; and

33. Do all acts necessary or advisable to carry out the purposes of this subtitle.

§ 4.1-605. Additional powers; mediation; alternative dispute resolution; confidentiality.

A. As used in this section:

"Appropriate case" means any alleged license *or permit* violation or objection to the application for a license *or permit* in which it is apparent that there are significant issues of disagreement among interested persons and for which the Board finds that the use of a mediation or dispute resolution proceeding is in the public interest.

"Dispute resolution proceeding" means the same as that term is defined in § 8.01-576.4.

"Mediation" means the same as that term is defined in § 8.01-576.4.

"Neutral" means the same as that term is defined in § 8.01-576.4.

B. The Board may use mediation or a dispute resolution proceeding in appropriate cases to resolve underlying issues or reach a consensus or compromise on contested issues. Mediation and other dispute resolution proceedings as authorized by this section shall be voluntary procedures that supplement, rather than limit, other dispute resolution techniques available to the Board. Mediation or a dispute resolution proceeding may be used for an objection to the issuance of a license *or permit* only with the consent of, and participation by, the applicant for ~~licensure~~ *a license or permit* and shall be terminated at the request of such applicant.

C. Any resolution of a contested issue accepted by the Board under this section shall be considered a consent agreement as provided in § 4.1-604. The decision to use mediation or a dispute resolution proceeding is in the Board's sole discretion and shall not be subject to judicial review.

D. The Board may adopt rules and regulations, in accordance with the Administrative Process Act (§ 2.2-4000 et seq.), for the implementation of this section. Such rules and regulations may include (i) standards and procedures for the conduct of mediation and dispute resolution proceedings, including an opportunity for interested persons identified by the Board to participate in the proceeding; (ii) the appointment and function of a neutral to encourage and assist parties to voluntarily compromise or settle contested issues; and (iii) procedures to protect the confidentiality of papers, work products, or other materials.

E. The provisions of § 8.01-576.10 concerning the confidentiality of a mediation or dispute resolution proceeding shall govern all such proceedings held pursuant to this section except where the Board uses or relies on information obtained in the course of such proceeding in granting ~~a~~ ~~license~~, suspending, *restricting,* or revoking a license *or permit*, or *in* accepting payment of a civil penalty or investigative costs. ~~However,~~ ~~a~~ ~~consent~~ ~~agreement~~ *Consent agreements shall be* signed by ~~the~~ *all* parties *and* shall ~~not be~~ *include provisions regarding whether the terms of the consent agreement are* confidential.

§ 4.1-627. Hearings; representation by counsel.

Any licensee, *permittee,* or applicant for ~~any~~ *a* license ~~granted by the Board~~ *or permit authorized by this subtitle* shall have the right to be represented by counsel at any Board hearing for which he has received notice. The licensee, *permittee,* or applicant shall not be required to be represented by counsel during such hearing. Any officer or director of a corporation may examine, cross-examine, and question witnesses, present evidence on behalf of the corporation, and draw conclusions and make arguments

before the Board or hearing officers without being in violation of the provisions of § 54.1-3904.

CHAPTER 16.

MEDICAL CANNABIS PROGRAM.

§ 4.1-1600. Definitions.

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

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

"Cannabis dispensing facility" means a facility that (i) has obtained a permit from the Board pursuant to § 4.1-1602; (ii) is owned, at least in part, by a pharmaceutical processor; and (iii) dispenses cannabis products produced by a pharmaceutical processor 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.

"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 § 4.1-1602, or a dilution of the resin of the Cannabis plant that contains no more than 10 milligrams of delta-9-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.

"Dispense" means the same as that term is defined in § 54.1-3300.

"Pharmaceutical processor" means a facility that (i) has obtained a permit from the Board pursuant to § 4.1-1602 and (ii) cultivates Cannabis plants intended only for the production of cannabis oil, botanical cannabis, and usable cannabis, produces cannabis products, and dispenses cannabis products to a patient pursuant to a written certification, 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.

"Pharmacist" means the same as that term is defined in § 54.1-3300.

"Pharmacy intern" means the same as that term is defined in § 54.1-3300.

"Pharmacy technician" means the same as that term is defined in § 54.1-3300.

"Pharmacy technician trainee" means the same as that term is defined in § 54.1-3300.

"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 Boards of Nursing and Medicine.

"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 F of § 4.1-1601.

"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, derivative, mixture, or preparation of the mature stalks; or (iii) oil or cake made from the seeds of the plant.

§ 4.1-1601. Certification for use of cannabis for treatment.

A. 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 audiovisual 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.

B. The written certification shall be on a form provided by the Authority. 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 A 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.

C. 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 A. 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.

D. 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.

E. 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 Authority.

F. 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.

G. 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.

H. 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 patient's registered agent, but only with respect to information related to such patient.

§ 4.1-1602. 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 Authority 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; (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 processor 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 N; (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 products.

D. The Board shall require pharmaceutical processors, after processing and before dispensing any cannabis products, to make a sample available from each batch of cannabis product for testing by an independent laboratory that is located in Commonwealth and meets 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 of Pharmacy 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.

H. 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.

I. 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.

J. 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.

K. 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.

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

M. 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.

N. 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.

O. 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 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 shall consider and keep on file all public comments received for any regulation adopted pursuant to this section.

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

§ 4.1-1603. 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 § 4.1-1602. 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 on General Laws and the Senate Committee on Rehabilitation and Social Services on the operation of pharmaceutical processors and cannabis dispensing facilities issued a permit by the Board.

D. The concentration of delta-9-tetrahydrocannabinol in any cannabis product on site may be up to 10 percent greater than or less than the level of delta-9-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.

§ 4.1-1604. Criminal liability; exceptions.

No agent or employee of a pharmaceutical processor or cannabis dispensing facility shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) or § 18.2-248, 18.2-248.1, or 18.2-250 for possession or manufacture of marijuana or for possession, manufacture, or distribution of cannabis products, subject to any civil penalty, denied any right or privilege, or subject to any disciplinary action by a professional licensing board if such agent or employee (i) possessed or manufactured such marijuana for the purposes of producing cannabis products in accordance with the provisions of this chapter and Board regulations or (ii) possessed, manufactured, or distributed such cannabis products that are consistent with generally accepted cannabis industry standards in accordance with the provisions of this chapter and Board regulations.

§ 4.1-1605. Summary suspensions and restrictions.

A. The Board may summarily suspend or restrict a permit issued pursuant to § 4.1-1602 without a hearing if the Board finds that such suspension or restriction is necessary to prevent substantial danger to public health or safety. The Board shall make decisions to summarily suspend or restrict a permit only during an in-person meeting in which a quorum is present; however, if, after a good faith effort, the Board is unable to assemble a quorum and a majority of the Board members determine that continued operation by the permittee constitutes a substantial danger to public health or safety, the Board may summarily suspend the permit during a telephone, video, or other electronic conference. Institution of proceedings for a hearing shall be provided simultaneously with a summary suspension. The Board may summarily restrict a permit without proceeding simultaneously with notification of an informal conference pursuant to § 2.2-4019 or Board regulations. Such hearing or conference shall be held within a reasonable amount of time after the summary suspension or restriction is issued.

B. Allegations of violations of this subtitle shall be submitted to the Board in writing.

§ 18.2-251.1:1. Possession or distribution of cannabis oil; public schools.

No school nurse employed by a local school board, person employed by a local health department who is assigned to the public school pursuant to an agreement between the local health department and the school board, or other person employed by or contracted with a local school board to deliver health-related services shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-248, 18.2-248.1, 18.2-250, or 18.2-255 for the possession or distribution of cannabis oil for storing, dispensing, or administering cannabis oil, in accordance with a policy adopted by the local school board, to a student who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601*.

§ 18.2-251.1:2. Possession or distribution of cannabis oil; nursing homes and certified nursing facilities; hospice and hospice facilities; assisted living facilities.

No person employed by a nursing home, hospice, hospice facility, or assisted living facility and authorized to possess, distribute, or administer medications to patients or residents shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-248, 18.2-248.1, or 18.2-250 for the possession or distribution of cannabis oil for the purposes of storing, dispensing, or administering cannabis oil to a patient or resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*.

§ 22.1-277. Suspensions and expulsions of students generally.

A. Students may be suspended or expelled from attendance at school for sufficient cause; however, in no cases may sufficient cause for suspensions include only instances of truancy.

B. Except as provided in subsection C or § 22.1-277.07 or 22.1-277.08, no student in preschool through grade three shall be suspended for more than three school days or expelled from attendance at school, unless (i) the offense involves physical harm or credible threat of physical harm to others or (ii)

the local school board or the division superintendent or his designee finds that aggravating circumstances exist, as defined by the Department.

C. Any student for whom the division superintendent of the school division in which such student is enrolled has received a report pursuant to § 16.1-305.1 of an adjudication of delinquency or a conviction for an offense listed in subsection G of § 16.1-260 may be suspended or expelled from school attendance pursuant to this article.

D. The authority provided in § 22.1-276.2 for teachers to remove students from their classes in certain instances of disruptive behavior shall not be interpreted to affect the operation of § 22.1-277.04, 22.1-277.05, or 22.1-277.06.

E. Notwithstanding the provisions of § 22.1-277.08, no school board shall be required to suspend or expel any student who holds a valid written certification for the use of cannabis oil issued by a practitioner in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601* for the possession or use of such oil in accordance with the student's individualized health plan and in compliance with a policy adopted by the school board.

§ 32.1-127. Regulations.

A. The regulations promulgated by the Board to carry out the provisions of this article shall be in substantial conformity to the standards of health, hygiene, sanitation, construction and safety as established and recognized by medical and health care professionals and by specialists in matters of public health and safety, including health and safety standards established under provisions of Title XVIII and Title XIX of the Social Security Act, and to the provisions of Article 2 (§ 32.1-138 et seq.).

B. Such regulations:

1. Shall include minimum standards for (i) the construction and maintenance of hospitals, nursing homes and certified nursing facilities to ensure the environmental protection and the life safety of its patients, employees, and the public; (ii) the operation, staffing and equipping of hospitals, nursing homes and certified nursing facilities; (iii) qualifications and training of staff of hospitals, nursing homes and certified nursing facilities, except those professionals licensed or certified by the Department of Health Professions; (iv) conditions under which a hospital or nursing home may provide medical and nursing services to patients in their places of residence; and (v) policies related to infection prevention, disaster preparedness, and facility security of hospitals, nursing homes, and certified nursing facilities;

2. Shall provide that at least one physician who is licensed to practice medicine in this Commonwealth shall be on call at all times, though not necessarily physically present on the premises, at each hospital which operates or holds itself out as operating an emergency service;

3. May classify hospitals and nursing homes by type of specialty or service and may provide for licensing hospitals and nursing homes by bed capacity and by type of specialty or service;

4. Shall also require that each hospital establish a protocol for organ donation, in compliance with federal law and the regulations of the Centers for Medicare and Medicaid Services (CMS), particularly 42 C.F.R. § 482.45. Each hospital shall have an agreement with an organ procurement organization designated in CMS regulations for routine contact, whereby the provider's designated organ procurement organization certified by CMS (i) is notified in a timely manner of all deaths or imminent deaths of patients in the hospital and (ii) is authorized to determine the suitability of the decedent or patient for organ donation and, in the absence of a similar arrangement with any eye bank or tissue bank in Virginia certified by the Eye Bank Association of America or the American Association of Tissue Banks, the suitability for tissue and eye donation. The hospital shall also have an agreement with at least one tissue bank and at least one eye bank to cooperate in the retrieval, processing, preservation, storage, and distribution of tissues and eyes to ensure that all usable tissues and eyes are obtained from potential donors and to avoid interference with organ procurement. The protocol shall ensure that the hospital collaborates with the designated organ procurement organization to inform the family of each potential donor of the option to donate organs, tissues, or eyes or to decline to donate. The individual making contact with the family shall have completed a course in the methodology for approaching potential donor families and requesting organ or tissue donation that (a) is offered or approved by the organ procurement organization and designed in conjunction with the tissue and eye bank community and (b) encourages discretion and sensitivity according to the specific circumstances, views, and beliefs of the relevant family. In addition, the hospital shall work cooperatively with the designated organ procurement organization in educating the staff responsible for contacting the organ procurement organization's personnel on donation issues, the proper review of death records to improve identification of potential donors, and the proper procedures for maintaining potential donors while necessary testing and placement of potential donated organs, tissues, and eyes takes place. This process shall be followed, without exception, unless the family of the relevant decedent or patient has expressed opposition to organ donation, the chief administrative officer of the hospital or his designee knows of such opposition, and no donor card or other relevant document, such as an advance directive, can be found;

5. Shall require that each hospital that provides obstetrical services establish a protocol for admission or transfer of any pregnant woman who presents herself while in labor;

6. Shall also require that each licensed hospital develop and implement a protocol requiring written discharge plans for identified, substance-abusing, postpartum women and their infants. The protocol shall

require that the discharge plan be discussed with the patient and that appropriate referrals for the mother and the infant be made and documented. Appropriate referrals may include, but need not be limited to, treatment services, comprehensive early intervention services for infants and toddlers with disabilities and their families pursuant to Part H of the Individuals with Disabilities Education Act, 20 U.S.C. § 1471 et seq., and family-oriented prevention services. The discharge planning process shall involve, to the extent possible, the other parent of the infant and any members of the patient's extended family who may participate in the follow-up care for the mother and the infant. Immediately upon identification, pursuant to § 54.1-2403.1, of any substance-abusing, postpartum woman, the hospital shall notify, subject to federal law restrictions, the community services board of the jurisdiction in which the woman resides to appoint a discharge plan manager. The community services board shall implement and manage the discharge plan;

7. Shall require that each nursing home and certified nursing facility fully disclose to the applicant for admission the home's or facility's admissions policies, including any preferences given;

8. Shall require that each licensed hospital establish a protocol relating to the rights and responsibilities of patients which shall include a process reasonably designed to inform patients of such rights and responsibilities. Such rights and responsibilities of patients, a copy of which shall be given to patients on admission, shall be consistent with applicable federal law and regulations of the Centers for Medicare and Medicaid Services;

9. Shall establish standards and maintain a process for designation of levels or categories of care in neonatal services according to an applicable national or state-developed evaluation system. Such standards may be differentiated for various levels or categories of care and may include, but need not be limited to, requirements for staffing credentials, staff/patient ratios, equipment, and medical protocols;

10. Shall require that each nursing home and certified nursing facility train all employees who are mandated to report adult abuse, neglect, or exploitation pursuant to § 63.2-1606 on such reporting procedures and the consequences for failing to make a required report;

11. Shall permit hospital personnel, as designated in medical staff bylaws, rules and regulations, or hospital policies and procedures, to accept emergency telephone and other verbal orders for medication or treatment for hospital patients from physicians, and other persons lawfully authorized by state statute to give patient orders, subject to a requirement that such verbal order be signed, within a reasonable period of time not to exceed 72 hours as specified in the hospital's medical staff bylaws, rules and regulations or hospital policies and procedures, by the person giving the order, or, when such person is not available within the period of time specified, co-signed by another physician or other person authorized to give the order;

12. Shall require, unless the vaccination is medically contraindicated or the resident declines the offer of the vaccination, that each certified nursing facility and nursing home provide or arrange for the administration to its residents of (i) an annual vaccination against influenza and (ii) a pneumococcal vaccination, in accordance with the most recent recommendations of the Advisory Committee on Immunization Practices of the Centers for Disease Control and Prevention;

13. Shall require that each nursing home and certified nursing facility register with the Department of State Police to receive notice of the registration, reregistration, or verification of registration information of any person required to register with the Sex Offender and Crimes Against Minors Registry pursuant to Chapter 9 (§ 9.1-900 et seq.) of Title 9.1 within the same or a contiguous zip code area in which the home or facility is located, pursuant to § 9.1-914;

14. Shall require that each nursing home and certified nursing facility ascertain, prior to admission, whether a potential patient is required to register with the Sex Offender and Crimes Against Minors Registry pursuant to Chapter 9 (§ 9.1-900 et seq.) of Title 9.1, if the home or facility anticipates the potential patient will have a length of stay greater than three days or in fact stays longer than three days;

15. Shall require that each licensed hospital include in its visitation policy a provision allowing each adult patient to receive visits from any individual from whom the patient desires to receive visits, subject to other restrictions contained in the visitation policy including, but not limited to, those related to the patient's medical condition and the number of visitors permitted in the patient's room simultaneously;

16. Shall require that each nursing home and certified nursing facility shall, upon the request of the facility's family council, send notices and information about the family council mutually developed by the family council and the administration of the nursing home or certified nursing facility, and provided to the facility for such purpose, to the listed responsible party or a contact person of the resident's choice up to six times per year. Such notices may be included together with a monthly billing statement or other regular communication. Notices and information shall also be posted in a designated location within the nursing home or certified nursing facility. No family member of a resident or other resident representative shall be restricted from participating in meetings in the facility with the families or resident representatives of other residents in the facility;

17. Shall require that each nursing home and certified nursing facility maintain liability insurance coverage in a minimum amount of $1 million, and professional liability coverage in an amount at least

equal to the recovery limit set forth in § 8.01-581.15, to compensate patients or individuals for injuries and losses resulting from the negligent or criminal acts of the facility. Failure to maintain such minimum insurance shall result in revocation of the facility's license;

18. Shall require each hospital that provides obstetrical services to establish policies to follow when a stillbirth, as defined in § 32.1-69.1, occurs that meet the guidelines pertaining to counseling patients and their families and other aspects of managing stillbirths as may be specified by the Board in its regulations;

19. Shall require each nursing home to provide a full refund of any unexpended patient funds on deposit with the facility following the discharge or death of a patient, other than entrance-related fees paid to a continuing care provider as defined in § 38.2-4900, within 30 days of a written request for such funds by the discharged patient or, in the case of the death of a patient, the person administering the person's estate in accordance with the Virginia Small Estates Act (§ 64.2-600 et seq.);

20. Shall require that each hospital that provides inpatient psychiatric services establish a protocol that requires, for any refusal to admit (i) a medically stable patient referred to its psychiatric unit, direct verbal communication between the on-call physician in the psychiatric unit and the referring physician, if requested by such referring physician, and prohibits on-call physicians or other hospital staff from refusing a request for such direct verbal communication by a referring physician and (ii) a patient for whom there is a question regarding the medical stability or medical appropriateness of admission for inpatient psychiatric services due to a situation involving results of a toxicology screening, the on-call physician in the psychiatric unit to which the patient is sought to be transferred to participate in direct verbal communication, either in person or via telephone, with a clinical toxicologist or other person who is a Certified Specialist in Poison Information employed by a poison control center that is accredited by the American Association of Poison Control Centers to review the results of the toxicology screen and determine whether a medical reason for refusing admission to the psychiatric unit related to the results of the toxicology screen exists, if requested by the referring physician;

21. Shall require that each hospital that is equipped to provide life-sustaining treatment shall develop a policy governing determination of the medical and ethical appropriateness of proposed medical care, which shall include (i) a process for obtaining a second opinion regarding the medical and ethical appropriateness of proposed medical care in cases in which a physician has determined proposed care to be medically or ethically inappropriate; (ii) provisions for review of the determination that proposed medical care is medically or ethically inappropriate by an interdisciplinary medical review committee and a determination by the interdisciplinary medical review committee regarding the medical and ethical appropriateness of the proposed health care; and (iii) requirements for a written explanation of the decision reached by the interdisciplinary medical review committee, which shall be included in the patient's medical record. Such policy shall ensure that the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986 (a) are informed of the patient's right to obtain his medical record and to obtain an independent medical opinion and (b) afforded reasonable opportunity to participate in the medical review committee meeting. Nothing in such policy shall prevent the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986 from obtaining legal counsel to represent the patient or from seeking other remedies available at law, including seeking court review, provided that the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986, or legal counsel provides written notice to the chief executive officer of the hospital within 14 days of the date on which the physician's determination that proposed medical treatment is medically or ethically inappropriate is documented in the patient's medical record;

22. Shall require every hospital with an emergency department to establish protocols to ensure that security personnel of the emergency department, if any, receive training appropriate to the populations served by the emergency department, which may include training based on a trauma-informed approach in identifying and safely addressing situations involving patients or other persons who pose a risk of harm to themselves or others due to mental illness or substance abuse or who are experiencing a mental health crisis;

23. Shall require that each hospital establish a protocol requiring that, before a health care provider arranges for air medical transportation services for a patient who does not have an emergency medical condition as defined in 42 U.S.C. § 1395dd(e)(1), the hospital shall provide the patient or his authorized representative with written or electronic notice that the patient (i) may have a choice of transportation by an air medical transportation provider or medically appropriate ground transportation by an emergency medical services provider and (ii) will be responsible for charges incurred for such transportation in the event that the provider is not a contracted network provider of the patient's health insurance carrier or such charges are not otherwise covered in full or in part by the patient's health insurance plan;

24. Shall establish an exemption from the requirement to obtain a license to add temporary beds in an existing hospital or nursing home, including beds located in a temporary structure or satellite location operated by the hospital or nursing home, provided that the ability remains to safely staff services across the existing hospital or nursing home, (i) for a period of no more than the duration of the Commissioner's determination plus 30 days when the Commissioner has determined that a natural or man-made disaster has caused the evacuation of a hospital or nursing home and that a public health

emergency exists due to a shortage of hospital or nursing home beds or (ii) for a period of no more than the duration of the emergency order entered pursuant to § 32.1-13 or 32.1-20 plus 30 days when the Board, pursuant to § 32.1-13, or the Commissioner, pursuant to § 32.1-20, has entered an emergency order for the purpose of suppressing a nuisance dangerous to public health or a communicable, contagious, or infectious disease or other danger to the public life and health;

25. Shall establish protocols to ensure that any patient scheduled to receive an elective surgical procedure for which the patient can reasonably be expected to require outpatient physical therapy as a follow-up treatment after discharge is informed that he (i) is expected to require outpatient physical therapy as a follow-up treatment and (ii) will be required to select a physical therapy provider prior to being discharged from the hospital;

26. Shall permit nursing home staff members who are authorized to possess, distribute, or administer medications to residents to store, dispense, or administer cannabis oil to a resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*;

27. Shall require each hospital with an emergency department to establish a protocol for the treatment and discharge of individuals experiencing a substance use-related emergency, which shall include provisions for (i) appropriate screening and assessment of individuals experiencing substance use-related emergencies to identify medical interventions necessary for the treatment of the individual in the emergency department and (ii) recommendations for follow-up care following discharge for any patient identified as having a substance use disorder, depression, or mental health disorder, as appropriate, which may include, for patients who have been treated for substance use-related emergencies, including opioid overdose, or other high-risk patients, (a) the dispensing of naloxone or other opioid antagonist used for overdose reversal pursuant to subsection X of § 54.1-3408 at discharge or (b) issuance of a prescription for and information about accessing naloxone or other opioid antagonist used for overdose reversal, including information about accessing naloxone or other opioid antagonist used for overdose reversal at a community pharmacy, including any outpatient pharmacy operated by the hospital, or through a community organization or pharmacy that may dispense naloxone or other opioid antagonist used for overdose reversal without a prescription pursuant to a statewide standing order. Such protocols may also provide for referrals of individuals experiencing a substance use-related emergency to peer recovery specialists and community-based providers of behavioral health services, or to providers of pharmacotherapy for the treatment of drug or alcohol dependence or mental health diagnoses;

28. During a public health emergency related to COVID-19, shall require each nursing home and certified nursing facility to establish a protocol to allow each patient to receive visits, consistent with guidance from the Centers for Disease Control and Prevention and as directed by the Centers for Medicare and Medicaid Services and the Board. Such protocol shall include provisions describing (i) the conditions, including conditions related to the presence of COVID-19 in the nursing home, certified nursing facility, and community, under which in-person visits will be allowed and under which in-person visits will not be allowed and visits will be required to be virtual; (ii) the requirements with which in-person visitors will be required to comply to protect the health and safety of the patients and staff of the nursing home or certified nursing facility; (iii) the types of technology, including interactive audio or video technology, and the staff support necessary to ensure visits are provided as required by this subdivision; and (iv) the steps the nursing home or certified nursing facility will take in the event of a technology failure, service interruption, or documented emergency that prevents visits from occurring as required by this subdivision. Such protocol shall also include (a) a statement of the frequency with which visits, including virtual and in-person, where appropriate, will be allowed, which shall be at least once every 10 calendar days for each patient; (b) a provision authorizing a patient or the patient's personal representative to waive or limit visitation, provided that such waiver or limitation is included in the patient's health record; and (c) a requirement that each nursing home and certified nursing facility publish on its website or communicate to each patient or the patient's authorized representative, in writing or via electronic means, the nursing home's or certified nursing facility's plan for providing visits to patients as required by this subdivision;

29. Shall require each hospital, nursing home, and certified nursing facility to establish and implement policies to ensure the permissible access to and use of an intelligent personal assistant provided by a patient, in accordance with such regulations, while receiving inpatient services. Such policies shall ensure protection of health information in accordance with the requirements of the federal Health Insurance Portability and Accountability Act of 1996, 42 U.S.C. § 1320d et seq., as amended. For the purposes of this subdivision, "intelligent personal assistant" means a combination of an electronic device and a specialized software application designed to assist users with basic tasks using a combination of natural language processing and artificial intelligence, including such combinations known as "digital assistants" or "virtual assistants";

30. During a declared public health emergency related to a communicable disease of public health threat, shall require each hospital, nursing home, and certified nursing facility to establish a protocol to allow patients to receive visits from a rabbi, priest, minister, or clergy of any religious denomination or sect consistent with guidance from the Centers for Disease Control and Prevention and the Centers for

Medicare and Medicaid Services and subject to compliance with any executive order, order of public health, Department guidance, or any other applicable federal or state guidance having the effect of limiting visitation. Such protocol may restrict the frequency and duration of visits and may require visits to be conducted virtually using interactive audio or video technology. Any such protocol may require the person visiting a patient pursuant to this subdivision to comply with all reasonable requirements of the hospital, nursing home, or certified nursing facility adopted to protect the health and safety of the person, patients, and staff of the hospital, nursing home, or certified nursing facility; and

31. Shall require that every hospital that makes health records, as defined in § 32.1-127.1:03, of patients who are minors available to such patients through a secure website shall make such health records available to such patient's parent or guardian through such secure website, unless the hospital cannot make such health record available in a manner that prevents disclosure of information, the disclosure of which has been denied pursuant to subsection F of § 32.1-127.1:03 or for which consent required in accordance with subsection E of § 54.1-2969 has not been provided.

C. Upon obtaining the appropriate license, if applicable, licensed hospitals, nursing homes, and certified nursing facilities may operate adult day care centers.

D. All facilities licensed by the Board pursuant to this article which provide treatment or care for hemophiliacs and, in the course of such treatment, stock clotting factors, shall maintain records of all lot numbers or other unique identifiers for such clotting factors in order that, in the event the lot is found to be contaminated with an infectious agent, those hemophiliacs who have received units of this contaminated clotting factor may be apprised of this contamination. Facilities which have identified a lot that is known to be contaminated shall notify the recipient's attending physician and request that he notify the recipient of the contamination. If the physician is unavailable, the facility shall notify by mail, return receipt requested, each recipient who received treatment from a known contaminated lot at the individual's last known address.

E. Hospitals in the Commonwealth may enter into agreements with the Department of Health for the provision to uninsured patients of naloxone or other opioid antagonists used for overdose reversal.

§ 32.1-162.6:1. Possession or administration of cannabis oil.

Hospice and hospice facility employees who are authorized to possess, distribute, or administer medications to patients shall be permitted to store, dispense, or administer cannabis oil to a patient who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*.

§ 40.1-27.4. Discipline for employee's medicinal use of cannabis oil prohibited.

A. As used in this section, "cannabis oil" means the same as that term is defined in § ~~54.1-3408.3~~ *4.1-1600*.

B. No employer shall discharge, discipline, or discriminate against an employee for such employee's lawful use of cannabis oil pursuant to a valid written certification issued by a practitioner for the treatment or to eliminate the symptoms of the employee's diagnosed condition or disease pursuant to § ~~54.1-3408.3~~ *4.1-1601*.

C. Notwithstanding the provisions of subsection B, nothing in this section shall (i) restrict an employer's ability to take any adverse employment action for any work impairment caused by the use of cannabis oil or to prohibit possession during work hours, (ii) require an employer to commit any act that would cause the employer to be in violation of federal law or that would result in the loss of a federal contract or federal funding, or (iii) require any defense industrial base sector employer or prospective employer, as defined by the U.S. Cybersecurity and Infrastructure Security Agency, to hire or retain any applicant or employee who tests positive for tetrahydrocannabinol (THC) in excess of 50 ng/ml for a urine test or 10 pg/mg for a hair test.

§ 46.2-341.20:7. Possession of marijuana in commercial motor vehicle unlawful; civil penalty.

A. It is unlawful for any person to knowingly or intentionally possess marijuana in a commercial motor vehicle as defined in § 46.2-341.4. The attorney for the Commonwealth or the county, city, or town attorney may prosecute such a case.

Upon the prosecution of a person for a violation of this section, ownership or occupancy of the vehicle in which marijuana was found shall not create a presumption that such person either knowingly or intentionally possessed such marijuana.

Any person who violates this section is subject to a civil penalty of no more than $25. A violation of this section is a civil offence. Any civil penalties collected pursuant to this section shall be deposited into the Drug Offender Assessment and Treatment Fund established pursuant to § 18.2-251.02. Violations of this section by an adult shall be prepayable according to the procedures in § 16.1-69.40:2.

B. Any violation of this section shall be charged by summons. A summons for a violation of this section may be executed by a law-enforcement officer when such violation is observed by such officer. The summons used by a law-enforcement officer pursuant to this section shall be in form the same as the uniform summons for motor vehicle law violations as prescribed pursuant to § 46.2-388. No court costs shall be assessed for violations of this section. A person's criminal history record information as defined in § 9.1-101 shall not include records of any charges or judgments for a violation of this section, and records of such charges or judgments shall not be reported to the Central Criminal Records

Exchange; however, such violation shall be reported to the Department of Motor Vehicles and shall be included on such individual's driving record.

C. The procedure for appeal and trial of any violation of this section shall be the same as provided by law for misdemeanors; if requested by either party on appeal to the circuit court, trial by jury shall be as provided in Article 4 (§ 19.2-260 et seq.) of Chapter 15 of Title 19.2, and the Commonwealth shall be required to prove its case beyond a reasonable doubt.

D. The provisions of this section shall not apply to members of state, federal, county, city, or town law-enforcement agencies, jail officers, or correctional officers, as defined in § 53.1-1, certified as handlers of dogs trained in the detection of controlled substances when possession of marijuana is necessary for the performance of their duties.

E. The provisions of this section involving marijuana in the form of cannabis products as that term is defined in § ~~54.1-3408.3~~ *4.1-1600* shall not apply to any person who possesses such cannabis product pursuant to a valid written certification issued by a practitioner in the course of his professional practice pursuant to § ~~54.1-3408.3~~ *4.1-1601* for treatment or to alleviate the symptoms of (i) the person's diagnosed condition or disease, (ii) if such person is the parent or guardian of a minor or of a vulnerable adult as defined in § 18.2-369, such minor's or vulnerable adult's diagnosed condition or disease, or (iii) if such person has been designated as a registered agent pursuant to § ~~54.1-3408.3~~ *4.1-1601,* the diagnosed condition or disease of his principal or, if the principal is the parent or legal guardian of a minor or of a vulnerable adult as defined in § 18.2-369, such minor's or vulnerable adult's diagnosed condition or disease.

§ 54.1-2522.1. (Effective until July 1, 2027) Requirements of practitioners.

A. Any prescriber who is licensed in the Commonwealth to treat human patients and is authorized pursuant to §§ 54.1-3303 and 54.1-3408 to issue a prescription for a covered substance shall be registered with the Prescription Monitoring Program by the Department of Health Professions.

B. A prescriber registered with the Prescription Monitoring Program or a person to whom he has delegated authority to access information in the possession of the Prescription Monitoring Program pursuant to § 54.1-2523.2 shall, at the time of initiating a new course of treatment to a human patient that includes the prescribing of opioids anticipated at the onset of treatment to last more than seven consecutive days, request information from the Director for the purpose of determining what, if any, other covered substances are currently prescribed to the patient. In addition, any prescriber who holds a special identification number from the Drug Enforcement Administration authorizing the prescribing of controlled substances approved for use in opioid addiction therapy shall, prior to or as a part of execution of a treatment agreement with the patient, request information from the Director for the purpose of determining what, if any, other covered substances the patient is currently being prescribed. Nothing in this section shall prohibit prescribers from making additional periodic requests for information from the Director as may be required by routine prescribing practices.

C. A prescriber shall not be required to meet the provisions of subsection B if:

1. The opioid is prescribed to a patient currently receiving hospice or palliative care;

2. The opioid is prescribed to a patient during an inpatient hospital admission or at discharge;

3. The opioid is prescribed to a patient in a nursing home or a patient in an assisted living facility that uses a sole source pharmacy;

4. The Prescription Monitoring Program is not operational or available due to temporary technological or electrical failure or natural disaster; or

5. The prescriber is unable to access the Prescription Monitoring Program due to emergency or disaster and documents such circumstances in the patient's medical record.

D. Prior to issuing a written certification for the use of cannabis oil in accordance with § ~~54.1-3408.3~~ *4.1-1601*, a practitioner shall request information from the Director for the purpose of determining what, if any, other covered substances have been dispensed to the patient.

§ 54.1-2522.1. (Effective July 1, 2027) Requirements of practitioners.

A. Any prescriber who is licensed in the Commonwealth to treat human patients and is authorized pursuant to §§ 54.1-3303 and 54.1-3408 to issue a prescription for a covered substance shall be registered with the Prescription Monitoring Program by the Department of Health Professions.

B. Prescribers registered with the Prescription Monitoring Program shall, at the time of initiating a new course of treatment to a human patient that includes the prescribing of benzodiazepine or an opiate anticipated at the onset of treatment to last more than 90 consecutive days, request information from the Director for the purpose of determining what, if any, other covered substances are currently prescribed to the patient. In addition, any prescriber who holds a special identification number from the Drug Enforcement Administration authorizing the prescribing of controlled substances approved for use in opioid addiction therapy shall, prior to or as a part of execution of a treatment agreement with the patient, request information from the Director for the purpose of determining what, if any, other covered substances the patient is currently being prescribed. Nothing in this section shall prohibit prescribers from making additional periodic requests for information from the Director as may be required by routine prescribing practices.

C. The Secretary of Health and Human Resources may identify and publish a list of benzodiazepines

or opiates that have a low potential for abuse by human patients. Prescribers who prescribe such identified benzodiazepines or opiates shall not be required to meet the provisions of subsection B. In addition, a prescriber shall not be required to meet the provisions of subsection B if the course of treatment arises from pain management relating to dialysis or cancer treatments.

D. Prior to issuing a written certification for the use of cannabis oil in accordance with § ~~54.1-3408.3~~ *4.1-1601*, a practitioner shall request information from the Director for the purpose of determining what, if any, other covered substances have been dispensed to the patient.

§ 54.1-2903. What constitutes practice; advertising in connection with medical practice.

A. Any person shall be regarded as practicing the healing arts who actually engages in such practice as defined in this chapter, or who opens an office for such purpose, or who advertises or announces to the public in any manner a readiness to practice or who uses in connection with his name the words or letters "Doctor," "Dr.," "M.D.," "D.O.," "D.P.M.," "D.C.," "Healer," "N.P.," or any other title, word, letter or designation intending to designate or imply that he is a practitioner of the healing arts or that he is able to heal, cure or relieve those suffering from any injury, deformity or disease.

Signing a birth or death certificate, or signing any statement certifying that the person so signing has rendered professional service to the sick or injured, or signing or issuing a prescription for drugs or other remedial agents, shall be prima facie evidence that the person signing or issuing such writing is practicing the healing arts within the meaning of this chapter except where persons other than physicians are required to sign birth certificates.

B. No person regulated under this chapter shall use the title "Doctor" or the abbreviation "Dr." in writing or in advertising in connection with his practice unless he simultaneously uses words, initials, an abbreviation or designation, or other language that identifies the type of practice for which he is licensed. No person regulated under this chapter shall include in any advertisement a reference to marijuana, as defined in § 18.2-247, unless such advertisement is for the treatment of addiction or substance abuse. However, nothing in this subsection shall prevent a person from including in any advertisement that such person is registered with the Board of ~~Pharmacy~~ *Directors of the Virginia Cannabis Control Authority* to issue written certifications for the use of cannabis products, as defined in § ~~54.1-3408.3~~ *4.1-1600*.

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

A. As used in this section:~~"Botanical~~, *"botanical* cannabis," ~~means cannabis that is composed wholly of usable cannabis from the same parts of the same chemovar of cannabis plant~~ *"cannabis oil," "cannabis product," and "practitioner" mean the same as those terms are defined in § 4.1-1600*.

~~"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 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 *in accordance with the provisions of § 4.1-1601*. ~~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.

§ 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 of~~ Chapter ~~34~~ *16 (§ 4.1-1600 et seq.)* of Title ~~54.1 of the Code of Virginia~~ *4.1*;
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 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 of~~ Chapter ~~34~~ *16 (§ 4.1-1600 et seq.)* of Title ~~54.1 of the Code of Virginia~~ *4.1*;
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.

§ 63.2-1803.01. Possession or administration of cannabis oil.

Assisted living facility staff members who are authorized to possess, distribute, or administer medications to residents in accordance with the facility's written plan for medication management shall be permitted to store, dispense, or administer cannabis oil to a resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601* and has registered with the Board of ~~Pharmacy~~ *Directors of the Virginia Cannabis Control Authority*.

2. That Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia is repealed.

3. That the provisions of the first and second enactments of this act shall become effective on January 1, 2024.

4. That the twenty-first enactment of Chapter 550 and the twenty-first enactment of Chapter 551 of the Acts of Assembly of 2021, Special Session I, are repealed.

5. That the Regulations Governing Pharmaceutical Processors (18VAC110-60) as promulgated or amended by the Board of Pharmacy prior to January 1, 2024, shall remain in full force and effect and shall be administered by the Virginia Cannabis Control Authority (the Authority) until the Board of Directors (the Board) of the Authority promulgates regulations to implement the provisions of this act, which shall model, to the greatest extent practicable, the Regulations Governing Pharmaceutical Processors (18VAC110-60) promulgated by the Board of Pharmacy. With the exception of § 2.2-4031 of the Code of Virginia, neither the provisions of the

Administrative Process Act (§ 2.2-4000 et seq. of the Code of Virginia) nor public participation guidelines adopted pursuant thereto shall apply to the Board's initial adoption of regulations to implement the provisions of this act. The Authority shall be vested with all powers and duties held by the Board of Pharmacy prior to January 1, 2024, in its administration of the provisions set forth in § 54.1-3408.3 of the Code of Virginia, as amended by this act, Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia, as repealed by this act, and any regulations promulgated pursuant thereto.

6. That any valid, active permits, certifications, and registrations issued by the Board of Pharmacy pursuant to § 54.1-3408.3 of the Code of Virginia, as amended by this act, Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia, as repealed by this act, or regulations promulgated pursuant thereto prior to January 1, 2024, shall remain valid until their expiration date and be considered to have been issued by the Board of Directors of the Virginia Cannabis Control Authority.

7. That the Virginia Cannabis Control Authority may assess and collect regulatory fees from each pharmaceutical processor and cannabis dispensing facility in an amount sufficient to implement this act.

CHAPTER 780

An Act to amend and reenact §§ 54.1-3408.3, 54.1-3442.5, 54.1-3442.6, and 54.1-3442.7 of the Code of Virginia and to amend the Code of Virginia by adding sections numbered 54.1-3442.7:1, 54.1-3442.7:2, and 54.1-3442.7:3, relating to medical marijuana program; product, registration, dispensing, and recordkeeping requirements; advertising.

[H 1846]

Approved April 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That §§ 54.1-3408.3, 54.1-3442.5, 54.1-3442.6, and 54.1-3442.7 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding sections numbered 54.1-3442.7:1, 54.1-3442.7:2, and 54.1-3442.7:3 as follows:

§ 54.1-3408.3. Certification for use of cannabis products 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 dispensing facility" means the same as that term is defined in § 54.1-3442.5.

"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, *except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.),* 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) *is formulated with cannabis oil or botanical cannabis; (ii) is* produced by a pharmaceutical processor~~,~~ *and sold by a pharmaceutical processor or cannabis dispensing facility; (iii) is* registered with the Board~~,~~*; (iv) contains, except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.), no more than 10 milligrams of tetrahydrocannabinol per dose;* and *(v) is* 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.

"Pharmaceutical processor" means the same as that term is defined in § 54.1-3442.5.

"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. *No practitioner may issue a written certification while such practitioner is on the premises of a pharmaceutical processor or cannabis dispensing facility. A pharmaceutical processor shall not endorse or promote any practitioner who issues certifications to patients.* 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. *A*

practitioner who issues written certifications shall not directly or indirectly accept, solicit, or receive anything of value from a pharmaceutical processor, cannabis dispensing facility, or any person associated with a pharmaceutical processor, cannabis dispensing facility, or provider of paraphernalia, excluding information on products or educational materials on the benefits and risks of cannabis products.

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~~ *a practitioner's professional licensing board* from sanctioning ~~a~~ *the* 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~~ *(i)* shall hold sufficient education and training to exercise appropriate professional judgment in the certification of patients*; (ii) shall not offer a discount or any other thing of value to a patient or a patient's parent, guardian, or registered agent that is contingent on or encourages the person's decision to use a particular pharmaceutical processor or cannabis product; (iii) shall not issue a certification to himself or his family members, employees, or coworkers; (iv) shall not provide product samples containing cannabis other than those approved by the U.S. Food and Drug Administration; and (v) shall not accept compensation from a pharmaceutical processor or cannabis dispensing facility*. 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 *patient certification or agent* 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.

§ 54.1-3442.5. Definitions.

As used in this article:

"Botanical cannabis," "cannabis oil," "cannabis product," *"designated caregiver facility," "practitioner," "registered agent,"* and "usable cannabis" have the same meanings as specified in § 54.1-3408.3.

"Cannabis dispensing facility" means a facility that (i) has obtained a permit from the Board pursuant to § 54.1-3442.6; (ii) is owned, at least in part, by a pharmaceutical processor; and (iii) dispenses

cannabis products produced by a pharmaceutical processor 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.

~~"Designated caregiver facility" has the same meaning as defined in § 54.1-3408.3.~~

"Pharmaceutical processor" means a facility that (i) has obtained a permit from the Board pursuant to § ~~54.1-3408.3~~ *54.1-3442.6* and (ii) cultivates Cannabis plants intended only for the production of cannabis oil, botanical cannabis, and usable cannabis, produces cannabis products, and dispenses cannabis products to a patient pursuant to a written certification, 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.

~~"Practitioner" has the same meaning as specified in § 54.1-3408.3.~~

~~"Registered agent" has the same meaning as specified in § 54.1-3408.3.~~

§ 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~~ *products* that provide that each dispensed dose of *a* cannabis ~~oil~~ *product* not exceed 10 milligrams of ~~delta-9-tetrahydrocannabinol~~ *total tetrahydrocannabinol, except as permitted under § 54.1-3442.7:2*; (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. *Remediated botanical cannabis or cannabis oil that passes such quality testing may be packaged and labeled.* 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~~ *15* percent deviation basis, of ~~active~~ ~~ingredients~~ *total THC and total CBD. No cannabis product shall have an expiration date longer than six months from the date of the cannabis product registration approval unless supported by stability testing.*

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 *unless all cannabis products are contained in a vault or other similar container to which only the pharmacist has access controls*. 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~~ *and cannabis* 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~~ *total tetrahydrocannabinol* in any cannabis product on site may be up to ~~10~~ *15* percent greater than or less than the level of ~~delta-9-tetrahydrocannabinol measured for labeling~~ *total tetrahydrocannabinol listed in the approved cannabis product registration*. 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 *that have an expiration date longer than six months*.

§ 54.1-3442.7:1. Packaging and labeling; corrections; records.

A. Pharmaceutical processors shall comply with all packaging and labeling requirements set forth in this article and Board regulations.

B. No cannabis product shall be 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 cannabis product.

C. Pharmaceutical processors may correct typographical errors made on cannabis product labels and any documents generated as the result of a wholesale transaction.

§ 54.1-3442.7:2. Cannabis product registration; approval, deviation, and modification.

A. A pharmaceutical processor shall register with the Board each cannabis product it manufactures. Applications for cannabis product registration shall be submitted to the Board on a form prescribed by the Board.

B. An application for cannabis product registration shall include:

1. The total tetrahydrocannabinol and total cannabidiol in such cannabis product, based on laboratory testing results for the cannabis product formulation;

2. A product name;

3. A proposed product package; and

4. A proposed product label, which shall not be required to contain an expiration date at the time of application.

C. The Board shall register all cannabis products that meet testing, labeling, and packaging standards after an application for registration is submitted. If the cannabis product fails to meet such standards or the application was deficient, the Board shall notify the applicant of the specific reasons for such failure or deficiency.

D. Within two business days of the Board's approval or deemed approval, the Board shall enter the cannabis product's national drug code number into the Prescription Monitoring Program.

E. The following cannabis product deviations from an approved cannabis product registration shall be permitted without any requirement for a new cannabis product registration or notice to the Board:

1. A deviation in the concentration of total tetrahydrocannabinol (THC) or total cannabidiol (CBD) in a cannabis product or dose thereof of up to 15 percent greater than or less than the concentration of total tetrahydrocannabinol or total cannabidiol, either or both, listed in the approved cannabis product registration; however, for a cannabis product with five milligrams or less of total THC or total CBD per dose, the total THC or total CBD concentration shall be within 0.5 milligrams of the single dose total THC or total CBD concentrations approved for that cannabis product;

2. A variation in packaging, provided that the packaging is substantially similar to the approved packaging and otherwise complies with applicable packaging requirements;

3. A deviation in labeling, including a variation made in accordance with § 54.1-3442.7:1, that reflects allowable deviations in total THC or total CBD or that makes a minor text, font, design, or similar modification, provided that the labeling is substantially similar to the approved labeling and otherwise complies with applicable labeling requirements; and

4. Any other insignificant changes.

F. A pharmaceutical processor may submit a request to modify an existing cannabis product registration in the event of a cannabis product deviation that is not set forth in subsection E. Upon receipt, the Board shall respond to such request. The Board may grant or deny the request, propose a reasonable revision, or require the pharmaceutical processor to provide additional information.

§ 54.1-3442.7:3. Advertising and marketing.

A. Pharmaceutical processors and cannabis dispensing facilities may (i) advertise and promote products and operations and (ii) provide educational material to practitioners, patients, and the public.

B. Pharmaceutical processors and cannabis dispensing facilities may engage in advertising or marketing that does not:

1. Include false or misleading statements;

2. Promote overconsumption;

3. Depict a person younger than 21 years of age;

4. Appeal particularly to persons younger than 21 years of age, including by using cartoons in any way;

5. Associate cannabis products with candy or similar products or depicts any images that bear a reasonable resemblance to a candy or similar product; or

6. Contain any seal, flag, crest, coat of arms, or other insignia that is likely to mislead patients or the public to believe that the cannabis product is made or endorsed by the Commonwealth.

C. All advertising and marketing by pharmaceutical processors and cannabis dispensing facilities shall (i) accurately and legibly identify the pharmaceutical processor or cannabis dispensing facility responsible for its content, (ii) include a statement that cannabis products are for use by certified patients only, and (iii) comply with Board regulations.

2. That pharmaceutical processors and cannabis dispensing facilities shall collect and provide to the Board of Pharmacy by July 1, 2024, data regarding the impact of this act on program participation, reductions in the price of cannabis products, and improved operational efficiencies.

3. That the Board of Pharmacy shall amend its regulations, including 18VAC110-60-270, 18VAC110-60-285, 18VAC110-60-290, and 18VAC110-60-310, to replace any references to "brand" with "registered cannabis product name."

4. That the Board of Pharmacy may assess and collect regulatory fees from each pharmaceutical processor and cannabis dispensing facility in an amount sufficient to implement the provisions of this act.

VIRGINIA ACTS OF ASSEMBLY -- 2023 RECONVENED SESSION

CHAPTER 799

An Act to amend and reenact §§ 54.1-2521, 54.1-3408.3, and 54.1-3442.6 of the Code of Virginia, relating to medical marijuana program; product requirements; certifications; reporting.

[H 2368]

Approved April 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That §§ 54.1-2521, 54.1-3408.3, and 54.1-3442.6 of the Code of Virginia are amended and reenacted as follows:

§ 54.1-2521. Reporting requirements.

A. The failure by any person subject to the reporting requirements set forth in this section and the Department's regulations to report the dispensing of covered substances shall constitute grounds for disciplinary action by the relevant health regulatory board.

B. Upon dispensing a covered substance, a dispenser of such covered substance shall report the following information:

1. The recipient's name and address.

2. The recipient's date of birth.

3. The covered substance that was dispensed to the recipient, *which, in the case of a cannabis product, shall be listed as the primary cannabinoid of such cannabis product.*

4. The quantity of the covered substance that was dispensed.

5. The date of the dispensing.

6. The prescriber's identifier number and, in cases in which the covered substance is a cannabis product, the *product's national drug code and the* expiration date of the written certification.

7. The dispenser's identifier number.

8. The method of payment for the prescription.

9. Any other non-clinical information that is designated by the Director as necessary for the implementation of this chapter in accordance with the Department's regulations.

10. Any other information specified in regulations promulgated by the Director as required in order for the Prescription Monitoring Program to be eligible to receive federal funds.

C. Except as provided in subdivision 7 of § 54.1-2522, in cases where the ultimate user of a covered substance is an animal, the dispenser shall report the relevant information required by subsection B for the owner of the animal.

D. The reports required herein shall be made to the Department or its agent within 24 hours or the dispenser's next business day, whichever comes later, and shall be made and transmitted in such manner and format and according to the standards and schedule established in the Department's regulations.

§ 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 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 *(i)* 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 *(ii)* registered with the Board *or listed on the patient's written certification* 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 *unless the individual's name is listed on the patient's written certification. An individual may, on the basis of medical need and in the discretion of the patient's registered practitioner, be listed on the patient's written certification upon the patient's request.* 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; (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. *Product labels for all cannabis products and botanical cannabis shall be complete, accurate, easily discernable, and uniform among different products and brands. Pharmaceutical processors shall affix to all cannabis products and botanical cannabis a label, which shall also be accessible on the pharmaceutical processor's website, that includes:*

1. The product name;

2. All active and inactive ingredients, including cannabinoids, terpenes, additives, preservatives, flavorings, sweeteners, and carrier oils;

3. The total percentage and milligrams of tetrahydrocannabinol and cannabidiol included in the product and the number of milligrams of tetrahydrocannabinol and cannabidiol in each serving;

4. The amount of product that constitutes a single serving and the amount recommended for use by the practitioner or dispensing pharmacist;

5. Information regarding the product's purpose and detailed usage directions;

6. Child and safety warnings in a conspicuous font; and

7. Such other information required by the Board.

O. A pharmaceutical processor or cannabis dispensing facility shall maintain an adequate supply of cannabis products that (i) contain cannabidiol as their primary cannabinoid and (ii) have low levels of or no tetrahydrocannabinol.

P. 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.~~ *Q.* The Board shall register all cannabis products that meet testing, labeling, and packaging standards.

CHAPTER 740

An Act to amend and reenact §§ 4.1-604, 4.1-605, 4.1-627, 18.2-251.1:1, 18.2-251.1:2, 22.1-277, 32.1-127, 32.1-162.6:1, 40.1-27.4, 46.2-341.20:7, 54.1-2522.1, as it is currently effective and as it shall become effective, 54.1-2903, 54.1-3408.3, 59.1-200, and 63.2-1803.01 of the Code of Virginia; to amend the Code of Virginia by adding in Title 4.1 a chapter numbered 16, consisting of sections numbered 4.1-1600 through 4.1-1605; and to repeal Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia and the twenty-first enactment of Chapter 550 and the twenty-first enactment of Chapter 551 of the Acts of Assembly of 2021, Special Session I, relating to medical cannabis program; transition from Board of Pharmacy to Virginia Cannabis Control Authority.

[S 788]

Approved April 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That §§ 4.1-604, 4.1-605, 4.1-627, 18.2-251.1:1, 18.2-251.1:2, 22.1-277, 32.1-127, 32.1-162.6:1, 40.1-27.4, 46.2-341.20:7, 54.1-2522.1, as it is currently effective and as it shall become effective, 54.1-2903, 54.1-3408.3, 59.1-200, and 63.2-1803.01 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding in Title 4.1 a chapter numbered 16, consisting of sections numbered 4.1-1600 through 4.1-1605, as follows:

§ 4.1-604. Powers and duties of the Board.

The Board shall have the following powers and duties:

1. Promulgate regulations in accordance with the Administrative Process Act (§ 2.2-4000 et seq.) and § 4.1-606;

2. Control the possession, sale, transportation, and delivery of marijuana and marijuana products;

3. Grant, suspend, ~~and~~ *restrict,* revoke ~~licenses for the cultivation, manufacture, distribution, sale, and testing of marijuana and marijuana products as provided by law~~, *or refuse to grant or renew any license or permit issued or authorized pursuant to this subtitle*;

4. Determine the nature, form, and capacity of all containers used for holding marijuana products to be kept or sold and prescribe the form and content of all labels and seals to be placed thereon;

5. Maintain actions to enjoin common nuisances as defined in § 4.1-1113;

6. Establish standards and implement an online course for employees of retail marijuana stores that trains employees on how to educate consumers on the potential risks of marijuana use;

7. Establish a plan to develop and disseminate to retail marijuana store licensees a pamphlet or similar document regarding the potential risks of marijuana use to be prominently displayed and made available to consumers;

8. Establish a position for a Cannabis Social Equity Liaison who shall lead the Cannabis Business Equity and Diversity Support Team and liaise with the Director of Diversity, Equity, and Inclusion on matters related to diversity, equity, and inclusion standards in the marijuana industry;

9. Establish a Cannabis Business Equity and Diversity Support Team, which shall (i) develop requirements for the creation and submission of diversity, equity, and inclusion plans by persons who wish to possess a license in more than one license category pursuant to subsection C of § 4.1-805, which may include a requirement that the licensee participate in social equity apprenticeship plan, and an approval process and requirements for implementation of such plans; (ii) be responsible for conducting an analysis of potential barriers to entry for small, women-owned, and minority-owned businesses and veteran-owned businesses interested in participating in the marijuana industry and recommending strategies to effectively mitigate such potential barriers; (iii) provide assistance with business planning for potential marijuana establishment licensees; (iv) spread awareness of business opportunities related to the marijuana marketplace in areas disproportionately impacted by marijuana prohibition and enforcement; (v) provide technical assistance in navigating the administrative process to potential marijuana establishment licensees; and (vi) conduct other outreach initiatives in areas disproportionately impacted by marijuana prohibition and enforcement as necessary;

10. Establish a position for an individual with professional experience in a health related field who shall staff the Cannabis Public Health Advisory Council, established pursuant to § 4.1-603, liaise with the Office of the Secretary of Health and Human Resources and relevant health and human services agencies and organizations, and perform other duties as needed;

11. Establish and implement a plan, in coordination with the Cannabis Social Equity Liaison and the Director of Diversity, Equity, and Inclusion to promote and encourage participation in the marijuana industry by people from communities that have been disproportionately impacted by marijuana prohibition and enforcement and to positively impact those communities;

12. Sue and be sued, implead and be impleaded, and complain and defend in all courts;

13. Adopt, use, and alter at will a common seal;

14. Fix, alter, charge, and collect rates, rentals, fees, and other charges for the use of property of, the sale of products of, or services rendered by the Authority at rates to be determined by the Authority for the purpose of providing for the payment of the expenses of the Authority;

15. Make and enter into all contracts and agreements necessary or incidental to the performance of its duties, the furtherance of its purposes, and the execution of its powers under this subtitle, including agreements with any person or federal agency;

16. Employ, at its discretion, consultants, researchers, architects, engineers, accountants, financial experts, investment bankers, superintendents, managers, and such other employees and special agents as may be necessary and fix their compensation to be payable from funds made available to the Authority. Legal services for the Authority shall be provided by the Attorney General in accordance with Chapter 5 (§ 2.2-500 et seq.) of Title 2.2;

17. Receive and accept from any federal or private agency, foundation, corporation, association, or person grants or other aid to be expended in accomplishing the objectives of the Authority, and receive and accept from the Commonwealth or any state and any municipality, county, or other political subdivision thereof or from any other source aid or contributions of either money, property, or other things of value, to be held, used, and applied only for the purposes for which such grants and contributions may be made. All federal moneys accepted under this section shall be accepted and expended by the Authority upon such terms and conditions as are prescribed by the United States and as are consistent with state law, and all state moneys accepted under this section shall be expended by the Authority upon such terms and conditions as are prescribed by the Commonwealth;

18. Adopt, alter, and repeal bylaws, rules, and regulations governing the manner in which its business shall be transacted and the manner in which the powers of the Authority shall be exercised and its duties performed. The Board may delegate or assign any duty or task to be performed by the Authority to any officer or employee of the Authority. The Board shall remain responsible for the performance of any such duties or tasks. Any delegation pursuant to this subdivision shall, where appropriate, be accompanied by written guidelines for the exercise of the duties or tasks delegated. Where appropriate, the guidelines shall require that the Board receive summaries of actions taken. Such delegation or assignment shall not relieve the Board of the responsibility to ensure faithful performance of the duties and tasks;

19. Conduct or engage in any lawful business, activity, effort, or project consistent with the Authority's purposes or necessary or convenient to exercise its powers;

20. Develop policies and procedures generally applicable to the procurement of goods, services, and construction, based upon competitive principles;

21. Develop policies and procedures consistent with Article 4 (§ 2.2-4347 et seq.) of Chapter 43 of Title 2.2;

22. Acquire, purchase, hold, use, lease, or otherwise dispose of any property, real, personal or mixed, tangible or intangible, or any interest therein necessary or desirable for carrying out the purposes of the Authority; lease as lessee any property, real, personal or mixed, tangible or intangible, or any interest therein, at such annual rental and on such terms and conditions as may be determined by the Board; lease as lessor to any person any property, real, personal or mixed, tangible or intangible, or any interest therein, at any time acquired by the Authority, whether wholly or partially completed, at such annual rental and on such terms and conditions as may be determined by the Board; sell, transfer, or convey any property, real, personal or mixed, tangible or intangible, or any interest therein, at any time acquired or held by the Authority on such terms and conditions as may be determined by the Board; and occupy and improve any land or building required for the purposes of this subtitle;

23. Purchase, lease, or acquire the use of, by any manner, any plant or equipment that may be considered necessary or useful in carrying into effect the purposes of this subtitle, including rectifying, blending, and processing plants;

24. Appoint every agent and employee required for its operations, require any or all of them to give bonds payable to the Commonwealth in such penalty as shall be fixed by the Board, and engage the services of experts and professionals;

25. Hold and conduct hearings, issue subpoenas requiring the attendance of witnesses and the production of records, memoranda, papers, and other documents before the Board or any agent of the Board, and administer oaths and take testimony thereunder. The Board may authorize any Board member or agent of the Board to hold and conduct hearings, issue subpoenas, administer oaths and take testimony thereunder, and decide cases, subject to final decision by the Board, on application of any party aggrieved. The Board may enter into consent agreements and may request and accept from any applicant ~~or~~, licensee, *or permittee* a consent agreement in lieu of proceedings on (i) objections to the issuance of a license *or permit* or (ii) disciplinary action. Any such consent agreement *(a)* shall include findings of fact *and provisions regarding whether the terms of the consent agreement are confidential* and *(b)* may include an admission or a finding of a violation. A consent agreement shall not be considered a case decision of the Board and shall not be subject to judicial review under the provisions

of the Administrative Process Act (§ 2.2-4000 et seq.), but may be considered by the Board in future disciplinary proceedings;

26. Make a reasonable charge for preparing and furnishing statistical information and compilations to persons other than (i) officials, including court and police officials, of the Commonwealth and of its subdivisions if the information requested is for official use and (ii) persons who have a personal or legal interest in obtaining the information requested if such information is not to be used for commercial or trade purposes;

27. ~~Assess~~ *Take appropriate disciplinary action and assess* and collect civil penalties and civil charges for violations of this subtitle and Board regulations;

28. Review and approve any proposed legislative or regulatory changes suggested by the Chief Executive Officer as the Board deems appropriate;

29. Report quarterly to the Secretary of Public Safety and Homeland Security on the law-enforcement activities undertaken to enforce the provisions of this subtitle;

30. Establish and collect fees for all permits set forth in this subtitle, including fees associated with applications for such permits;

31. Develop and make available on its website guidance documents regarding compliance and safe practices for persons who cultivate marijuana at home for personal use, which shall include information regarding cultivation practices that promote personal and public safety, including child protection, and discourage practices that create a nuisance;

32. Develop and make available on its website a resource that provides information regarding (i) responsible marijuana consumption; (ii) health risks and other dangers associated with marijuana consumption, including inability to operate a motor vehicle and other types of transportation and equipment; and (iii) ancillary effects of marijuana consumption, including ineligibility for certain employment opportunities. The Board shall require that the web address for such resource be included on the label of all retail marijuana and retail marijuana product as provided in § 4.1-1402; and

33. Do all acts necessary or advisable to carry out the purposes of this subtitle.

§ 4.1-605. Additional powers; mediation; alternative dispute resolution; confidentiality.

A. As used in this section:

"Appropriate case" means any alleged license *or permit* violation or objection to the application for a license *or permit* in which it is apparent that there are significant issues of disagreement among interested persons and for which the Board finds that the use of a mediation or dispute resolution proceeding is in the public interest.

"Dispute resolution proceeding" means the same as that term is defined in § 8.01-576.4.

"Mediation" means the same as that term is defined in § 8.01-576.4.

"Neutral" means the same as that term is defined in § 8.01-576.4.

B. The Board may use mediation or a dispute resolution proceeding in appropriate cases to resolve underlying issues or reach a consensus or compromise on contested issues. Mediation and other dispute resolution proceedings as authorized by this section shall be voluntary procedures that supplement, rather than limit, other dispute resolution techniques available to the Board. Mediation or a dispute resolution proceeding may be used for an objection to the issuance of a license *or permit* only with the consent of, and participation by, the applicant for ~~licensure~~ *a license or permit* and shall be terminated at the request of such applicant.

C. Any resolution of a contested issue accepted by the Board under this section shall be considered a consent agreement as provided in § 4.1-604. The decision to use mediation or a dispute resolution proceeding is in the Board's sole discretion and shall not be subject to judicial review.

D. The Board may adopt rules and regulations, in accordance with the Administrative Process Act (§ 2.2-4000 et seq.), for the implementation of this section. Such rules and regulations may include (i) standards and procedures for the conduct of mediation and dispute resolution proceedings, including an opportunity for interested persons identified by the Board to participate in the proceeding; (ii) the appointment and function of a neutral to encourage and assist parties to voluntarily compromise or settle contested issues; and (iii) procedures to protect the confidentiality of papers, work products, or other materials.

E. The provisions of § 8.01-576.10 concerning the confidentiality of a mediation or dispute resolution proceeding shall govern all such proceedings held pursuant to this section except where the Board uses or relies on information obtained in the course of such proceeding in granting ~~a~~ ~~license~~, suspending, *restricting,* or revoking a license *or permit*, or *in* accepting payment of a civil penalty or investigative costs. ~~However,~~ ~~a~~ ~~consent~~ ~~agreement~~ *Consent agreements shall be* signed by ~~the~~ *all* parties *and* shall ~~not be~~ *include provisions regarding whether the terms of the consent agreement are* confidential.

§ 4.1-627. Hearings; representation by counsel.

Any licensee, *permittee,* or applicant for ~~any~~ *a* license ~~granted by the Board~~ *or permit authorized by this subtitle* shall have the right to be represented by counsel at any Board hearing for which he has received notice. The licensee, *permittee,* or applicant shall not be required to be represented by counsel during such hearing. Any officer or director of a corporation may examine, cross-examine, and question witnesses, present evidence on behalf of the corporation, and draw conclusions and make arguments

before the Board or hearing officers without being in violation of the provisions of § 54.1-3904.

CHAPTER 16.

MEDICAL CANNABIS PROGRAM.

§ 4.1-1600. Definitions.

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

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

"Cannabis dispensing facility" means a facility that (i) has obtained a permit from the Board pursuant to § 4.1-1602; (ii) is owned, at least in part, by a pharmaceutical processor; and (iii) dispenses cannabis products produced by a pharmaceutical processor 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.

"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 § 4.1-1602, or a dilution of the resin of the Cannabis plant that contains no more than 10 milligrams of delta-9-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.

"Dispense" means the same as that term is defined in § 54.1-3300.

"Pharmaceutical processor" means a facility that (i) has obtained a permit from the Board pursuant to § 4.1-1602 and (ii) cultivates Cannabis plants intended only for the production of cannabis oil, botanical cannabis, and usable cannabis, produces cannabis products, and dispenses cannabis products to a patient pursuant to a written certification, 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.

"Pharmacist" means the same as that term is defined in § 54.1-3300.

"Pharmacy intern" means the same as that term is defined in § 54.1-3300.

"Pharmacy technician" means the same as that term is defined in § 54.1-3300.

"Pharmacy technician trainee" means the same as that term is defined in § 54.1-3300.

"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 Boards of Nursing and Medicine.

"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 F of § 4.1-1601.

"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, derivative, mixture, or preparation of the mature stalks; or (iii) oil or cake made from the seeds of the plant.

§ 4.1-1601. Certification for use of cannabis for treatment.

A. 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 audiovisual 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.

B. The written certification shall be on a form provided by the Authority. 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 A 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.

C. 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 A. 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.

D. 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.

E. 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 Authority.

F. 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.

G. 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.

H. 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 patient's registered agent, but only with respect to information related to such patient.

§ 4.1-1602. 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 Authority 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; (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 processor 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 N; (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 products.

D. The Board shall require pharmaceutical processors, after processing and before dispensing any cannabis products, to make a sample available from each batch of cannabis product for testing by an independent laboratory that is located in Commonwealth and meets 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 of Pharmacy 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.

H. 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.

I. 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.

J. 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.

K. 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.

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

M. 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.

N. 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.

O. 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 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 shall consider and keep on file all public comments received for any regulation adopted pursuant to this section.

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

§ 4.1-1603. 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 § 4.1-1602. 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 on General Laws and the Senate Committee on Rehabilitation and Social Services on the operation of pharmaceutical processors and cannabis dispensing facilities issued a permit by the Board.

D. The concentration of delta-9-tetrahydrocannabinol in any cannabis product on site may be up to 10 percent greater than or less than the level of delta-9-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.

§ 4.1-1604. Criminal liability; exceptions.

No agent or employee of a pharmaceutical processor or cannabis dispensing facility shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) or § 18.2-248, 18.2-248.1, or 18.2-250 for possession or manufacture of marijuana or for possession, manufacture, or distribution of cannabis products, subject to any civil penalty, denied any right or privilege, or subject to any disciplinary action by a professional licensing board if such agent or employee (i) possessed or manufactured such marijuana for the purposes of producing cannabis products in accordance with the provisions of this chapter and Board regulations or (ii) possessed, manufactured, or distributed such cannabis products that are consistent with generally accepted cannabis industry standards in accordance with the provisions of this chapter and Board regulations.

§ 4.1-1605. Summary suspensions and restrictions.

A. The Board may summarily suspend or restrict a permit issued pursuant to § 4.1-1602 without a hearing if the Board finds that such suspension or restriction is necessary to prevent substantial danger to public health or safety. The Board shall make decisions to summarily suspend or restrict a permit only during an in-person meeting in which a quorum is present; however, if, after a good faith effort, the Board is unable to assemble a quorum and a majority of the Board members determine that continued operation by the permittee constitutes a substantial danger to public health or safety, the Board may summarily suspend the permit during a telephone, video, or other electronic conference. Institution of proceedings for a hearing shall be provided simultaneously with a summary suspension. The Board may summarily restrict a permit without proceeding simultaneously with notification of an informal conference pursuant to § 2.2-4019 or Board regulations. Such hearing or conference shall be held within a reasonable amount of time after the summary suspension or restriction is issued.

B. Allegations of violations of this subtitle shall be submitted to the Board in writing.

§ 18.2-251.1:1. Possession or distribution of cannabis oil; public schools.

No school nurse employed by a local school board, person employed by a local health department who is assigned to the public school pursuant to an agreement between the local health department and the school board, or other person employed by or contracted with a local school board to deliver health-related services shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-248, 18.2-248.1, 18.2-250, or 18.2-255 for the possession or distribution of cannabis oil for storing, dispensing, or administering cannabis oil, in accordance with a policy adopted by the local school board, to a student who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601*.

§ 18.2-251.1:2. Possession or distribution of cannabis oil; nursing homes and certified nursing facilities; hospice and hospice facilities; assisted living facilities.

No person employed by a nursing home, hospice, hospice facility, or assisted living facility and authorized to possess, distribute, or administer medications to patients or residents shall be prosecuted under Chapter 11 (§ 4.1-1100 et seq.) of Title 4.1 or § 18.2-248, 18.2-248.1, or 18.2-250 for the possession or distribution of cannabis oil for the purposes of storing, dispensing, or administering cannabis oil to a patient or resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*.

§ 22.1-277. Suspensions and expulsions of students generally.

A. Students may be suspended or expelled from attendance at school for sufficient cause; however, in no cases may sufficient cause for suspensions include only instances of truancy.

B. Except as provided in subsection C or § 22.1-277.07 or 22.1-277.08, no student in preschool through grade three shall be suspended for more than three school days or expelled from attendance at school, unless (i) the offense involves physical harm or credible threat of physical harm to others or (ii)

the local school board or the division superintendent or his designee finds that aggravating circumstances exist, as defined by the Department.

C. Any student for whom the division superintendent of the school division in which such student is enrolled has received a report pursuant to § 16.1-305.1 of an adjudication of delinquency or a conviction for an offense listed in subsection G of § 16.1-260 may be suspended or expelled from school attendance pursuant to this article.

D. The authority provided in § 22.1-276.2 for teachers to remove students from their classes in certain instances of disruptive behavior shall not be interpreted to affect the operation of § 22.1-277.04, 22.1-277.05, or 22.1-277.06.

E. Notwithstanding the provisions of § 22.1-277.08, no school board shall be required to suspend or expel any student who holds a valid written certification for the use of cannabis oil issued by a practitioner in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601* for the possession or use of such oil in accordance with the student's individualized health plan and in compliance with a policy adopted by the school board.

§ 32.1-127. Regulations.

A. The regulations promulgated by the Board to carry out the provisions of this article shall be in substantial conformity to the standards of health, hygiene, sanitation, construction and safety as established and recognized by medical and health care professionals and by specialists in matters of public health and safety, including health and safety standards established under provisions of Title XVIII and Title XIX of the Social Security Act, and to the provisions of Article 2 (§ 32.1-138 et seq.).

B. Such regulations:

1. Shall include minimum standards for (i) the construction and maintenance of hospitals, nursing homes and certified nursing facilities to ensure the environmental protection and the life safety of its patients, employees, and the public; (ii) the operation, staffing and equipping of hospitals, nursing homes and certified nursing facilities; (iii) qualifications and training of staff of hospitals, nursing homes and certified nursing facilities, except those professionals licensed or certified by the Department of Health Professions; (iv) conditions under which a hospital or nursing home may provide medical and nursing services to patients in their places of residence; and (v) policies related to infection prevention, disaster preparedness, and facility security of hospitals, nursing homes, and certified nursing facilities;

2. Shall provide that at least one physician who is licensed to practice medicine in this Commonwealth shall be on call at all times, though not necessarily physically present on the premises, at each hospital which operates or holds itself out as operating an emergency service;

3. May classify hospitals and nursing homes by type of specialty or service and may provide for licensing hospitals and nursing homes by bed capacity and by type of specialty or service;

4. Shall also require that each hospital establish a protocol for organ donation, in compliance with federal law and the regulations of the Centers for Medicare and Medicaid Services (CMS), particularly 42 C.F.R. § 482.45. Each hospital shall have an agreement with an organ procurement organization designated in CMS regulations for routine contact, whereby the provider's designated organ procurement organization certified by CMS (i) is notified in a timely manner of all deaths or imminent deaths of patients in the hospital and (ii) is authorized to determine the suitability of the decedent or patient for organ donation and, in the absence of a similar arrangement with any eye bank or tissue bank in Virginia certified by the Eye Bank Association of America or the American Association of Tissue Banks, the suitability for tissue and eye donation. The hospital shall also have an agreement with at least one tissue bank and at least one eye bank to cooperate in the retrieval, processing, preservation, storage, and distribution of tissues and eyes to ensure that all usable tissues and eyes are obtained from potential donors and to avoid interference with organ procurement. The protocol shall ensure that the hospital collaborates with the designated organ procurement organization to inform the family of each potential donor of the option to donate organs, tissues, or eyes or to decline to donate. The individual making contact with the family shall have completed a course in the methodology for approaching potential donor families and requesting organ or tissue donation that (a) is offered or approved by the organ procurement organization and designed in conjunction with the tissue and eye bank community and (b) encourages discretion and sensitivity according to the specific circumstances, views, and beliefs of the relevant family. In addition, the hospital shall work cooperatively with the designated organ procurement organization in educating the staff responsible for contacting the organ procurement organization's personnel on donation issues, the proper review of death records to improve identification of potential donors, and the proper procedures for maintaining potential donors while necessary testing and placement of potential donated organs, tissues, and eyes takes place. This process shall be followed, without exception, unless the family of the relevant decedent or patient has expressed opposition to organ donation, the chief administrative officer of the hospital or his designee knows of such opposition, and no donor card or other relevant document, such as an advance directive, can be found;

5. Shall require that each hospital that provides obstetrical services establish a protocol for admission or transfer of any pregnant woman who presents herself while in labor;

6. Shall also require that each licensed hospital develop and implement a protocol requiring written discharge plans for identified, substance-abusing, postpartum women and their infants. The protocol shall

require that the discharge plan be discussed with the patient and that appropriate referrals for the mother and the infant be made and documented. Appropriate referrals may include, but need not be limited to, treatment services, comprehensive early intervention services for infants and toddlers with disabilities and their families pursuant to Part H of the Individuals with Disabilities Education Act, 20 U.S.C. § 1471 et seq., and family-oriented prevention services. The discharge planning process shall involve, to the extent possible, the other parent of the infant and any members of the patient's extended family who may participate in the follow-up care for the mother and the infant. Immediately upon identification, pursuant to § 54.1-2403.1, of any substance-abusing, postpartum woman, the hospital shall notify, subject to federal law restrictions, the community services board of the jurisdiction in which the woman resides to appoint a discharge plan manager. The community services board shall implement and manage the discharge plan;

7. Shall require that each nursing home and certified nursing facility fully disclose to the applicant for admission the home's or facility's admissions policies, including any preferences given;

8. Shall require that each licensed hospital establish a protocol relating to the rights and responsibilities of patients which shall include a process reasonably designed to inform patients of such rights and responsibilities. Such rights and responsibilities of patients, a copy of which shall be given to patients on admission, shall be consistent with applicable federal law and regulations of the Centers for Medicare and Medicaid Services;

9. Shall establish standards and maintain a process for designation of levels or categories of care in neonatal services according to an applicable national or state-developed evaluation system. Such standards may be differentiated for various levels or categories of care and may include, but need not be limited to, requirements for staffing credentials, staff/patient ratios, equipment, and medical protocols;

10. Shall require that each nursing home and certified nursing facility train all employees who are mandated to report adult abuse, neglect, or exploitation pursuant to § 63.2-1606 on such reporting procedures and the consequences for failing to make a required report;

11. Shall permit hospital personnel, as designated in medical staff bylaws, rules and regulations, or hospital policies and procedures, to accept emergency telephone and other verbal orders for medication or treatment for hospital patients from physicians, and other persons lawfully authorized by state statute to give patient orders, subject to a requirement that such verbal order be signed, within a reasonable period of time not to exceed 72 hours as specified in the hospital's medical staff bylaws, rules and regulations or hospital policies and procedures, by the person giving the order, or, when such person is not available within the period of time specified, co-signed by another physician or other person authorized to give the order;

12. Shall require, unless the vaccination is medically contraindicated or the resident declines the offer of the vaccination, that each certified nursing facility and nursing home provide or arrange for the administration to its residents of (i) an annual vaccination against influenza and (ii) a pneumococcal vaccination, in accordance with the most recent recommendations of the Advisory Committee on Immunization Practices of the Centers for Disease Control and Prevention;

13. Shall require that each nursing home and certified nursing facility register with the Department of State Police to receive notice of the registration, reregistration, or verification of registration information of any person required to register with the Sex Offender and Crimes Against Minors Registry pursuant to Chapter 9 (§ 9.1-900 et seq.) of Title 9.1 within the same or a contiguous zip code area in which the home or facility is located, pursuant to § 9.1-914;

14. Shall require that each nursing home and certified nursing facility ascertain, prior to admission, whether a potential patient is required to register with the Sex Offender and Crimes Against Minors Registry pursuant to Chapter 9 (§ 9.1-900 et seq.) of Title 9.1, if the home or facility anticipates the potential patient will have a length of stay greater than three days or in fact stays longer than three days;

15. Shall require that each licensed hospital include in its visitation policy a provision allowing each adult patient to receive visits from any individual from whom the patient desires to receive visits, subject to other restrictions contained in the visitation policy including, but not limited to, those related to the patient's medical condition and the number of visitors permitted in the patient's room simultaneously;

16. Shall require that each nursing home and certified nursing facility shall, upon the request of the facility's family council, send notices and information about the family council mutually developed by the family council and the administration of the nursing home or certified nursing facility, and provided to the facility for such purpose, to the listed responsible party or a contact person of the resident's choice up to six times per year. Such notices may be included together with a monthly billing statement or other regular communication. Notices and information shall also be posted in a designated location within the nursing home or certified nursing facility. No family member of a resident or other resident representative shall be restricted from participating in meetings in the facility with the families or resident representatives of other residents in the facility;

17. Shall require that each nursing home and certified nursing facility maintain liability insurance coverage in a minimum amount of $1 million, and professional liability coverage in an amount at least

equal to the recovery limit set forth in § 8.01-581.15, to compensate patients or individuals for injuries and losses resulting from the negligent or criminal acts of the facility. Failure to maintain such minimum insurance shall result in revocation of the facility's license;

18. Shall require each hospital that provides obstetrical services to establish policies to follow when a stillbirth, as defined in § 32.1-69.1, occurs that meet the guidelines pertaining to counseling patients and their families and other aspects of managing stillbirths as may be specified by the Board in its regulations;

19. Shall require each nursing home to provide a full refund of any unexpended patient funds on deposit with the facility following the discharge or death of a patient, other than entrance-related fees paid to a continuing care provider as defined in § 38.2-4900, within 30 days of a written request for such funds by the discharged patient or, in the case of the death of a patient, the person administering the person's estate in accordance with the Virginia Small Estates Act (§ 64.2-600 et seq.);

20. Shall require that each hospital that provides inpatient psychiatric services establish a protocol that requires, for any refusal to admit (i) a medically stable patient referred to its psychiatric unit, direct verbal communication between the on-call physician in the psychiatric unit and the referring physician, if requested by such referring physician, and prohibits on-call physicians or other hospital staff from refusing a request for such direct verbal communication by a referring physician and (ii) a patient for whom there is a question regarding the medical stability or medical appropriateness of admission for inpatient psychiatric services due to a situation involving results of a toxicology screening, the on-call physician in the psychiatric unit to which the patient is sought to be transferred to participate in direct verbal communication, either in person or via telephone, with a clinical toxicologist or other person who is a Certified Specialist in Poison Information employed by a poison control center that is accredited by the American Association of Poison Control Centers to review the results of the toxicology screen and determine whether a medical reason for refusing admission to the psychiatric unit related to the results of the toxicology screen exists, if requested by the referring physician;

21. Shall require that each hospital that is equipped to provide life-sustaining treatment shall develop a policy governing determination of the medical and ethical appropriateness of proposed medical care, which shall include (i) a process for obtaining a second opinion regarding the medical and ethical appropriateness of proposed medical care in cases in which a physician has determined proposed care to be medically or ethically inappropriate; (ii) provisions for review of the determination that proposed medical care is medically or ethically inappropriate by an interdisciplinary medical review committee and a determination by the interdisciplinary medical review committee regarding the medical and ethical appropriateness of the proposed health care; and (iii) requirements for a written explanation of the decision reached by the interdisciplinary medical review committee, which shall be included in the patient's medical record. Such policy shall ensure that the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986 (a) are informed of the patient's right to obtain his medical record and to obtain an independent medical opinion and (b) afforded reasonable opportunity to participate in the medical review committee meeting. Nothing in such policy shall prevent the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986 from obtaining legal counsel to represent the patient or from seeking other remedies available at law, including seeking court review, provided that the patient, his agent, or the person authorized to make medical decisions pursuant to § 54.1-2986, or legal counsel provides written notice to the chief executive officer of the hospital within 14 days of the date on which the physician's determination that proposed medical treatment is medically or ethically inappropriate is documented in the patient's medical record;

22. Shall require every hospital with an emergency department to establish protocols to ensure that security personnel of the emergency department, if any, receive training appropriate to the populations served by the emergency department, which may include training based on a trauma-informed approach in identifying and safely addressing situations involving patients or other persons who pose a risk of harm to themselves or others due to mental illness or substance abuse or who are experiencing a mental health crisis;

23. Shall require that each hospital establish a protocol requiring that, before a health care provider arranges for air medical transportation services for a patient who does not have an emergency medical condition as defined in 42 U.S.C. § 1395dd(e)(1), the hospital shall provide the patient or his authorized representative with written or electronic notice that the patient (i) may have a choice of transportation by an air medical transportation provider or medically appropriate ground transportation by an emergency medical services provider and (ii) will be responsible for charges incurred for such transportation in the event that the provider is not a contracted network provider of the patient's health insurance carrier or such charges are not otherwise covered in full or in part by the patient's health insurance plan;

24. Shall establish an exemption from the requirement to obtain a license to add temporary beds in an existing hospital or nursing home, including beds located in a temporary structure or satellite location operated by the hospital or nursing home, provided that the ability remains to safely staff services across the existing hospital or nursing home, (i) for a period of no more than the duration of the Commissioner's determination plus 30 days when the Commissioner has determined that a natural or man-made disaster has caused the evacuation of a hospital or nursing home and that a public health

emergency exists due to a shortage of hospital or nursing home beds or (ii) for a period of no more than the duration of the emergency order entered pursuant to § 32.1-13 or 32.1-20 plus 30 days when the Board, pursuant to § 32.1-13, or the Commissioner, pursuant to § 32.1-20, has entered an emergency order for the purpose of suppressing a nuisance dangerous to public health or a communicable, contagious, or infectious disease or other danger to the public life and health;

25. Shall establish protocols to ensure that any patient scheduled to receive an elective surgical procedure for which the patient can reasonably be expected to require outpatient physical therapy as a follow-up treatment after discharge is informed that he (i) is expected to require outpatient physical therapy as a follow-up treatment and (ii) will be required to select a physical therapy provider prior to being discharged from the hospital;

26. Shall permit nursing home staff members who are authorized to possess, distribute, or administer medications to residents to store, dispense, or administer cannabis oil to a resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*;

27. Shall require each hospital with an emergency department to establish a protocol for the treatment and discharge of individuals experiencing a substance use-related emergency, which shall include provisions for (i) appropriate screening and assessment of individuals experiencing substance use-related emergencies to identify medical interventions necessary for the treatment of the individual in the emergency department and (ii) recommendations for follow-up care following discharge for any patient identified as having a substance use disorder, depression, or mental health disorder, as appropriate, which may include, for patients who have been treated for substance use-related emergencies, including opioid overdose, or other high-risk patients, (a) the dispensing of naloxone or other opioid antagonist used for overdose reversal pursuant to subsection X of § 54.1-3408 at discharge or (b) issuance of a prescription for and information about accessing naloxone or other opioid antagonist used for overdose reversal, including information about accessing naloxone or other opioid antagonist used for overdose reversal at a community pharmacy, including any outpatient pharmacy operated by the hospital, or through a community organization or pharmacy that may dispense naloxone or other opioid antagonist used for overdose reversal without a prescription pursuant to a statewide standing order. Such protocols may also provide for referrals of individuals experiencing a substance use-related emergency to peer recovery specialists and community-based providers of behavioral health services, or to providers of pharmacotherapy for the treatment of drug or alcohol dependence or mental health diagnoses;

28. During a public health emergency related to COVID-19, shall require each nursing home and certified nursing facility to establish a protocol to allow each patient to receive visits, consistent with guidance from the Centers for Disease Control and Prevention and as directed by the Centers for Medicare and Medicaid Services and the Board. Such protocol shall include provisions describing (i) the conditions, including conditions related to the presence of COVID-19 in the nursing home, certified nursing facility, and community, under which in-person visits will be allowed and under which in-person visits will not be allowed and visits will be required to be virtual; (ii) the requirements with which in-person visitors will be required to comply to protect the health and safety of the patients and staff of the nursing home or certified nursing facility; (iii) the types of technology, including interactive audio or video technology, and the staff support necessary to ensure visits are provided as required by this subdivision; and (iv) the steps the nursing home or certified nursing facility will take in the event of a technology failure, service interruption, or documented emergency that prevents visits from occurring as required by this subdivision. Such protocol shall also include (a) a statement of the frequency with which visits, including virtual and in-person, where appropriate, will be allowed, which shall be at least once every 10 calendar days for each patient; (b) a provision authorizing a patient or the patient's personal representative to waive or limit visitation, provided that such waiver or limitation is included in the patient's health record; and (c) a requirement that each nursing home and certified nursing facility publish on its website or communicate to each patient or the patient's authorized representative, in writing or via electronic means, the nursing home's or certified nursing facility's plan for providing visits to patients as required by this subdivision;

29. Shall require each hospital, nursing home, and certified nursing facility to establish and implement policies to ensure the permissible access to and use of an intelligent personal assistant provided by a patient, in accordance with such regulations, while receiving inpatient services. Such policies shall ensure protection of health information in accordance with the requirements of the federal Health Insurance Portability and Accountability Act of 1996, 42 U.S.C. § 1320d et seq., as amended. For the purposes of this subdivision, "intelligent personal assistant" means a combination of an electronic device and a specialized software application designed to assist users with basic tasks using a combination of natural language processing and artificial intelligence, including such combinations known as "digital assistants" or "virtual assistants";

30. During a declared public health emergency related to a communicable disease of public health threat, shall require each hospital, nursing home, and certified nursing facility to establish a protocol to allow patients to receive visits from a rabbi, priest, minister, or clergy of any religious denomination or sect consistent with guidance from the Centers for Disease Control and Prevention and the Centers for

Medicare and Medicaid Services and subject to compliance with any executive order, order of public health, Department guidance, or any other applicable federal or state guidance having the effect of limiting visitation. Such protocol may restrict the frequency and duration of visits and may require visits to be conducted virtually using interactive audio or video technology. Any such protocol may require the person visiting a patient pursuant to this subdivision to comply with all reasonable requirements of the hospital, nursing home, or certified nursing facility adopted to protect the health and safety of the person, patients, and staff of the hospital, nursing home, or certified nursing facility; and

31. Shall require that every hospital that makes health records, as defined in § 32.1-127.1:03, of patients who are minors available to such patients through a secure website shall make such health records available to such patient's parent or guardian through such secure website, unless the hospital cannot make such health record available in a manner that prevents disclosure of information, the disclosure of which has been denied pursuant to subsection F of § 32.1-127.1:03 or for which consent required in accordance with subsection E of § 54.1-2969 has not been provided.

C. Upon obtaining the appropriate license, if applicable, licensed hospitals, nursing homes, and certified nursing facilities may operate adult day care centers.

D. All facilities licensed by the Board pursuant to this article which provide treatment or care for hemophiliacs and, in the course of such treatment, stock clotting factors, shall maintain records of all lot numbers or other unique identifiers for such clotting factors in order that, in the event the lot is found to be contaminated with an infectious agent, those hemophiliacs who have received units of this contaminated clotting factor may be apprised of this contamination. Facilities which have identified a lot that is known to be contaminated shall notify the recipient's attending physician and request that he notify the recipient of the contamination. If the physician is unavailable, the facility shall notify by mail, return receipt requested, each recipient who received treatment from a known contaminated lot at the individual's last known address.

E. Hospitals in the Commonwealth may enter into agreements with the Department of Health for the provision to uninsured patients of naloxone or other opioid antagonists used for overdose reversal.

§ 32.1-162.6:1. Possession or administration of cannabis oil.

Hospice and hospice facility employees who are authorized to possess, distribute, or administer medications to patients shall be permitted to store, dispense, or administer cannabis oil to a patient who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of § 54.1-3408.3 and has registered with the Board of Pharmacy~~ *§ 4.1-1601*.

§ 40.1-27.4. Discipline for employee's medicinal use of cannabis oil prohibited.

A. As used in this section, "cannabis oil" means the same as that term is defined in § ~~54.1-3408.3~~ *4.1-1600*.

B. No employer shall discharge, discipline, or discriminate against an employee for such employee's lawful use of cannabis oil pursuant to a valid written certification issued by a practitioner for the treatment or to eliminate the symptoms of the employee's diagnosed condition or disease pursuant to § ~~54.1-3408.3~~ *4.1-1601*.

C. Notwithstanding the provisions of subsection B, nothing in this section shall (i) restrict an employer's ability to take any adverse employment action for any work impairment caused by the use of cannabis oil or to prohibit possession during work hours, (ii) require an employer to commit any act that would cause the employer to be in violation of federal law or that would result in the loss of a federal contract or federal funding, or (iii) require any defense industrial base sector employer or prospective employer, as defined by the U.S. Cybersecurity and Infrastructure Security Agency, to hire or retain any applicant or employee who tests positive for tetrahydrocannabinol (THC) in excess of 50 ng/ml for a urine test or 10 pg/mg for a hair test.

§ 46.2-341.20:7. Possession of marijuana in commercial motor vehicle unlawful; civil penalty.

A. It is unlawful for any person to knowingly or intentionally possess marijuana in a commercial motor vehicle as defined in § 46.2-341.4. The attorney for the Commonwealth or the county, city, or town attorney may prosecute such a case.

Upon the prosecution of a person for a violation of this section, ownership or occupancy of the vehicle in which marijuana was found shall not create a presumption that such person either knowingly or intentionally possessed such marijuana.

Any person who violates this section is subject to a civil penalty of no more than $25. A violation of this section is a civil offence. Any civil penalties collected pursuant to this section shall be deposited into the Drug Offender Assessment and Treatment Fund established pursuant to § 18.2-251.02. Violations of this section by an adult shall be prepayable according to the procedures in § 16.1-69.40:2.

B. Any violation of this section shall be charged by summons. A summons for a violation of this section may be executed by a law-enforcement officer when such violation is observed by such officer. The summons used by a law-enforcement officer pursuant to this section shall be in form the same as the uniform summons for motor vehicle law violations as prescribed pursuant to § 46.2-388. No court costs shall be assessed for violations of this section. A person's criminal history record information as defined in § 9.1-101 shall not include records of any charges or judgments for a violation of this section, and records of such charges or judgments shall not be reported to the Central Criminal Records

Exchange; however, such violation shall be reported to the Department of Motor Vehicles and shall be included on such individual's driving record.

C. The procedure for appeal and trial of any violation of this section shall be the same as provided by law for misdemeanors; if requested by either party on appeal to the circuit court, trial by jury shall be as provided in Article 4 (§ 19.2-260 et seq.) of Chapter 15 of Title 19.2, and the Commonwealth shall be required to prove its case beyond a reasonable doubt.

D. The provisions of this section shall not apply to members of state, federal, county, city, or town law-enforcement agencies, jail officers, or correctional officers, as defined in § 53.1-1, certified as handlers of dogs trained in the detection of controlled substances when possession of marijuana is necessary for the performance of their duties.

E. The provisions of this section involving marijuana in the form of cannabis products as that term is defined in § ~~54.1-3408.3~~ *4.1-1600* shall not apply to any person who possesses such cannabis product pursuant to a valid written certification issued by a practitioner in the course of his professional practice pursuant to § ~~54.1-3408.3~~ *4.1-1601* for treatment or to alleviate the symptoms of (i) the person's diagnosed condition or disease, (ii) if such person is the parent or guardian of a minor or of a vulnerable adult as defined in § 18.2-369, such minor's or vulnerable adult's diagnosed condition or disease, or (iii) if such person has been designated as a registered agent pursuant to § ~~54.1-3408.3~~ *4.1-1601,* the diagnosed condition or disease of his principal or, if the principal is the parent or legal guardian of a minor or of a vulnerable adult as defined in § 18.2-369, such minor's or vulnerable adult's diagnosed condition or disease.

§ 54.1-2522.1. (Effective until July 1, 2027) Requirements of practitioners.

A. Any prescriber who is licensed in the Commonwealth to treat human patients and is authorized pursuant to §§ 54.1-3303 and 54.1-3408 to issue a prescription for a covered substance shall be registered with the Prescription Monitoring Program by the Department of Health Professions.

B. A prescriber registered with the Prescription Monitoring Program or a person to whom he has delegated authority to access information in the possession of the Prescription Monitoring Program pursuant to § 54.1-2523.2 shall, at the time of initiating a new course of treatment to a human patient that includes the prescribing of opioids anticipated at the onset of treatment to last more than seven consecutive days, request information from the Director for the purpose of determining what, if any, other covered substances are currently prescribed to the patient. In addition, any prescriber who holds a special identification number from the Drug Enforcement Administration authorizing the prescribing of controlled substances approved for use in opioid addiction therapy shall, prior to or as a part of execution of a treatment agreement with the patient, request information from the Director for the purpose of determining what, if any, other covered substances the patient is currently being prescribed. Nothing in this section shall prohibit prescribers from making additional periodic requests for information from the Director as may be required by routine prescribing practices.

C. A prescriber shall not be required to meet the provisions of subsection B if:

1. The opioid is prescribed to a patient currently receiving hospice or palliative care;

2. The opioid is prescribed to a patient during an inpatient hospital admission or at discharge;

3. The opioid is prescribed to a patient in a nursing home or a patient in an assisted living facility that uses a sole source pharmacy;

4. The Prescription Monitoring Program is not operational or available due to temporary technological or electrical failure or natural disaster; or

5. The prescriber is unable to access the Prescription Monitoring Program due to emergency or disaster and documents such circumstances in the patient's medical record.

D. Prior to issuing a written certification for the use of cannabis oil in accordance with § ~~54.1-3408.3~~ *4.1-1601*, a practitioner shall request information from the Director for the purpose of determining what, if any, other covered substances have been dispensed to the patient.

§ 54.1-2522.1. (Effective July 1, 2027) Requirements of practitioners.

A. Any prescriber who is licensed in the Commonwealth to treat human patients and is authorized pursuant to §§ 54.1-3303 and 54.1-3408 to issue a prescription for a covered substance shall be registered with the Prescription Monitoring Program by the Department of Health Professions.

B. Prescribers registered with the Prescription Monitoring Program shall, at the time of initiating a new course of treatment to a human patient that includes the prescribing of benzodiazepine or an opiate anticipated at the onset of treatment to last more than 90 consecutive days, request information from the Director for the purpose of determining what, if any, other covered substances are currently prescribed to the patient. In addition, any prescriber who holds a special identification number from the Drug Enforcement Administration authorizing the prescribing of controlled substances approved for use in opioid addiction therapy shall, prior to or as a part of execution of a treatment agreement with the patient, request information from the Director for the purpose of determining what, if any, other covered substances the patient is currently being prescribed. Nothing in this section shall prohibit prescribers from making additional periodic requests for information from the Director as may be required by routine prescribing practices.

C. The Secretary of Health and Human Resources may identify and publish a list of benzodiazepines

or opiates that have a low potential for abuse by human patients. Prescribers who prescribe such identified benzodiazepines or opiates shall not be required to meet the provisions of subsection B. In addition, a prescriber shall not be required to meet the provisions of subsection B if the course of treatment arises from pain management relating to dialysis or cancer treatments.

D. Prior to issuing a written certification for the use of cannabis oil in accordance with § ~~54.1-3408.3~~ *4.1-1601*, a practitioner shall request information from the Director for the purpose of determining what, if any, other covered substances have been dispensed to the patient.

§ 54.1-2903. What constitutes practice; advertising in connection with medical practice.

A. Any person shall be regarded as practicing the healing arts who actually engages in such practice as defined in this chapter, or who opens an office for such purpose, or who advertises or announces to the public in any manner a readiness to practice or who uses in connection with his name the words or letters "Doctor," "Dr.," "M.D.," "D.O.," "D.P.M.," "D.C.," "Healer," "N.P.," or any other title, word, letter or designation intending to designate or imply that he is a practitioner of the healing arts or that he is able to heal, cure or relieve those suffering from any injury, deformity or disease.

Signing a birth or death certificate, or signing any statement certifying that the person so signing has rendered professional service to the sick or injured, or signing or issuing a prescription for drugs or other remedial agents, shall be prima facie evidence that the person signing or issuing such writing is practicing the healing arts within the meaning of this chapter except where persons other than physicians are required to sign birth certificates.

B. No person regulated under this chapter shall use the title "Doctor" or the abbreviation "Dr." in writing or in advertising in connection with his practice unless he simultaneously uses words, initials, an abbreviation or designation, or other language that identifies the type of practice for which he is licensed. No person regulated under this chapter shall include in any advertisement a reference to marijuana, as defined in § 18.2-247, unless such advertisement is for the treatment of addiction or substance abuse. However, nothing in this subsection shall prevent a person from including in any advertisement that such person is registered with the Board of ~~Pharmacy~~ *Directors of the Virginia Cannabis Control Authority* to issue written certifications for the use of cannabis products, as defined in § ~~54.1-3408.3~~ *4.1-1600*.

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

A. As used in this section:~~"Botanical~~, *"botanical* cannabis," ~~means cannabis that is composed wholly of usable cannabis from the same parts of the same chemovar of cannabis plant~~ *"cannabis oil," "cannabis product," and "practitioner" mean the same as those terms are defined in § 4.1-1600*.

~~"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 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 *in accordance with the provisions of § 4.1-1601*. ~~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.

§ 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 of~~ Chapter ~~34~~ *16 (§ 4.1-1600 et seq.)* of Title ~~54.1 of the Code of Virginia~~ *4.1*;
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 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 of~~ Chapter ~~34~~ *16 (§ 4.1-1600 et seq.)* of Title ~~54.1 of the Code of Virginia~~ *4.1*;
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.

§ 63.2-1803.01. Possession or administration of cannabis oil.

Assisted living facility staff members who are authorized to possess, distribute, or administer medications to residents in accordance with the facility's written plan for medication management shall be permitted to store, dispense, or administer cannabis oil to a resident who has been issued a valid written certification for the use of cannabis oil in accordance with ~~subsection B of~~ § ~~54.1-3408.3~~ *4.1-1601* and has registered with the Board of ~~Pharmacy~~ *Directors of the Virginia Cannabis Control Authority*.

2. That Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia is repealed.

3. That the provisions of the first and second enactments of this act shall become effective on January 1, 2024.

4. That the twenty-first enactment of Chapter 550 and the twenty-first enactment of Chapter 551 of the Acts of Assembly of 2021, Special Session I, are repealed.

5. That the Regulations Governing Pharmaceutical Processors (18VAC110-60) as promulgated or amended by the Board of Pharmacy prior to January 1, 2024, shall remain in full force and effect and shall be administered by the Virginia Cannabis Control Authority (the Authority) until the Board of Directors (the Board) of the Authority promulgates regulations to implement the provisions of this act, which shall model, to the greatest extent practicable, the Regulations Governing Pharmaceutical Processors (18VAC110-60) promulgated by the Board of Pharmacy. With the exception of § 2.2-4031 of the Code of Virginia, neither the provisions of the

Administrative Process Act (§ 2.2-4000 et seq. of the Code of Virginia) nor public participation guidelines adopted pursuant thereto shall apply to the Board's initial adoption of regulations to implement the provisions of this act. The Authority shall be vested with all powers and duties held by the Board of Pharmacy prior to January 1, 2024, in its administration of the provisions set forth in § 54.1-3408.3 of the Code of Virginia, as amended by this act, Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia, as repealed by this act, and any regulations promulgated pursuant thereto.

6. That any valid, active permits, certifications, and registrations issued by the Board of Pharmacy pursuant to § 54.1-3408.3 of the Code of Virginia, as amended by this act, Article 4.2 (§§ 54.1-3442.5 through 54.1-3442.8) of Chapter 34 of Title 54.1 of the Code of Virginia, as repealed by this act, or regulations promulgated pursuant thereto prior to January 1, 2024, shall remain valid until their expiration date and be considered to have been issued by the Board of Directors of the Virginia Cannabis Control Authority.

7. That the Virginia Cannabis Control Authority may assess and collect regulatory fees from each pharmaceutical processor and cannabis dispensing facility in an amount sufficient to implement this act.

CHAPTER 760

An Act to amend and reenact §§ 54.1-3408.3, 54.1-3442.5, 54.1-3442.6, and 54.1-3442.7 of the Code of Virginia and to amend the Code of Virginia by adding sections numbered 54.1-3442.7:1, 54.1-3442.7:2, and 54.1-3442.7:3, relating to medical marijuana program; product, registration, dispensing, and recordkeeping requirements; advertising.

[S 1337]

Approved April 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That §§ 54.1-3408.3, 54.1-3442.5, 54.1-3442.6, and 54.1-3442.7 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding sections numbered 54.1-3442.7:1, 54.1-3442.7:2, and 54.1-3442.7:3 as follows:

§ 54.1-3408.3. Certification for use of cannabis products 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 dispensing facility" means the same as that term is defined in § 54.1-3442.5.

"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, *except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.),* 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) *is formulated with cannabis oil or botanical cannabis; (ii) is* produced by a pharmaceutical processor~~,~~ *and sold by a pharmaceutical processor or cannabis dispensing facility; (iii) is* registered with the Board~~,~~*; (iv) contains, except as otherwise provided in Article 4.2 (§ 54.1-3442.5 et seq.), no more than 10 milligrams of tetrahydrocannabinol per dose;* and *(v) is* 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.

"Pharmaceutical processor" means the same as that term is defined in § 54.1-3442.5.

"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. *No practitioner may issue a written certification while such practitioner is on the premises of a pharmaceutical processor or cannabis dispensing facility. A pharmaceutical processor shall not endorse or promote any practitioner who issues certifications to patients.* 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. *A*

practitioner who issues written certifications shall not directly or indirectly accept, solicit, or receive anything of value from a pharmaceutical processor, cannabis dispensing facility, or any person associated with a pharmaceutical processor, cannabis dispensing facility, or provider of paraphernalia, excluding information on products or educational materials on the benefits and risks of cannabis products.

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~~ *a practitioner's professional licensing board* from sanctioning ~~a~~ *the* 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~~ *(i)* shall hold sufficient education and training to exercise appropriate professional judgment in the certification of patients*; (ii) shall not offer a discount or any other thing of value to a patient or a patient's parent, guardian, or registered agent that is contingent on or encourages the person's decision to use a particular pharmaceutical processor or cannabis product; (iii) shall not issue a certification to himself or his family members, employees, or coworkers; (iv) shall not provide product samples containing cannabis other than those approved by the U.S. Food and Drug Administration; and (v) shall not accept compensation from a pharmaceutical processor or cannabis dispensing facility*. 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 *patient certification or agent* 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.

§ 54.1-3442.5. Definitions.

As used in this article:

"Botanical cannabis," "cannabis oil," "cannabis product," *"designated caregiver facility," "practitioner," "registered agent,"* and "usable cannabis" have the same meanings as specified in § 54.1-3408.3.

"Cannabis dispensing facility" means a facility that (i) has obtained a permit from the Board pursuant to § 54.1-3442.6; (ii) is owned, at least in part, by a pharmaceutical processor; and (iii) dispenses

cannabis products produced by a pharmaceutical processor 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.

~~"Designated caregiver facility" has the same meaning as defined in § 54.1-3408.3.~~

"Pharmaceutical processor" means a facility that (i) has obtained a permit from the Board pursuant to § ~~54.1-3408.3~~ *54.1-3442.6* and (ii) cultivates Cannabis plants intended only for the production of cannabis oil, botanical cannabis, and usable cannabis, produces cannabis products, and dispenses cannabis products to a patient pursuant to a written certification, 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.

~~"Practitioner" has the same meaning as specified in § 54.1-3408.3.~~

~~"Registered agent" has the same meaning as specified in § 54.1-3408.3.~~

§ 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~~ *products* that provide that each dispensed dose of *a* cannabis ~~oil~~ *product* not exceed 10 milligrams of ~~delta-9-tetrahydrocannabinol~~ *total tetrahydrocannabinol, except as permitted under § 54.1-3442.7:2*; (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. *Remediated botanical cannabis or cannabis oil that passes such quality testing may be packaged and labeled.* 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~~ *15* percent deviation basis, of ~~active~~ ~~ingredients~~ *total THC and total CBD. No cannabis product shall have an expiration date longer than six months from the date of the cannabis product registration approval unless supported by stability testing.*

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 *unless all cannabis products are contained in a vault or other similar container to which only the pharmacist has access controls*. 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~~ *and cannabis* 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~~ *total tetrahydrocannabinol* in any cannabis product on site may be up to ~~10~~ *15* percent greater than or less than the level of ~~delta-9-tetrahydrocannabinol measured for labeling~~ *total tetrahydrocannabinol listed in the approved cannabis product registration*. 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 *that have an expiration date longer than six months*.

§ 54.1-3442.7:1. Packaging and labeling; corrections; records.

A. Pharmaceutical processors shall comply with all packaging and labeling requirements set forth in this article and Board regulations.

B. No cannabis product shall be 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 cannabis product.

C. Pharmaceutical processors may correct typographical errors made on cannabis product labels and any documents generated as the result of a wholesale transaction.

§ 54.1-3442.7:2. Cannabis product registration; approval, deviation, and modification.

A. A pharmaceutical processor shall register with the Board each cannabis product it manufactures. Applications for cannabis product registration shall be submitted to the Board on a form prescribed by the Board.

B. An application for cannabis product registration shall include:

1. The total tetrahydrocannabinol and total cannabidiol in such cannabis product, based on laboratory testing results for the cannabis product formulation;

2. A product name;

3. A proposed product package; and

4. A proposed product label, which shall not be required to contain an expiration date at the time of application.

C. The Board shall register all cannabis products that meet testing, labeling, and packaging standards after an application for registration is submitted. If the cannabis product fails to meet such standards or the application was deficient, the Board shall notify the applicant of the specific reasons for such failure or deficiency.

D. Within two business days of the Board's approval or deemed approval, the Board shall enter the cannabis product's national drug code number into the Prescription Monitoring Program.

E. The following cannabis product deviations from an approved cannabis product registration shall be permitted without any requirement for a new cannabis product registration or notice to the Board:

1. A deviation in the concentration of total tetrahydrocannabinol (THC) or total cannabidiol (CBD) in a cannabis product or dose thereof of up to 15 percent greater than or less than the concentration of total tetrahydrocannabinol or total cannabidiol, either or both, listed in the approved cannabis product registration; however, for a cannabis product with five milligrams or less of total THC or total CBD per dose, the total THC or total CBD concentration shall be within 0.5 milligrams of the single dose total THC or total CBD concentrations approved for that cannabis product;

2. A variation in packaging, provided that the packaging is substantially similar to the approved packaging and otherwise complies with applicable packaging requirements;

3. A deviation in labeling, including a variation made in accordance with § 54.1-3442.7:1, that reflects allowable deviations in total THC or total CBD or that makes a minor text, font, design, or similar modification, provided that the labeling is substantially similar to the approved labeling and otherwise complies with applicable labeling requirements; and

4. Any other insignificant changes.

F. A pharmaceutical processor may submit a request to modify an existing cannabis product registration in the event of a cannabis product deviation that is not set forth in subsection E. Upon receipt, the Board shall respond to such request. The Board may grant or deny the request, propose a reasonable revision, or require the pharmaceutical processor to provide additional information.

§ 54.1-3442.7:3. Advertising and marketing.

A. Pharmaceutical processors and cannabis dispensing facilities may (i) advertise and promote products and operations and (ii) provide educational material to practitioners, patients, and the public.

B. Pharmaceutical processors and cannabis dispensing facilities may engage in advertising or marketing that does not:

1. Include false or misleading statements;

2. Promote overconsumption;

3. Depict a person younger than 21 years of age;

4. Appeal particularly to persons younger than 21 years of age, including by using cartoons in any way;

5. Associate cannabis products with candy or similar products or depicts any images that bear a reasonable resemblance to a candy or similar product; or

6. Contain any seal, flag, crest, coat of arms, or other insignia that is likely to mislead patients or the public to believe that the cannabis product is made or endorsed by the Commonwealth.

C. All advertising and marketing by pharmaceutical processors and cannabis dispensing facilities shall (i) accurately and legibly identify the pharmaceutical processor or cannabis dispensing facility responsible for its content, (ii) include a statement that cannabis products are for use by certified patients only, and (iii) comply with Board regulations.

2. That pharmaceutical processors and cannabis dispensing facilities shall collect and provide to the Board of Pharmacy by July 1, 2024, data regarding the impact of this act on program participation, reductions in the price of cannabis products, and improved operational efficiencies.

3. That the Board of Pharmacy shall amend its regulations, including 18VAC110-60-270, 18VAC110-60-285, 18VAC110-60-290, and 18VAC110-60-310, to replace any references to "brand" with "registered cannabis product name."

4. That the Board of Pharmacy may assess and collect regulatory fees from each pharmaceutical processor and cannabis dispensing facility in an amount sufficient to implement the provisions of this act.

VIRGINIA ACTS OF ASSEMBLY -- 2023 RECONVENED SESSION

CHAPTER 812

An Act to amend and reenact § 54.1-3442.6 of the Code of Virginia, relating to medical marijuana program; additional cultivation facility.

[S 1533]

Approved May 12, 2023

Be it enacted by the General Assembly of Virginia:

1. That § 54.1-3442.6 of the Code of Virginia is amended and reenacted as follows:

§ 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; (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 *(i)* establish up to five cannabis dispensing facilities, *subject to the permit requirement set forth in subsection B,* for the dispensing of cannabis products that have been cultivated and produced on the premises of a pharmaceutical processor permitted by the Board *and (ii) establish, if authorized by the Board, one additional location at which the pharmaceutical processor may cultivate cannabis plants.* Each cannabis dispensing facility *and the additional cultivation location* 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.

2. That the Board of Pharmacy (the Board) shall consider the following factors when determining whether to authorize an additional cannabis cultivation location: (i) the location of the proposed additional cannabis cultivation location, (ii) the distance of the proposed additional cannabis cultivation location from the pharmaceutical processor, (iii) access to a secure transportation network between the proposed additional cannabis cultivation location and the pharmaceutical processor, (iv) whether the proposed location for the additional cannabis cultivation facility is economically viable, (v) whether there is a demonstrated demand for additional cannabis cultivation, and (vi) any additional factors determined by the Board to be relevant in making its determination.