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**HOUSE BILL NO. 1489**

Offered January 11, 2023

Prefiled January 3, 2023

*A BILL to amend and reenact §§ 54.1-2900, 54.1-2901, 54.1-2914, 54.1-2973.1, and 54.1-3401 of the Code of Virginia and to amend the Code of Virginia by adding sections numbered 54.1-2956.15 through 54.1-2956.18, relating to practice of naturopathic medicine.*

Patron—Davis

Referred to Committee on Health, Welfare and Institutions

**Be it enacted by the General Assembly of Virginia:**

**1. That §§ 54.1-2900, 54.1-2901, 54.1-2914, 54.1-2973.1, and 54.1-3401 of the Code of Virginia are amended and reenacted and that the Code of Virginia is amended by adding sections numbered 54.1-2956.15 through 54.1-2956.18 as follows:**

**§ 54.1-2900. Definitions.**

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

"Acupuncturist" means an individual approved by the Board to practice acupuncture. This is limited to "licensed acupuncturist" which means an individual other than a doctor of medicine, osteopathy, chiropractic or podiatry who has successfully completed the requirements for licensure established by the Board (approved titles are limited to: Licensed Acupuncturist, Lic.Ac., and L.Ac.).

"Auricular acupuncture" means the subcutaneous insertion of sterile, disposable acupuncture needles in predetermined, bilateral locations in the outer ear when used exclusively and specifically in the context of a chemical dependency treatment program.

"Birth control" means contraceptive methods that are approved by the U.S. Food and Drug Administration. "Birth control" shall not be considered abortion for the purposes of Title 18.2.

"Board" means the Board of Medicine.

"Certified nurse midwife" means an advanced practice registered nurse who is certified in the specialty of nurse midwifery and who is jointly licensed by the Boards of Medicine and Nursing as a nurse practitioner pursuant to § 54.1-2957.

"Certified registered nurse anesthetist" means an advanced practice registered nurse who is certified in the specialty of nurse anesthesia, who is jointly licensed by the Boards of Medicine and Nursing as a nurse practitioner pursuant to § 54.1-2957, and who practices under the supervision of a doctor of medicine, osteopathy, podiatry, or dentistry but is not subject to the practice agreement requirement described in § 54.1-2957.

"Clinical nurse specialist" means an advance practice registered nurse who is certified in the specialty of clinical nurse specialist and who is jointly licensed by the Boards of Medicine and Nursing as a nurse practitioner pursuant to § 54.1-2957.

"Collaboration" means the communication and decision-making process among health care providers who are members of a patient care team related to the treatment of a patient that includes the degree of cooperation necessary to provide treatment and care of the patient and includes (i) communication of data and information about the treatment and care of a patient, including the exchange of clinical observations and assessments, and (ii) development of an appropriate plan of care, including decisions regarding the health care provided, accessing and assessment of appropriate additional resources or expertise, and arrangement of appropriate referrals, testing, or studies.

"Consultation" means communicating data and information, exchanging clinical observations and assessments, accessing and assessing additional resources and expertise, problem-solving, and arranging for referrals, testing, or studies.

"Genetic counselor" means a person licensed by the Board to engage in the practice of genetic counseling.

"Healing arts" means the arts and sciences dealing with the prevention, diagnosis, treatment and cure or alleviation of human physical or mental ailments, conditions, diseases, pain or infirmities.

"Licensed certified midwife" means a person who is licensed as a certified midwife by the Boards of Medicine and Nursing.

"Medical malpractice judgment" means any final order of any court entering judgment against a licensee of the Board that arises out of any tort action or breach of contract action for personal injuries or wrongful death, based on health care or professional services rendered, or that should have been rendered, by a health care provider, to a patient.

"Medical malpractice settlement" means any written agreement and release entered into by or on

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59 behalf of a licensee of the Board in response to a written claim for money damages that arises out of  
60 any personal injuries or wrongful death, based on health care or professional services rendered, or that  
61 should have been rendered, by a health care provider, to a patient.

62 *"Naturopathic doctor" means an individual, other than a doctor of medicine, osteopathy,*  
63 *chiropractic, or podiatry, who is licensed by the Board to practice naturopathic medicine and authorized*  
64 *by the Board to diagnose, treat, and help prevent diseases using a system of practice that is based on*  
65 *the natural healing capacity of individuals, using physiological, psychological, or physical methods, and*  
66 *who may also use natural medicines, prescriptions, legend drugs, foods, herbs, or other natural*  
67 *remedies, including light and air.*

68 "Nurse practitioner" means an advanced practice registered nurse who is jointly licensed by the  
69 Boards of Medicine and Nursing pursuant to § 54.1-2957.

70 "Occupational therapy assistant" means an individual who has met the requirements of the Board for  
71 licensure and who works under the supervision of a licensed occupational therapist to assist in the  
72 practice of occupational therapy.

73 "Patient care team" means a multidisciplinary team of health care providers actively functioning as a  
74 unit with the management and leadership of one or more patient care team physicians for the purpose of  
75 providing and delivering health care to a patient or group of patients.

76 "Patient care team physician" means a physician who is actively licensed to practice medicine in the  
77 Commonwealth, who regularly practices medicine in the Commonwealth, and who provides management  
78 and leadership in the care of patients as part of a patient care team.

79 "Patient care team podiatrist" means a podiatrist who is actively licensed to practice podiatry in the  
80 Commonwealth, who regularly practices podiatry in the Commonwealth, and who provides management  
81 and leadership in the care of patients as part of a patient care team.

82 "Physician assistant" means a health care professional who has met the requirements of the Board for  
83 licensure as a physician assistant.

84 "Practice of acupuncture" means the stimulation of certain points on or near the surface of the body  
85 by the insertion of needles to prevent or modify the perception of pain or to normalize physiological  
86 functions, including pain control, for the treatment of certain ailments or conditions of the body and  
87 includes the techniques of electroacupuncture, cupping and moxibustion. The practice of acupuncture  
88 does not include the use of physical therapy, chiropractic, or osteopathic manipulative techniques; the  
89 use or prescribing of any drugs, medications, serums or vaccines; or the procedure of auricular  
90 acupuncture as exempted in § 54.1-2901 when used in the context of a chemical dependency treatment  
91 program for patients eligible for federal, state or local public funds by an employee of the program who  
92 is trained and approved by the National Acupuncture Detoxification Association or an equivalent  
93 certifying body.

94 "Practice of athletic training" means the prevention, recognition, evaluation, and treatment of injuries  
95 or conditions related to athletic or recreational activity that requires physical skill and utilizes strength,  
96 power, endurance, speed, flexibility, range of motion or agility or a substantially similar injury or  
97 condition resulting from occupational activity immediately upon the onset of such injury or condition;  
98 and subsequent treatment and rehabilitation of such injuries or conditions under the direction of the  
99 patient's physician or under the direction of any doctor of medicine, osteopathy, chiropractic, podiatry, or  
100 dentistry, while using heat, light, sound, cold, electricity, exercise or mechanical or other devices.

101 "Practice of behavior analysis" means the design, implementation, and evaluation of environmental  
102 modifications, using behavioral stimuli and consequences, to produce socially significant improvement in  
103 human behavior, including the use of direct observation, measurement, and functional analysis of the  
104 relationship between environment and behavior.

105 "Practice of chiropractic" means the adjustment of the 24 movable vertebrae of the spinal column,  
106 and assisting nature for the purpose of normalizing the transmission of nerve energy, but does not  
107 include the use of surgery, obstetrics, osteopathy, or the administration or prescribing of any drugs,  
108 medicines, serums, or vaccines. "Practice of chiropractic" shall include (i) requesting, receiving, and  
109 reviewing a patient's medical and physical history, including information related to past surgical and  
110 nonsurgical treatment of the patient and controlled substances prescribed to the patient, and (ii)  
111 documenting in a patient's record information related to the condition and symptoms of the patient, the  
112 examination and evaluation of the patient made by the doctor of chiropractic, and treatment provided to  
113 the patient by the doctor of chiropractic. "Practice of chiropractic" shall also include performing the  
114 physical examination of an applicant for a commercial driver's license or commercial learner's permit  
115 pursuant to § 46.2-341.12 if the practitioner has (i) applied for and received certification as a medical  
116 examiner pursuant to 49 C.F.R. Part 390, Subpart D and (ii) registered with the National Registry of  
117 Certified Medical Examiners.

118 "Practice of genetic counseling" means (i) obtaining and evaluating individual and family medical  
119 histories to assess the risk of genetic medical conditions and diseases in a patient, his offspring, and  
120 other family members; (ii) discussing the features, history, diagnosis, environmental factors, and risk

management of genetic medical conditions and diseases; (iii) ordering genetic laboratory tests and other diagnostic studies necessary for genetic assessment; (iv) integrating the results with personal and family medical history to assess and communicate risk factors for genetic medical conditions and diseases; (v) evaluating the patient's and family's responses to the medical condition or risk of recurrence and providing client-centered counseling and anticipatory guidance; (vi) identifying and utilizing community resources that provide medical, educational, financial, and psychosocial support and advocacy; and (vii) providing written documentation of medical, genetic, and counseling information for families and health care professionals.

"Practice of licensed certified midwifery" means the provision of primary health care for preadolescents, adolescents, and adults within the scope of practice of a certified midwife established in accordance with the Standards for the Practice of Midwifery set by the American College of Nurse-Midwives, including (i) providing sexual and reproductive care and care during pregnancy and childbirth, postpartum care, and care for the newborn for up to 28 days following the birth of the child; (ii) prescribing of pharmacological and non-pharmacological therapies within the scope of the practice of midwifery; (iii) consulting or collaborating with or referring patients to such other health care providers as may be appropriate for the care of the patients; and (iv) serving as an educator in the theory and practice of midwifery.

"Practice of medicine or osteopathic medicine" means the prevention, diagnosis, and treatment of human physical or mental ailments, conditions, diseases, pain, or infirmities by any means or method.

*"Practice of naturopathic medicine" means a system of primary health care for the (i) prevention, diagnosis, and treatment of human health conditions, injury, and disease; (ii) promotion or restoration of health; and (iii) support and stimulation of a patient's inherent self-healing process through patient education and use of naturopathic therapies and therapeutic substances.*

"Practice of occupational therapy" means the therapeutic use of occupations for habilitation and rehabilitation to enhance physical health, mental health, and cognitive functioning and includes the evaluation, analysis, assessment, and delivery of education and training in basic and instrumental activities of daily living; the design, fabrication, and application of orthoses (splints); the design, selection, and use of adaptive equipment and assistive technologies; therapeutic activities to enhance functional performance; vocational evaluation and training; and consultation concerning the adaptation of physical, sensory, and social environments.

"Practice of podiatry" means the prevention, diagnosis, treatment, and cure or alleviation of physical conditions, diseases, pain, or infirmities of the human foot and ankle, including the medical, mechanical and surgical treatment of the ailments of the human foot and ankle, but does not include amputation of the foot proximal to the transmetatarsal level through the metatarsal shafts. Amputations proximal to the metatarsal-phalangeal joints may only be performed in a hospital or ambulatory surgery facility accredited by an organization listed in § 54.1-2939. The practice includes the diagnosis and treatment of lower extremity ulcers; however, the treatment of severe lower extremity ulcers proximal to the foot and ankle may only be performed by appropriately trained, credentialed podiatrists in an approved hospital or ambulatory surgery center at which the podiatrist has privileges, as described in § 54.1-2939. The Board of Medicine shall determine whether a specific type of treatment of the foot and ankle is within the scope of practice of podiatry.

"Practice of radiologic technology" means the application of ionizing radiation to human beings for diagnostic or therapeutic purposes.

"Practice of respiratory care" means the (i) administration of pharmacological, diagnostic, and therapeutic agents related to respiratory care procedures necessary to implement a treatment, disease prevention, pulmonary rehabilitative, or diagnostic regimen prescribed by a practitioner of medicine or osteopathic medicine; (ii) transcription and implementation of the written or verbal orders of a practitioner of medicine or osteopathic medicine pertaining to the practice of respiratory care; (iii) observation and monitoring of signs and symptoms, general behavior, general physical response to respiratory care treatment and diagnostic testing, including determination of whether such signs, symptoms, reactions, behavior or general physical response exhibit abnormal characteristics; and (iv) implementation of respiratory care procedures, based on observed abnormalities, or appropriate reporting, referral, respiratory care protocols or changes in treatment pursuant to the written or verbal orders by a licensed practitioner of medicine or osteopathic medicine or the initiation of emergency procedures, pursuant to the Board's regulations or as otherwise authorized by law. The practice of respiratory care may be performed in any clinic, hospital, skilled nursing facility, private dwelling or other place deemed appropriate by the Board in accordance with the written or verbal order of a practitioner of medicine or osteopathic medicine, and shall be performed under qualified medical direction.

"Practice of surgical assisting" means the performance of significant surgical tasks, including manipulation of organs, suturing of tissue, placement of hemostatic agents, injection of local anesthetic, harvesting of veins, implementation of devices, and other duties as directed by a licensed doctor of

182 medicine, osteopathy, or podiatry under the direct supervision of a licensed doctor of medicine,  
183 osteopathy, or podiatry.

184 "Qualified medical direction" means, in the context of the practice of respiratory care, having readily  
185 accessible to the respiratory therapist a licensed practitioner of medicine or osteopathic medicine who  
186 has specialty training or experience in the management of acute and chronic respiratory disorders and  
187 who is responsible for the quality, safety, and appropriateness of the respiratory services provided by the  
188 respiratory therapist.

189 "Radiologic technologist" means an individual, other than a licensed doctor of medicine, osteopathy,  
190 podiatry, or chiropractic or a dentist licensed pursuant to Chapter 27 (§ 54.1-2700 et seq.), who (i)  
191 performs, may be called upon to perform, or is licensed to perform a comprehensive scope of diagnostic  
192 or therapeutic radiologic procedures employing ionizing radiation and (ii) is delegated or exercises  
193 responsibility for the operation of radiation-generating equipment, the shielding of patient and staff from  
194 unnecessary radiation, the appropriate exposure of radiographs, the administration of radioactive  
195 chemical compounds under the direction of an authorized user as specified by regulations of the  
196 Department of Health, or other procedures that contribute to any significant extent to the site or dosage  
197 of ionizing radiation to which a patient is exposed.

198 "Radiologic technologist, limited" means an individual, other than a licensed radiologic technologist,  
199 dental hygienist, or person who is otherwise authorized by the Board of Dentistry under Chapter 27  
200 (§ 54.1-2700 et seq.) and the regulations pursuant thereto, who performs diagnostic radiographic  
201 procedures employing equipment that emits ionizing radiation that is limited to specific areas of the  
202 human body.

203 "Radiologist assistant" means an individual who has met the requirements of the Board for licensure  
204 as an advanced-level radiologic technologist and who, under the direct supervision of a licensed doctor  
205 of medicine or osteopathy specializing in the field of radiology, is authorized to (i) assess and evaluate  
206 the physiological and psychological responsiveness of patients undergoing radiologic procedures; (ii)  
207 evaluate image quality, make initial observations, and communicate observations to the supervising  
208 radiologist; (iii) administer contrast media or other medications prescribed by the supervising radiologist;  
209 and (iv) perform, or assist the supervising radiologist to perform, any other procedure consistent with the  
210 guidelines adopted by the American College of Radiology, the American Society of Radiologic  
211 Technologists, and the American Registry of Radiologic Technologists.

212 "Respiratory care" means the practice of the allied health profession responsible for the direct and  
213 indirect services, including inhalation therapy and respiratory therapy, in the treatment, management,  
214 diagnostic testing, control, and care of patients with deficiencies and abnormalities associated with the  
215 cardiopulmonary system under qualified medical direction.

216 "Surgical assistant" means an individual who has met the requirements of the Board for licensure as  
217 a surgical assistant and who works under the direct supervision of a licensed doctor of medicine,  
218 osteopathy, or podiatry.

219 **§ 54.1-2901. Exceptions and exemptions generally.**

220 A. The provisions of this chapter shall not prevent or prohibit:

221 1. Any person entitled to practice his profession under any prior law on June 24, 1944, from  
222 continuing such practice within the scope of the definition of his particular school of practice;

223 2. Any person licensed to practice naturopathy prior to June 30, 1980, from continuing such practice  
224 in accordance with regulations promulgated by the Board;

225 3. Any licensed nurse practitioner from rendering care in accordance with the provisions of  
226 §§ 54.1-2957 and 54.1-2957.01, any nurse practitioner licensed by the Boards of Medicine and Nursing  
227 in the category of certified nurse midwife practicing pursuant to subsection H of § 54.1-2957, or any  
228 nurse practitioner licensed by the Boards of Medicine and Nursing in the category of clinical nurse  
229 specialist practicing pursuant to subsection J of § 54.1-2957 when such services are authorized by  
230 regulations promulgated jointly by the Boards of Medicine and Nursing;

231 4. 3. Any registered professional nurse, licensed nurse practitioner, graduate laboratory technician or  
232 other technical personnel who have been properly trained from rendering care or services within the  
233 scope of their usual professional activities which shall include the taking of blood, the giving of  
234 intravenous infusions and intravenous injections, and the insertion of tubes when performed under the  
235 orders of a person licensed to practice medicine or osteopathy, a nurse practitioner, or a physician  
236 assistant;

237 5. 4. Any dentist, pharmacist or optometrist from rendering care or services within the scope of his  
238 usual professional activities;

239 6. 5. Any practitioner licensed or certified by the Board from delegating to personnel supervised by  
240 him, such activities or functions as are nondiscretionary and do not require the exercise of professional  
241 judgment for their performance and which are usually or customarily delegated to such persons by  
242 practitioners of the healing arts, if such activities or functions are authorized by and performed for such  
243 practitioners of the healing arts and responsibility for such activities or functions is assumed by such

practitioners of the healing arts;

7. 6. The rendering of medical advice or information through telecommunications from a physician licensed to practice medicine in Virginia or an adjoining state, or from a licensed nurse practitioner, to emergency medical personnel acting in an emergency situation;

8. 7. The domestic administration of family remedies;

8. *Any person who sells vitamins and herbs from providing information about such products;*

9. The giving or use of massages, steam baths, dry heat rooms, infrared heat or ultraviolet lamps in public or private health clubs and spas;

10. The manufacture or sale of proprietary medicines in this Commonwealth by licensed pharmacists or druggists;

11. The advertising or sale of commercial appliances or remedies;

12. The fitting by nonitinerant persons or manufacturers of artificial eyes, limbs or other apparatus or appliances or the fitting of plaster cast counterparts of deformed portions of the body by a nonitinerant bracer or prosthetist for the purpose of having a three-dimensional record of the deformity, when such bracer or prosthetist has received a prescription from a licensed physician, licensed nurse practitioner, or licensed physician assistant directing the fitting of such casts and such activities are conducted in conformity with the laws of Virginia;

13. Any person from the rendering of first aid or medical assistance in an emergency in the absence of a person licensed to practice medicine or osteopathy under the provisions of this chapter;

14. The practice of the religious tenets of any church in the ministration to the sick and suffering by mental or spiritual means without the use of any drug or material remedy, whether gratuitously or for compensation;

15. Any legally qualified out-of-state or foreign practitioner from meeting in consultation with legally licensed practitioners in this Commonwealth;

16. Any practitioner of the healing arts licensed or certified and in good standing with the applicable regulatory agency in another state or Canada when that practitioner of the healing arts is in Virginia temporarily and such practitioner has been issued a temporary authorization by the Board from practicing medicine or the duties of the profession for which he is licensed or certified (i) in a summer camp or in conjunction with patients who are participating in recreational activities, (ii) while participating in continuing educational programs prescribed by the Board, or (iii) by rendering at any site any health care services within the limits of his license, voluntarily and without compensation, to any patient of any clinic which is organized in whole or in part for the delivery of health care services without charge as provided in § 54.1-106;

17. The performance of the duties of any active duty health care provider in active service in the army, navy, coast guard, marine corps, air force, or public health service of the United States at any public or private health care facility while such individual is so commissioned or serving and in accordance with his official military duties;

18. Any masseur, who publicly represents himself as such, from performing services within the scope of his usual professional activities and in conformance with state law;

19. Any person from performing services in the lawful conduct of his particular profession or business under state law;

20. Any person from rendering emergency care pursuant to the provisions of § 8.01-225;

21. Qualified emergency medical services personnel, when acting within the scope of their certification, and licensed health care practitioners, when acting within their scope of practice, from following Durable Do Not Resuscitate Orders issued in accordance with § 54.1-2987.1 and Board of Health regulations, or licensed health care practitioners from following any other written order of a physician not to resuscitate a patient in the event of cardiac or respiratory arrest;

22. Any commissioned or contract medical officer of the army, navy, coast guard or air force rendering services voluntarily and without compensation while deemed to be licensed pursuant to § 54.1-106;

23. Any provider of a chemical dependency treatment program who is certified as an "acupuncture detoxification specialist" by the National Acupuncture Detoxification Association or an equivalent certifying body, from administering auricular acupuncture treatment under the appropriate supervision of a National Acupuncture Detoxification Association certified licensed physician or licensed acupuncturist;

24. Any employee of any assisted living facility who is certified in cardiopulmonary resuscitation (CPR) acting in compliance with the patient's individualized service plan and with the written order of the attending physician not to resuscitate a patient in the event of cardiac or respiratory arrest;

25. Any person working as a health assistant under the direction of a licensed medical or osteopathic doctor within the Department of Corrections, the Department of Juvenile Justice or local correctional facilities;

26. Any employee of a school board, authorized by a prescriber and trained in the administration of

305 insulin and glucagon, when, upon the authorization of a prescriber and the written request of the parents  
306 as defined in § 22.1-1, assisting with the administration of insulin or administering glucagon to a  
307 student diagnosed as having diabetes and who requires insulin injections during the school day or for  
308 whom glucagon has been prescribed for the emergency treatment of hypoglycemia;

309 27. Any practitioner of the healing arts or other profession regulated by the Board from rendering  
310 free health care to an underserved population of Virginia who (i) does not regularly practice his  
311 profession in Virginia, (ii) holds a current valid license or certificate to practice his profession in another  
312 state, territory, district or possession of the United States, (iii) volunteers to provide free health care to  
313 an underserved area of the Commonwealth under the auspices of a publicly supported all volunteer,  
314 nonprofit organization that sponsors the provision of health care to populations of underserved people,  
315 (iv) files a copy of the license or certification issued in such other jurisdiction with the Board, (v)  
316 notifies the Board at least five business days prior to the voluntary provision of services of the dates and  
317 location of such service, and (vi) acknowledges, in writing, that such licensure exemption shall only be  
318 valid, in compliance with the Board's regulations, during the limited period that such free health care is  
319 made available through the volunteer, nonprofit organization on the dates and at the location filed with  
320 the Board. The Board may deny the right to practice in Virginia to any practitioner of the healing arts  
321 whose license or certificate has been previously suspended or revoked, who has been convicted of a  
322 felony or who is otherwise found to be in violation of applicable laws or regulations. However, the  
323 Board shall allow a practitioner of the healing arts who meets the above criteria to provide volunteer  
324 services without prior notice for a period of up to three days, provided the nonprofit organization  
325 verifies that the practitioner has a valid, unrestricted license in another state;

326 28. Any registered nurse, acting as an agent of the Department of Health, from obtaining specimens  
327 of sputum or other bodily fluid from persons in whom the diagnosis of active tuberculosis disease, as  
328 defined in § 32.1-49.1, is suspected and submitting orders for testing of such specimens to the Division  
329 of Consolidated Laboratories or other public health laboratories, designated by the State Health  
330 Commissioner, for the purpose of determining the presence or absence of tubercle bacilli as defined in  
331 § 32.1-49.1;

332 29. Any physician of medicine or osteopathy or nurse practitioner from delegating to a registered  
333 nurse under his supervision the screening and testing of children for elevated blood-lead levels when  
334 such testing is conducted (i) in accordance with a written protocol between the physician or nurse  
335 practitioner and the registered nurse and (ii) in compliance with the Board of Health's regulations  
336 promulgated pursuant to §§ 32.1-46.1 and 32.1-46.2. Any follow-up testing or treatment shall be  
337 conducted at the direction of a physician or nurse practitioner;

338 30. Any practitioner of one of the professions regulated by the Board of Medicine who is in good  
339 standing with the applicable regulatory agency in another state or Canada from engaging in the practice  
340 of that profession when the practitioner is in Virginia temporarily with an out-of-state athletic team or  
341 athlete for the duration of the athletic tournament, game, or event in which the team or athlete is  
342 competing;

343 31. Any person from performing state or federally funded health care tasks directed by the consumer,  
344 which are typically self-performed, for an individual who lives in a private residence and who, by  
345 reason of disability, is unable to perform such tasks but who is capable of directing the appropriate  
346 performance of such tasks;

347 32. Any practitioner of one of the professions regulated by the Board of Medicine who is in good  
348 standing with the applicable regulatory agency in another state from engaging in the practice of that  
349 profession in Virginia with a patient who is being transported to or from a Virginia hospital for care;

350 33. Any doctor of medicine or osteopathy, physician assistant, or nurse practitioner who would  
351 otherwise be subject to licensure by the Board who holds an active, unrestricted license in another state,  
352 the District of Columbia, or a United States territory or possession and who is in good standing with the  
353 applicable regulatory agency in that state, the District of Columbia, or that United States territory or  
354 possession who provides behavioral health services, as defined in § 37.2-100, from engaging in the  
355 practice of his profession and providing behavioral health services to a patient located in the  
356 Commonwealth in accordance with the standard of care when (i) such practice is for the purpose of  
357 providing continuity of care through the use of telemedicine services as defined in § 38.2-3418.16 and  
358 (ii) the practitioner has previously established a practitioner-patient relationship with the patient and has  
359 performed an in-person evaluation of the patient within the previous year. A practitioner who provides  
360 behavioral health services to a patient located in the Commonwealth through use of telemedicine  
361 services pursuant to this subdivision may provide such services for a period of no more than one year  
362 from the date on which the practitioner began providing such services to such patient;

363 34. Any employee of a program licensed by the Department of Behavioral Health and Developmental  
364 Services who is certified in cardiopulmonary resuscitation from acting in compliance with a program  
365 participant's valid written order not to resuscitate issued in accordance with § 54.1-2987.1 if such valid  
366 written order not to resuscitate is included in the program participant's individualized service plan; or

35. Any practitioner of a profession regulated by the Board of Medicine who is licensed in another state or the District of Columbia and who is in good standing with the applicable regulatory agency in that state or the District of Columbia from engaging in the practice of that profession in the Commonwealth with a patient located in the Commonwealth when (i) such practice is for the purpose of providing continuity of care through the use of telemedicine services as defined in § 38.2-3418.16 and (ii) the patient is a current patient of the practitioner with whom the practitioner has previously established a practitioner-patient relationship and the practitioner has performed an in-person examination of the patient within the previous 12 months. For the purposes of this subdivision, if a patient is (a) an enrollee of a health maintenance organization that contracts with a multispecialty group of practitioners, each of whom is licensed by the Board of Medicine, and (b) a current patient of at least one practitioner who is a member of the multispecialty group with whom such practitioner has previously established a practitioner-patient relationship and of whom such practitioner has performed an in-person examination within the previous 12 months, the patient shall be deemed to be a current patient of each practitioner in the multispecialty group with whom each such practitioner has established a practitioner-patient relationship.

B. Notwithstanding any provision of law or regulation to the contrary, military medical personnel, as defined in § 2.2-2001.4, while participating in a program established by the Department of Veterans Services pursuant to § 2.2-2001.4, may practice under the supervision of a licensed physician or podiatrist or the chief medical officer of an organization participating in such program, or his designee who is a licensee of the Board and supervising within his scope of practice.

**§ 54.1-2914. Sale of controlled substances and medical devices or appliances; requirements for vision care services.**

A. A practitioner of the healing arts shall not engage in selling controlled substances unless he is licensed to do so by the Board of Pharmacy. However, this prohibition shall not apply to a doctor of medicine, osteopathy or podiatry who administers controlled substances to his patients or provides controlled substances to his patient in a bona fide medical emergency or when pharmaceutical services are not available. Practitioners who sell or dispense controlled substances shall be subject to inspection by the Department of Health Professions to ensure compliance with Chapters 33 (§ 54.1-3300 et seq.) and 34 (§ 54.1-3400 et seq.) of this title and the Board of Pharmacy's regulations. This subsection shall not apply to physicians acting on behalf of the Virginia Department of Health or local health departments.

B. A practitioner of the healing arts who may lawfully sell medical appliances or devices shall not sell such appliances or devices to persons who are not his own patients and shall not sell such articles to his own patients either for his own convenience or for the purpose of supplementing his income. This subsection shall not apply to physicians acting on behalf of the Virginia Department of Health or local health departments.

C. A practitioner of the healing arts may, from within the practitioner's office, engage in selling or promoting the sale of eyeglasses and may dispense contact lenses. Only those practitioners of the healing arts who engage in the examination of eyes and prescribing of eyeglasses may engage in the sale or promotion of eyeglasses. Practitioners shall not employ any unlicensed person to fill prescriptions for eyeglasses within the practitioner's office except as provided in subdivision A 6 5 of § 54.1-2901. A practitioner may also own, in whole or in part, an optical dispensary located adjacent to or at a distance from his office.

D. Any practitioner of the healing arts engaging in the examination of eyes and prescribing of eyeglasses shall give the patient a copy of any prescription for eyeglasses and inform the patient of his right to have the prescription filled at the establishment of his choice. No practitioner who owns, in whole or in part, an establishment dispensing eyeglasses shall make any statement or take any action, directly or indirectly, that infringes on the patient's right to have a prescription filled at an establishment other than the one in which the practitioner has an ownership interest.

Disclosure of ownership interest by a practitioner as required by § 54.1-2964 or participation by the practitioner in contractual arrangements with third-party payors or purchasers of vision care services shall not constitute a violation of this subsection.

**§ 54.1-2956.15. Regulations relating to practice of naturopathy or naturopathic medicine.**

*The Board shall promulgate regulations governing the practice of naturopathic medicine. The regulations shall include, at a minimum, standards for (i) performing and ordering physical and laboratory examinations for diagnostic purposes, consistent with naturopathic medical education and training, including, but not limited to, phlebotomy, clinical laboratory tests, orificial examinations, physiological function tests, and diagnostic imaging studies; (ii) repair and care incidental to superficial lacerations and abrasions; (iii) removal of foreign bodies located in the superficial tissues; (iv) prescribing, dispensing, ordering, administering, or performing, as applicable, (a) food, extracts of food, nutraceuticals, vitamins, amino acids, minerals, enzymes, botanicals and their extracts, botanical*

428 medicines, homeopathic medicines, any dietary supplements or nonprescription drugs as defined by the  
429 Federal Food, Drug, and Cosmetic Act, 21 U.S.C. § 301 et seq.; (b) hot or cold hydrotherapy,  
430 naturopathic physical medicine, electromagnetic energy, colon hydrotherapy, and therapeutic exercise;  
431 or (c) devices, including therapeutic devices, barrier contraception, and durable medical equipment; (v)  
432 utilization of routes of administration that include oral, nasal, auricular, ocular, rectal, vaginal,  
433 transdermal, intradermal, subcutaneous, intravenous, and intramuscular consistent with naturopathic  
434 medical education and training; (vi) naturopathic childbirth attendance; and (vii) performing other  
435 therapies consistent with naturopathic medical education and training that are approved by the Board.

436 **§ 54.1-2956.16. Unlawful to practice naturopathic medicine without license; unlawful designation**  
437 **as naturopath; Board to regulate naturopathic doctors; scope of practice.**

438 A. It shall be unlawful for a person not holding a current and valid license from the Board to  
439 practice naturopathic medicine or to claim to be a naturopath, naturopathic doctor, naturopathic  
440 medical doctor, naturopathic physician, doctor of naturopathy, or doctor of naturopathic medicine or to  
441 assume the title doctor of naturopathic medicine, naturopathic doctor, naturopathic medical doctor,  
442 naturopathic physician, or to use the designations "N.D.," "ND," "N.M.D.," or "NMD" or any variation  
443 thereof.

444 In addition, it shall be unlawful for any person who is not licensed under this chapter, whose  
445 licensure has been suspended or revoked, or whose licensure has lapsed and has not been renewed to  
446 use in conjunction with his name the words "naturopath," "doctor of naturopathic medicine,"  
447 "naturopathic doctor," "naturopathic medical doctor," "naturopathic physician," or to otherwise by  
448 letters, words, representations, or insignias assert or imply that he is licensed to practice naturopathic  
449 medicine.

450 B. The Board shall prescribe by regulation the qualifications governing the licensure of naturopathic  
451 doctors. The regulations shall at a minimum require (i) graduation from a naturopathic medical  
452 education program in the United States accredited by the Council on Naturopathic Medical Education  
453 or an equivalent accrediting body for the naturopathic medical profession recognized by the U.S.  
454 Secretary of Education and the Board that offers graduate-level, full-time didactic and supervised  
455 clinical training leading to the degree of Doctor of Naturopathy or Doctor of Naturopathic Medicine  
456 and (ii) successful completion of a competency-based national naturopathic medicine licensing  
457 examination administered by the North American Board of Naturopathic Examiners, or an equivalent  
458 agency recognized by the Board. In lieu of graduation from an accredited naturopathic medical  
459 education program and the national naturopathic medicine licensing examination, the Board may  
460 require graduation from (a) an accredited institution of higher education or one that has received  
461 provisional accreditation from a regional accrediting body recognized by the U.S. Secretary of  
462 Education or (b) an accredited degree-granting institution of higher education that offers a full-time  
463 structured curriculum in basic sciences and supervised patient care consisting of a program of doctoral  
464 naturopathic medical education approved by the Board that requires the completion of 132 weeks or  
465 more of instruction within a period of not less than three years as a condition of graduation. The Board  
466 may license graduates of approved alternative naturopathic medical education programs upon evidence  
467 of successful completion of a Board-approved, competency-based state naturopathic medicine licensing  
468 examination or an equivalent Canadian provincial licensing examination for the practice of naturopathic  
469 medicine.

470 **§ 54.1-2956.17. Advisory Board on Naturopathy; composition; appointments.**

471 There is hereby established the Advisory Board on Naturopathy (the Advisory Board), which shall  
472 consist of five nonlegislative citizen members appointed by the Governor for four-year terms as follows:  
473 three members who shall be, at the time of appointment, naturopathic doctors licensed in the  
474 Commonwealth by the Board for not less than three years; one member who shall be licensed by the  
475 Board to practice either medicine, osteopathy, chiropractic, or podiatry; and one member who shall be  
476 a citizen of the Commonwealth at large. Appointments to fill vacancies, other than by expiration of a  
477 term, shall be for the unexpired terms. All members may be reappointed; however, no member shall  
478 serve more than two consecutive four-year terms. The remainder of any term to which a member is  
479 appointed to fill a vacancy shall not constitute a term in determining the member's eligibility for  
480 reappointment.

481 **§ 54.1-2956.18. Advisory Board on Naturopathy; powers.**

482 A. The Advisory Board shall, under the authority of the Board:

483 1. Recommend to the Board, for its promulgation into regulation, the criteria for licensure as a  
484 naturopathic doctors and the standards of professional conduct for holders of licenses.

485 2. Assess the qualifications of applicants for licensure and recommend licensure when applicants  
486 meet the required criteria. The recommendations of the Advisory Board relating to the licensure of  
487 applicants shall be presented to the Board, which shall then issue or deny licenses. Any applicant who  
488 is aggrieved by a denial of recommendation on licensure of the Advisory Board may appeal to the  
489 Board.



3. Recommend to the Board, for its promulgation into regulation, accrediting agencies, institutions of higher education, and competency-based examinations to be approved for naturopathic medical education and licensure.

4. Receive investigative reports of professional misconduct and unlawful acts and recommend sanctions when appropriate. Any recommendation of sanctions shall be presented to the Board, which may then impose sanctions or take such other action as may be warranted by law.

5. Recommend to the Board, for its promulgation into regulation, a formulary for use by naturopathic doctors.

6. Assist in such other matters dealing with naturopathic medicine as the Board may in its discretion direct.

B. Nothing in this chapter shall be construed to authorize the Advisory Board to advise the Board in matters pertaining to regulations governing the practice of medicine, osteopathy, chiropractic, or podiatry, or matters pertaining to doctors of medicine, osteopathy, chiropractic, or podiatry who are also licensed by the Board to practice naturopathy or naturopathic medicine.

#### **§ 54.1-2973.1. Practice of laser hair removal.**

The practice of laser hair removal shall be performed by a properly trained person licensed to practice medicine or osteopathic medicine or a physician assistant as authorized pursuant to § 54.1-2952 or a nurse practitioner as authorized pursuant to § 54.1-2957 or by a properly trained person under the direction and supervision of a licensed doctor of medicine or osteopathic medicine or a physician assistant as authorized pursuant to § 54.1-2952 or a nurse practitioner as authorized pursuant to § 54.1-2957 who may delegate such practice in accordance with subdivision A 6 5 of § 54.1-2901.

#### **§ 54.1-3401. Definitions.**

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

"Administer" means the direct application of a controlled substance, whether by injection, inhalation, ingestion, or any other means, to the body of a patient or research subject by (i) a practitioner or by his authorized agent and under his direction or (ii) the patient or research subject at the direction and in the presence of the practitioner.

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

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

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

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

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

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

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

"Board" means the Board of Pharmacy.

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

"Change of ownership" of an existing entity permitted, registered, or licensed by the Board means (i) the sale or transfer of all or substantially all of the assets of the entity or of any corporation that owns or controls the entity; (ii) the creation of a partnership by a sole proprietor, the dissolution of a partnership, or change in partnership composition; (iii) the acquisition or disposal of 50 percent or more of the outstanding shares of voting stock of a corporation owning the entity or of the parent corporation of a wholly owned subsidiary owning the entity, except that this shall not apply to any corporation the

551 voting stock of which is actively traded on any securities exchange or in any over-the-counter market;  
552 (iv) the merger of a corporation owning the entity or of the parent corporation of a wholly-owned  
553 subsidiary owning the entity with another business or corporation; or (v) the expiration or forfeiture of a  
554 corporation's charter.

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

557 "Compounding" means the combining of two or more ingredients to fabricate such ingredients into a  
558 single preparation and includes the mixing, assembling, packaging, or labeling of a drug or device (i) by  
559 a pharmacist, or within a permitted pharmacy, pursuant to a valid prescription issued for a medicinal or  
560 therapeutic purpose in the context of a bona fide practitioner-patient-pharmacist relationship, or in  
561 expectation of receiving a valid prescription based on observed historical patterns of prescribing and  
562 dispensing; (ii) by a practitioner of medicine, osteopathy, podiatry, dentistry, or veterinary medicine as  
563 an incident to his administering or dispensing, if authorized to dispense, a controlled substance in the  
564 course of his professional practice; or (iii) for the purpose of, or as incident to, research, teaching, or  
565 chemical analysis and not for sale or for dispensing. The mixing, diluting, or reconstituting of a  
566 manufacturer's product drugs for the purpose of administration to a patient, when performed by a  
567 practitioner of medicine or osteopathy licensed under Chapter 29 (§ 54.1-2900 et seq.), a person  
568 supervised by such practitioner pursuant to subdivision A 6 5 or 19 of § 54.1-2901, or a person  
569 supervised by such practitioner or a licensed nurse practitioner or physician assistant pursuant to  
570 subdivision A 4 3 of § 54.1-2901 shall not be considered compounding.

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

576 "Controlled substance analog" means a substance the chemical structure of which is substantially  
577 similar to the chemical structure of a controlled substance in Schedule I or II and either (i) which has a  
578 stimulant, depressant, or hallucinogenic effect on the central nervous system that is substantially similar  
579 to or greater than the stimulant, depressant, or hallucinogenic effect on the central nervous system of a  
580 controlled substance in Schedule I or II or (ii) with respect to a particular person, which such person  
581 represents or intends to have a stimulant, depressant, or hallucinogenic effect on the central nervous  
582 system that is substantially similar to or greater than the stimulant, depressant, or hallucinogenic effect  
583 on the central nervous system of a controlled substance in Schedule I or II. "Controlled substance  
584 analog" does not include (a) any substance for which there is an approved new drug application as  
585 defined under § 505 of the federal Food, Drug, and Cosmetic Act (21 U.S.C. § 355) or that is generally  
586 recognized as safe and effective pursuant to §§ 501, 502, and 503 of the federal Food, Drug, and  
587 Cosmetic Act (21 U.S.C. §§ 351, 352, and 353) and 21 C.F.R. Part 330; (b) with respect to a particular  
588 person, any substance for which an exemption is in effect for investigational use for that person under  
589 § 505 of the federal Food, Drug, and Cosmetic Act to the extent that the conduct with respect to that  
590 substance is pursuant to such exemption; or (c) any substance to the extent not intended for human  
591 consumption before such an exemption takes effect with respect to that substance.

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

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

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

603 "Dialysis care technician" or "dialysis patient care technician" means an individual who is certified  
604 by an organization approved by the Board of Health Professions pursuant to Chapter 27.01 (§  
605 54.1-2729.1 et seq.) and who, under the supervision of a licensed physician, nurse practitioner, physician  
606 assistant, or a registered nurse, assists in the care of patients undergoing renal dialysis treatments in a  
607 Medicare-certified renal dialysis facility.

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

612 "Dispense" means to deliver a drug to an ultimate user or research subject by or pursuant to the

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

"Dispenser" means a practitioner who dispenses.

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

"Distributor" means a person who distributes.

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

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

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

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

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

"Immediate precursor" means a substance which the Board of Pharmacy has found to be and by regulation designates as being the principal compound commonly used or produced primarily for use, and which is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance, the control of which is necessary to prevent, curtail, or limit manufacture.

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

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

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

"Manufacture" means the production, preparation, propagation, conversion, or processing of any item regulated by this chapter, either directly or indirectly by extraction from substances of natural origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis, and includes any packaging or repackaging of the substance or labeling or relabeling of its container. This term does not include compounding.

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

"Marijuana" means any part of a plant of the genus Cannabis whether growing or not, its seeds, or its resin; and every compound, manufacture, salt, derivative, mixture, or preparation of such plant, its seeds, its resin, or any extract containing one or more cannabinoids. Marijuana does not include the mature stalks of such plant, fiber produced from such stalk, or oil or cake made from the seeds of such plant, unless such stalks, fiber, oil, or cake is combined with other parts of plants of the genus Cannabis. Marijuana does not include (i) industrial hemp, as defined in § 3.2-4112, that is possessed by a person registered pursuant to subsection A of § 3.2-4115 or his agent, (ii) industrial hemp, as defined in § 3.2-4112, that is possessed by a person who holds a hemp producer license issued by the U.S. Department of Agriculture pursuant to 7 C.F.R. Part 990, or (iii) a hemp product, as defined in § 3.2-4112, containing a tetrahydrocannabinol concentration of no greater than 0.3 percent that is derived from industrial hemp, as defined in § 3.2-4112, that is grown, dealt, or processed in compliance with state or federal law.

"Medical equipment supplier" means any person, as defined in § 1-230, engaged in the delivery to the ultimate consumer, pursuant to the lawful order of a practitioner, of hypodermic syringes and needles, medicinal oxygen, Schedule VI controlled devices, those Schedule VI controlled substances with

674 no medicinal properties that are used for the operation and cleaning of medical equipment, solutions for  
675 peritoneal dialysis, and sterile water or saline for irrigation.

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

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

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

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

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

707 "Opiate" means any substance having an addiction-forming or addiction-sustaining liability similar to  
708 morphine or being capable of conversion into a drug having such addiction-forming or  
709 addiction-sustaining liability. It does not include, unless specifically designated as controlled under  
710 Article 4 (§ 54.1-3437 et seq.), the dextrorotatory isomer of 3-methoxy-n-methylmorphinan and its salts  
711 (dextromethorphan). It does include its racemic and levorotatory forms.

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

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

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

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

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

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

728 "Practitioner" means a physician, dentist, licensed nurse practitioner pursuant to § 54.1-2957.01,  
729 licensed physician assistant pursuant to § 54.1-2952.1, pharmacist pursuant to § 54.1-3300, TPA-certified  
730 optometrist pursuant to Article 5 (§ 54.1-3222 et seq.) of Chapter 32, veterinarian, scientific investigator,  
731 or other person licensed, registered, or otherwise permitted to distribute, dispense, prescribe and  
732 administer, or conduct research with respect to a controlled substance in the course of professional  
733 practice or research in the Commonwealth.

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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