SUBSTITUTE SENATE BILL 5112

State of Washington 69th Legislature 2025 Regular Session

By Senate Health & Long-Term Care (originally sponsored by Senators Bateman, Harris, Shewmake, Trudeau, Conway, Nobles, Riccelli, C. Wilson, Robinson, and Hasegawa)

READ FIRST TIME 02/21/25.

- AN ACT Relating to establishing a prescribing psychologist certification in Washington state; amending RCW 18.83.010, 18.83.035, 18.83.050, 18.83.050, 18.83.080, 18.83.080, 18.83.090, 18.79.260, 18.79.260, 69.50.101, 69.41.010, and 69.41.030; reenacting and amending RCW 18.64.011; adding new sections to chapter 18.83 RCW; creating a new section; providing effective dates; and providing expiration dates.
- 8 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF WASHINGTON:
- 9 NEW SECTION. Sec. 1. The legislature finds that:
- 10 (1) One in five adults in Washington experience a mental illness 11 each year;
- 12 (2) More than half of people with a mental health condition did 13 not receive any treatment in the last year;
- 14 (3) The association of American medical colleges forecasts a 15 long-term and persistent shortage of doctors, including specialty 16 providers, in a 2021 report;
- 17 (4) Other states, the department of defense, and the Indian 18 health service have all successfully credentialed prescribing 19 psychologists to safely prescribe psychotropic medications; and

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- 1 (5) Washington residents will benefit from increased access and 2 decreased costs by creating a new credential for prescribing 3 psychology.
- 4 **Sec. 2.** RCW 18.83.010 and 1994 c 35 s 1 are each amended to read 5 as follows:
 - ((When used in this chapter:

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- 7 (1) The "practice)) The definitions in this section apply
 8 throughout this chapter unless the context clearly requires
 9 otherwise.
 - (1) "Board" means the examining board of psychology.
- 11 (2) "Clinical experience" means a period of supervised clinical 12 training and practice conducted as part of a training program in 13 which clinical diagnoses and interventions are learned.
 - (3) "Department" means the department of health.
 - (4) "Practice of psychology" means the observation, evaluation, interpretation, and modification of human behavior by the application of psychological principles, methods, and procedures for the purposes of preventing or eliminating symptomatic or maladaptive behavior and promoting mental and behavioral health. It includes, but is not limited to, providing the following services to individuals, families, groups, organizations, and the public, whether or not payment is received for services rendered:
- 23 (a) Psychological measurement, assessment, and evaluation by 24 means of psychological, neuropsychological, and psychoeducational 25 testing;
 - (b) Diagnosis and treatment of mental, emotional, and behavioral disorders, and psychological aspects of illness, injury, and disability; and
 - (c) Counseling and guidance, psychotherapeutic techniques, remediation, health promotion, and consultation within the context of established psychological principles and theories.

This definition does not include the teaching of principles of psychology for accredited educational institutions, or the conduct of research in problems of human or animal behavior.

Nothing in this definition shall be construed as permitting the administration or prescribing of drugs <u>except as allowed in this chapter</u> or in any way infringing upon the practice of medicine and surgery as defined in chapter 18.71 RCW.

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- 1 $((\frac{2}{2}))$ (5) "Prescribing psychologist" means a person who holds an active license to engage in the practice of psychology under this chapter and an active certificate as a prescribing psychologist under 3 section 3 of this act, and is limited by the restrictions under 4 section 4 of this act. 5
- 6 (6) "Prescription" has the same meaning as defined in RCW 7 18.64.011.
 - (7) "Prescriptive authority" means the authority of a prescribing psychologist to prescribe, administer, discontinue, and distribute psychotropic medications recognized or customarily used in the diagnosis, treatment, and management of individuals with psychiatric, mental, cognitive, nervous, emotional, developmental, or behavioral disorders identified in the most recent edition of a widely accepted classification system of mental disorders, as identified by the board. The term includes ordering and obtaining necessary laboratory tests, procedures, imaging, and diagnostic examinations.
 - (8) "Psychotropic medication" means substances recognized as drugs, including controlled substances used to treat mental illnesses, in the official United States pharmacopoeia, official homeopathic pharmacopeia of the United States, official national formulary, or any respective supplement to those publications.
- 22 (9) "Secretary" means the secretary of health.

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- 23 (((3) "Board" means the examining board of psychology.
- 24 (4) "Department" means the department of health.))
- 25 <u>NEW SECTION.</u> **Sec. 3.** A new section is added to chapter 18.83 RCW to read as follows: 26
- 27 (1) A psychologist licensed under this chapter may apply for 28 certification as a prescribing psychologist.
 - The board shall certify an applicant as a prescribing psychologist if the applicant demonstrates to the board, by official transcript or other official evidence satisfactory to the board, that the applicant:
 - (a) Holds a current license as a psychologist under this chapter;
- (b) Holds a doctorate degree obtained from an integrated program 34 of graduate study in psychology, as defined in rule by the board; 35
- Has successfully completed a designated master's degree 36 program in clinical psychopharmacology that meets the criteria 37 38 established in subsection (3) of this section;

p. 3 SSB 5112 (d) Has successfully completed at least 80 hours of supervised clinical experience in physical assessment, including physical examinations with instruction in the proper use of instruments used in physical examination, supervised by a medical provider licensed to conduct independent physical assessments;

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- (e) Has successfully completed a clinical prescribing fellowship under the supervision of a qualified supervisor including clinical experience sufficient to attain competency in the psychopharmacological treatment of a diverse patient population, to be comprised of no less than 500 hours and 100 individual patients. Qualified supervisors are licensed health care providers with specialized training and experience in the management of psychotropic medication who are licensed in Washington state or pursuant to a substantially equivalent licensing provision of the law of another state, as established by the board, including physicians, physician assistants, osteopathic physicians, psychiatric nurse practitioners, or prescribing psychologists; and
- (f) Has passed an examination relevant to establishing competence for prescribing as developed by a nationally recognized organization and approved by the board.
- Α designated master's degree program in clinical psychopharmacology must be an accredited program within a regionally accredited institution of higher education approved by the United States department of education, satisfy requirements for designation established by the board, and be substantially equivalent to the training required of advanced practice psychiatric nurses. The board may use the standards of an association which publishes relevant education and training program standards such as the American psychological association. The didactic portion of the program shall include at least two years of education, a minimum of 400 contact hours, or the equivalent thereof, and include sufficient biomedical education to ensure the necessary knowledge and skills to prescribe psychotropic medications in a safe and effective manner, including but not limited to:
- 35 (a) Science prerequisites, including human anatomy and human 36 physiology, and a course in biology;
- 37 (b) Basic science, including human anatomy, human physiology, 38 biochemistry, and genetics;
- 39 (c) Functional neuroscience, including neuroanatomy, 40 neurophysiology, and neurochemistry;

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(d) Physical examinations, including the measurement and interpretation of vital signs and neurological, cardiovascular, respiratory, abdominal, eye, ear, nose, throat, gastrointestinal, genitourinary, integumentary, allergic and immunologic, and musculoskeletal examinations;

- (e) Interpretation of laboratory tests, including therapeutic drug monitoring, blood and urine tests, radiology, electrocardiogram, brain electrophysiology, neuroimaging techniques, and applied genetics;
- 10 (f) Pathological basis of disease, including pathophysiology of 11 common clinical cardiovascular, respiratory, gastrointestinal, 12 hepatic, neurological, and endocrine conditions;
 - (g) Clinical medicine, including clinical manifestations, differential diagnosis, laboratory or radiological evaluation of commonly encountered medical conditions such as patients with complex medical needs and comorbidities, and medical emergencies and their management;
 - (h) Clinical neurotherapeutics, including electrophysiology, electroconvulsive therapy, and noninvasive interventions, such as transcranial magnetic stimulation, neurofeedback, and biofeedback;
 - (i) Systems of care, including coordination of care with other medical specialties, consultations and referrals, and coordination and consultation in long-term care;
 - (j) Pharmacology, including pharmacokinetics and drug delivery systems, pharmacodynamics, neuropharmacology, toxicology, and mechanisms of medication interactions;
 - (k) Clinical pharmacology, including major drug classes and nutritional supplements;
 - (1) Psychopharmacology, including sedatives and hypnotics, antidepressants, antipsychotics, mood stabilizers, anxiolytics, stimulants, medications for substance use disorders, medications for drug adverse effects, pediatric psychopharmacology, geriatric psychopharmacology, medications for cognitive impairment and polypharmacy, issues of diversity and cultural competence in pharmacological practice, clinical decision making and standard practice guidelines, and guidelines for prescribing controlled substances;
- 38 (m) Psychopharmacology research, including phases of drug 39 development, clinical trials in psychiatry, and critical evaluation 40 of evidence; and

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(n) Professional, ethical, and legal issues, including conflicts of interest and relationships with the industry, scope of practice issues, diversity and equity issues related to treatment access and adherence, and documentation issues, including nomenclature, abbreviations, and prescription writing.

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- (4) The board may waive certain requirements for applicants who have obtained relevant training and experience, including psychologists who are dually licensed as physicians, nurse practitioners, or other health professionals with comparable prescriptive authority in Washington.
- (5) The board may offer a certificate in prescriptive authority 11 12 by endorsement to an applicant who has a current and unrestricted license to practice psychology and either a current and unrestricted 13 certificate in prescriptive authority from another state, or training 14 from the United States department of defense demonstration project or 15 16 other similar program developed and operated by any branch of the 17 armed forces that imposes substantially equivalent educational and training requirements as those contained in this chapter and required 18 19 by the board. Upon payment of the required fees, compliance with relevant statutory provisions, and the approval of the application, 20 21 the applicant may be certified by endorsement pursuant to this chapter. The board may consider an applicant's experience in 22 prescribing in another state as meeting a portion of the requirements 23 necessary to obtain provisional certification or certification under 24 25 this chapter, but also shall require additional education and supervision if the board deems it necessary to meet the education and 26 training requirements imposed by this chapter. 27
- 28 (6) A certificate issued under this section may be renewed in accordance with RCW 18.83.090.
- 30 (7) The secretary shall establish the administrative procedures, 31 administrative requirements, and fees for the certificate as provided 32 in RCW 43.70.250 and 43.70.280.
- NEW SECTION. Sec. 4. A new section is added to chapter 18.83 RCW to read as follows:
- 35 (1) The prescriptive authority of a prescribing psychologist is 36 limited to prescribing, administering, discontinuing, and 37 distributing psychotropic medications recognized or customarily used 38 in the diagnosis, treatment, and management of individuals with 39 psychiatric, mental, cognitive, nervous, emotional, developmental, or

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behavioral disorders identified in the most recent edition of a widely accepted classification system of mental disorders, as identified by the board. A prescribing psychologist may order and obtain necessary laboratory tests, procedures, imaging, and diagnostic examinations necessary to exercise this prescriptive authority. A psychologist who is not a prescribing psychologist may not exercise this prescriptive authority.

- (2) When prescribing psychotropic medication for a patient, a prescribing psychologist must maintain an ongoing collaborative relationship with a health care practitioner who oversees the patient's general medical care to ensure that necessary medical examinations are conducted and that the psychotropic medication is appropriate for the patient's medical condition. The prescribing psychologist and the health care practitioner must coordinate the patient's ongoing care. The prescribing psychologist may not issue a new prescription for a psychotropic medication without agreement, in a form and manner determined by the board in rule by January 1, 2027, from the health care practitioner who oversees the patient's general medical care.
- (3) A prescribing psychologist may not prescribe opioid medications except for medications appropriate for treatment of an opioid use disorder which are prescribed for the purpose of treatment of such a disorder. A prescribing psychologist may not prescribe any antipsychotic medication or mood stabilizer.
- (4) Each prescription issued by a prescribing psychologist must comply with applicable state and federal laws and regulations and be identified as written by the prescribing psychologist in a manner determined by the board.
- (5) A prescribing psychologist shall ensure that a record of all prescriptions made by the prescribing psychologist is maintained in the patient's record.
 - (6) A prescribing psychologist may not delegate the authority to prescribe drugs or controlled substances to any other person.
- 34 (7) A prescribing psychologist may not prescribe to any 35 individual under the age of 25 or over the age of 65.
- **Sec. 5.** RCW 18.83.035 and 2022 c 240 s 10 are each amended to read as follows:
- There is created the examining board of psychology which shall examine the qualifications of applicants for licensing. The board

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1 shall consist of nine psychologists, one expert on psychiatric prescribing, and two public members, all appointed by the governor. 2 3 The public members shall not be and have never been psychologists or in training to be psychologists; they may not have any household 4 member who is a psychologist or in training to be a psychologist; 5 6 they may not participate or ever have participated in a commercial or professional field related to psychology, nor have a household member 7 who has so participated; and they may not have had within two years 8 before appointment a substantial financial interest in a person 9 regulated by the board. Each psychologist member of the board shall 10 11 have actively practiced psychology in the state of Washington for at 12 least three years immediately preceding appointment and ((who is)) be licensed under this chapter. The member who is an expert on 13 psychiatric prescribing must have specialized training and experience 14 15 in the management of psychotropic medication and be a prescribing psychologist, physician, osteopathic physician with special knowledge 16 of psychopharmacology, psychiatric nurse practitioner, or pharmacist 17 with expertise in psychopharmacology. Board members 18 shall 19 appointed for a term of five years, except that the terms of the existing appointees shall be adjusted by the governor so that no more 20 21 than two members' terms expire each year with all subsequent 22 appointments for a five-year term. Upon the death, resignation, or 23 removal of a member, the governor shall appoint a successor to serve for the unexpired term. The board shall elect one of its members to 24 25 serve as chairperson.

- 26 **Sec. 6.** RCW 18.83.050 and 2004 c 262 s 8 are each amended to 27 read as follows:
- 28 (1) The board shall adopt such rules as it deems necessary to 29 carry out its functions.

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- (2) The board shall examine the qualifications of applicants for licensing under this chapter, to determine which applicants are eligible for licensing under this chapter and shall forward to the secretary the names of applicants so eligible.
- (3) The board shall administer examinations to qualified applicants on at least an annual basis. The board shall determine the subject matter and scope of the examination, except as provided in RCW 18.83.170. The board may allow applicants to take the examination upon the granting of their doctoral degree before completion of their internship for supervised experience.

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(4) The board shall:

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- 2 (a) Develop and implement procedures for reviewing the education 3 and training credentials of applicants for certification as a 4 prescribing psychologist;
 - (b) Adopt rules, in consultation with the Washington medical commission, to establish standards for the certification of prescribing psychologists in accordance with section 3 of this act and for their exercise of prescriptive authority under this chapter; and
 - (c) Adopt rules for denying, modifying, suspending, or revoking the certification of a prescribing psychologist. The board may require remediation of any deficiencies in the training or practice pattern of the prescribing psychologist when, in the judgment of the board, such deficiencies could reasonably be expected to jeopardize the health, safety, or welfare of the public.
- 16 <u>(5) The board shall maintain a current list of each prescribing</u> 17 <u>psychologist's license and certification numbers.</u>
 - (6) The board shall keep a complete record of its own proceedings, of the questions given in examinations, of the names and qualifications of all applicants, and the names and addresses of all licensed psychologists. The examination paper of such applicant shall be kept on file for a period of at least one year after examination.
- 23 (((+5))) (7) The board shall, by rule, adopt a code of ethics for psychologists which is designed to protect the public interest.
- (((6))) <u>(8)</u> The board may require that persons licensed under this chapter as psychologists obtain and maintain professional liability insurance in amounts determined by the board to be practicable and reasonably available.
- 29 **Sec. 7.** RCW 18.83.050 and 2024 c 371 s 3 are each amended to 30 read as follows:
- 31 (1) The board shall adopt such rules as it deems necessary to 32 carry out its functions.
 - (2) The board shall examine the qualifications of applicants for licensing under this chapter, to determine which applicants are eligible for licensing under this chapter and shall forward to the secretary the names of applicants so eligible.
- 37 (3) The board shall administer examinations to qualified 38 applicants on at least an annual basis. The board shall determine the 39 subject matter and scope of the examination, except as provided in

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1 RCW 18.83.170. The board may allow applicants to take the examination 2 upon the granting of their doctoral degree before completion of their 3 internship for supervised experience.

(4) The board shall:

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- (a) Develop and implement procedures for reviewing the education and training credentials of applicants for certification as a prescribing psychologist;
- (b) Adopt rules, in consultation with the Washington medical commission, to establish standards for the certification of prescribing psychologists in accordance with section 3 of this act and for their exercise of prescriptive authority under this chapter; and
- (c) Adopt rules for denying, modifying, suspending, or revoking the certification of a prescribing psychologist. The board may require remediation of any deficiencies in the training or practice pattern of the prescribing psychologist when, in the judgment of the board, such deficiencies could reasonably be expected to jeopardize the health, safety, or welfare of the public.
- 19 <u>(5) The board shall maintain a current list of each prescribing</u> 20 <u>psychologist's license and certification numbers.</u>
 - (6) The board shall keep a complete record of its own proceedings, of the questions given in examinations, of the names and qualifications of all applicants, and the names and addresses of all licensed psychologists and licensed psychological associates. The examination paper of such applicant shall be kept on file for a period of at least one year after examination.
 - $((\frac{5}{1}))$ The board shall, by rule, adopt a code of ethics for psychologists and licensed psychological associates which is designed to protect the public interest.
- 30 (((6))) <u>(8)</u> The board may require that persons licensed under 31 this chapter as psychologists or licensed psychological associates 32 obtain and maintain professional liability insurance in amounts 33 determined by the board to be practicable and reasonably available.
- 34 **Sec. 8.** RCW 18.83.080 and 1996 c 191 s 66 are each amended to read as follows:
- The board shall forward to the secretary the name of each applicant entitled to a license <u>or certificate</u> under this chapter. The secretary shall promptly issue to such applicant a license authorizing such applicant to use the title "psychologist" $((\cdot))$ or a

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- 1 certificate authorizing the applicant to use the title "prescribing
- 2 <u>psychologist."</u> Each licensed psychologist shall keep his or her
- 3 license and, if applicable, prescribing psychologist certificate
- 4 displayed in a conspicuous place in his or her principal place of
- 5 business.
- 6 **Sec. 9.** RCW 18.83.080 and 2024 c 371 s 4 are each amended to read as follows:
- 8 The board shall forward to the secretary the name of each
- 9 applicant entitled to a license or certificate under this chapter.
- 10 The secretary shall promptly issue to such applicant a license
- 11 authorizing such applicant to use the title "psychologist" or
- "licensed psychological associate((\cdot))" or a certificate authorizing
- 13 the applicant to use the title "prescribing psychologist." Each
- 14 licensed psychologist or licensed psychological associate shall keep
- 15 his or her license and, if applicable, prescribing psychologist
- 16 <u>certificate</u> displayed in a conspicuous place in his or her principal
- 17 place of business.

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- 18 **Sec. 10.** RCW 18.83.090 and 2009 c 492 s 6 are each amended to 19 read as follows:
- 20 (1) The board shall establish rules governing mandatory 21 continuing education requirements which shall be met by any 22 psychologist applying for a license or prescribing psychologist 23 certificate renewal.
 - (2) The office of crime victims advocacy shall supply the board with information on methods of recognizing victims of human trafficking, what services are available for these victims, and where to report potential trafficking situations. The information supplied must be culturally sensitive and must include information relating to minor victims. The board shall disseminate this information to licensees by: Providing the information on the board's website; including the information in newsletters; holding trainings at meetings attended by organization members; or ((through another)) using other distribution methods determined by the board. The board shall report to the office of crime victims advocacy on the method or methods it uses to distribute information under this subsection.
- 36 (3) Administrative procedures, administrative requirements, and 37 fees for renewal and reissue of licenses <u>and certificates</u> shall be 38 established as provided in RCW 43.70.250 and 43.70.280.

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(4) (a) The board shall establish rules for the renewal of a prescribing psychologist certificate issued under section 3 of this act at the time of the renewal of the psychologist's license to practice psychology.

- 5 (b) Each applicant for renewal of a prescribing psychologist
 6 certificate shall present satisfactory evidence to the board
 7 demonstrating the completion of continuing education instruction
 8 relevant to prescriptive authority during the previous three-year
 9 renewal period.
- Sec. 11. RCW 18.64.011 and 2024 c 121 s 30 are each reenacted and amended to read as follows:

The definitions in this section apply throughout this chapter unless the context clearly requires otherwise.

- (1) "Administer" means the direct application of a drug or device, whether by injection, inhalation, ingestion, or any other means, to the body of a patient or research subject.
- (2) "Business licensing system" means the mechanism established by chapter 19.02 RCW by which business licenses, endorsed for individual state-issued licenses, are issued and renewed utilizing a business license application and a business license expiration date common to each renewable license endorsement.
- (3) "Chart order" means a lawful order for a drug or device entered on the chart or medical record of an inpatient or resident of an institutional facility by a practitioner or his or her designated agent.
- (4) "Closed door long-term care pharmacy" means a pharmacy that provides pharmaceutical care to a defined and exclusive group of patients who have access to the services of the pharmacy because they are treated by or have an affiliation with a long-term care facility or hospice program, and that is not a retailer of goods to the general public.
 - (5) "Commission" means the pharmacy quality assurance commission.
- (6) "Compounding" means the act of combining two or more ingredients in the preparation of a prescription. Reconstitution and mixing of (a) sterile products according to federal food and drug administration-approved labeling does not constitute compounding if prepared pursuant to a prescription and administered immediately or in accordance with package labeling, and (b) nonsterile products according to federal food and drug administration-approved labeling

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1 does not constitute compounding if prepared pursuant to a 2 prescription.

- (7) "Controlled substance" means a drug or substance, or an immediate precursor of such drug or substance, so designated under or pursuant to the provisions of chapter 69.50 RCW.
- (8) "Deliver" or "delivery" means the actual, constructive, or attempted transfer from one person to another of a drug or device, whether or not there is an agency relationship.
 - (9) "Department" means the department of health.

- (10) "Device" means instruments, apparatus, and contrivances, including their components, parts, and accessories, intended (a) for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in human beings or other animals, or (b) to affect the structure or any function of the body of human beings or other animals.
- (11) "Directed plan of correction" means a plan devised by the commission that includes specific actions that must be taken to correct identified unresolved deficiencies with time frames to complete them.
- (12) "Dispense" means the interpretation of a prescription or order for a drug, biological, or device and, pursuant to that prescription or order, the proper selection, measuring, compounding, labeling, or packaging necessary to prepare that prescription or order for delivery.
- (13) "Distribute" means the delivery of a drug or device other than by administering or dispensing.
- (14) "Drug" and "devices" do not include surgical or dental instruments or laboratory materials, gas and oxygen, therapy equipment, X-ray apparatus or therapeutic equipment, their component parts or accessories, or equipment, instruments, apparatus, or contrivances used to render such articles effective in medical, surgical, or dental treatment, or for use or consumption in or for mechanical, industrial, manufacturing, or scientific applications or purposes. "Drug" also does not include any article or mixture covered by the Washington pesticide control act (chapter 15.58 RCW), as enacted or hereafter amended, nor medicated feed intended for and used exclusively as a feed for animals other than human beings.
 - (15) "Drugs" means:

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1 (a) Articles recognized in the official United States 2 pharmacopoeia or the official homeopathic pharmacopoeia of the United 3 States;

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- (b) Substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in human beings or other animals;
- (c) Substances (other than food) intended to affect the structure or any function of the body of human beings or other animals; or
- (d) Substances intended for use as a component of any substances specified in (a), (b), or (c) of this subsection, but not including devices or their component parts or accessories.
- (16) "Health care entity" means an organization that provides health care services in a setting that is not otherwise licensed by the state to acquire or possess legend drugs. Health care entity includes a freestanding outpatient surgery center, a residential treatment facility, and a freestanding cardiac care center. "Health care entity" does not include an individual practitioner's office or a multipractitioner clinic, regardless of ownership, unless the owner elects licensure as a health care entity. "Health care entity" also does not include an individual practitioner's office or multipractitioner clinic identified by a hospital on a pharmacy application or renewal pursuant to RCW 18.64.043.
- (17) "Hospice program" means a hospice program certified or paid by medicare under Title XVIII of the federal social security act, or a hospice program licensed under chapter 70.127 RCW.
- (18) "Immediate jeopardy" means a situation in which a licensee's noncompliance with one or more statutory or regulatory requirements has placed the health and safety of individuals or animals at risk for serious injury, serious harm, serious impairment, or death.
- 30 (19) "Institutional facility" means any organization whose 31 primary purpose is to provide a physical environment for patients to 32 obtain health care services including, but not limited to, services 33 in a hospital, long-term care facility, hospice program, mental 34 health facility, drug abuse treatment center, residential 35 habilitation center, or a local, state, or federal correction 36 facility.
- 37 (20) "Labeling" means the process of preparing and affixing a 38 label to any drug or device container. The label must include all 39 information required by current federal and state law and pharmacy 40 rules.

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(21) "Legend drugs" means any drugs which are required by any applicable federal or state law or regulation to be dispensed on prescription only or are restricted to use by practitioners only.

- (22) "License," "licensing," and "licensure" shall be deemed equivalent to the terms "approval," "credential," "certificate," "certification," "permit," and "registration" and an "exemption" issued under chapter 69.50 RCW.
- (23) "Long-term care facility" means a nursing home licensed under chapter 18.51 RCW, an assisted living facility licensed under chapter 18.20 RCW, or an adult family home licensed under chapter 70.128 RCW.
- "Manufacture" means the production, preparation, propagation, compounding, or processing of a drug or other substance or device or the packaging or repackaging of such substance or device, or the labeling or relabeling of the commercial container of such substance or device, but does not include the activities of a practitioner who, as an incident to his or her administration or dispensing such substance or device in the course of his or her professional practice, personally prepares, compounds, packages, or labels such substance or device. "Manufacture" includes the distribution of a licensed pharmacy compounded drug product to other state licensed persons or commercial entities for subsequent resale or distribution, unless a specific product item has approval of the commission. The term does not include:
- (a) The activities of a licensed pharmacy that compounds a product on or in anticipation of an order of a licensed practitioner for use in the course of their professional practice to administer to patients, either personally or under their direct supervision;
- (b) The practice of a licensed pharmacy when repackaging commercially available medication in small, reasonable quantities for a practitioner legally authorized to prescribe the medication for office use only;
- (c) The distribution of a drug product that has been compounded by a licensed pharmacy to other appropriately licensed entities under common ownership or control of the facility in which the compounding takes place; or
- (d) The delivery of finished and appropriately labeled compounded products dispensed pursuant to a valid prescription to alternate delivery locations, other than the patient's residence, when

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requested by the patient, or the prescriber to administer to the patient, or to another licensed pharmacy to dispense to the patient.

- (25) "Manufacturer" means a person, corporation, or other entity engaged in the manufacture of drugs or devices.
- (26) "Nonlegend" or "nonprescription" drugs means any drugs which may be lawfully sold without a prescription.
- (27) "Person" means an individual, corporation, government, governmental subdivision or agency, business trust, estate, trust, partnership or association, or any other legal entity.
- 10 (28) "Pharmacist" means a person duly licensed by the commission 11 to engage in the practice of pharmacy.
 - (29) "Pharmacy" means every place properly licensed by the commission where the practice of pharmacy is conducted.
 - (30) "Plan of correction" means a proposal devised by the applicant or licensee that includes specific actions that must be taken to correct identified unresolved deficiencies with the time frames to complete them.
 - (31) "Poison" does not include any article or mixture covered by the Washington pesticide control act (chapter 15.58 RCW), as enacted or hereafter amended.
 - (32) "Practice of pharmacy" includes the practice of and responsibility for: Interpreting prescription orders; the compounding, dispensing, labeling, administering, and distributing of drugs and devices; the monitoring of drug therapy and use; the initiating or modifying of drug therapy in accordance with written guidelines or protocols previously established and approved for his or her practice by a practitioner authorized to prescribe drugs; the participating in drug utilization reviews and drug product selection; the proper and safe storing and distributing of drugs and devices and maintenance of proper records thereof; the providing of information on legend drugs which may include, but is not limited to, the advising of therapeutic values, hazards, and the uses of drugs and devices.
 - (33) "Practitioner" means a physician, dentist, veterinarian, advanced practice registered nurse, prescribing psychologist, or other person duly authorized by law or rule in the state of Washington to prescribe drugs.
- 38 (34) "Prescription" means an order for drugs or devices issued by 39 a practitioner duly authorized by law or rule in the state of

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Washington to prescribe drugs or devices in the course of his or her professional practice for a legitimate medical purpose.

- (35) "Secretary" means the secretary of health or the secretary's designee.
 - (36) "Shared pharmacy services" means a system that allows a participating pharmacist or pharmacy pursuant to a request from another participating pharmacist or pharmacy to process or fill a prescription or drug order, which may include but is not necessarily limited to preparing, packaging, labeling, data entry, compounding for specific patients, dispensing, performing drug utilization reviews, conducting claims adjudication, obtaining refill authorizations, reviewing therapeutic interventions, or reviewing chart orders.
 - (37) "Statement of deficiency" means a written statement of the deficiencies prepared by the commission, or its designee, identifying one or more violations of law. The report clearly identifies the specific law or rule that has been violated along with a description of the reasons for noncompliance.
- 19 (38) "Wholesaler" means a corporation, individual, or other 20 entity which buys drugs or devices for resale and distribution to 21 corporations, individuals, or entities other than consumers.
- **Sec. 12.** RCW 18.79.260 and 2022 c 14 s 2 are each amended to 23 read as follows:
 - (1) A registered nurse under his or her license may perform for compensation nursing care, as that term is usually understood, to individuals with illnesses, injuries, or disabilities.
 - (2) A registered nurse may, at or under the general direction of a licensed physician and surgeon, dentist, osteopathic physician and surgeon, naturopathic physician, optometrist, podiatric physician and surgeon, physician assistant, advanced registered nurse practitioner, prescribing psychologist, or midwife acting within the scope of his or her license, administer medications, treatments, tests, and inoculations, whether or not the severing or penetrating of tissues is involved and whether or not a degree of independent judgment and skill is required. Such direction must be for acts which are within the scope of registered nursing practice.
- 37 (3) A registered nurse may delegate tasks of nursing care to 38 other individuals where the registered nurse determines that it is in 39 the best interest of the patient.

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(a) The delegating nurse shall:

- 2 (i) Determine the competency of the individual to perform the 3 tasks;
 - (ii) Evaluate the appropriateness of the delegation;
 - (iii) Supervise the actions of the person performing the delegated task; and
 - (iv) Delegate only those tasks that are within the registered nurse's scope of practice.
 - (b) A registered nurse, working for a home health or hospice agency regulated under chapter 70.127 RCW, may delegate the application, instillation, or insertion of medications to a registered or certified nursing assistant under a plan of care.
 - (c) Except as authorized in (b) or (e) of this subsection, a registered nurse may not delegate the administration of medications. Except as authorized in (e) or (f) of this subsection, a registered nurse may not delegate acts requiring substantial skill, and may not delegate piercing or severing of tissues. Acts that require nursing judgment shall not be delegated.
 - (d) No person may coerce a nurse into compromising patient safety by requiring the nurse to delegate if the nurse determines that it is inappropriate to do so. Nurses shall not be subject to any employer reprisal or disciplinary action by the ((nursing care quality assurance commission)) board for refusing to delegate tasks or refusing to provide the required training for delegation if the nurse determines delegation may compromise patient safety.
 - (e) For delegation in community-based care settings or in-home care settings, a registered nurse may delegate nursing care tasks only to registered or certified nursing assistants under chapter 18.88A RCW or home care aides certified under chapter 18.88B RCW. Simple care tasks such as blood pressure monitoring, personal care service, diabetic insulin device set up, verbal verification of insulin dosage for sight-impaired individuals, or other tasks as defined by the ((nursing care quality assurance commission)) board are exempted from this requirement.
 - (i) "Community-based care settings" includes: Community residential programs for people with developmental disabilities, certified by the department of social and health services under chapter 71A.12 RCW; adult family homes licensed under chapter 70.128 RCW; and assisted living facilities licensed under chapter 18.20 RCW.

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Community-based care settings do not include acute care or skilled nursing facilities.

- (ii) "In-home care settings" include an individual's place of temporary or permanent residence, but does not include acute care or skilled nursing facilities, and does not include community-based care settings as defined in (e)(i) of this subsection.
- (iii) Delegation of nursing care tasks in community-based care settings and in-home care settings is only allowed for individuals who have a stable and predictable condition. "Stable and predictable condition" means a situation in which the individual's clinical and behavioral status is known and does not require the frequent presence and evaluation of a registered nurse.
- (iv) The determination of the appropriateness of delegation of a nursing task is at the discretion of the registered nurse. Other than delegation of the administration of insulin by injection for the purpose of caring for individuals with diabetes, the administration of medications by injection, sterile procedures, and central line maintenance may never be delegated.
- (v) When delegating insulin injections under this section, the registered nurse delegator must instruct the individual regarding proper injection procedures and the use of insulin, demonstrate proper injection procedures, and must supervise and evaluate the individual performing the delegated task as required by the ((commission)) board by rule. If the registered nurse delegator determines that the individual is competent to perform the injection properly and safely, supervision and evaluation shall occur at an interval determined by the ((commission)) board by rule.
- (vi)(A) The registered nurse shall verify that the nursing assistant or home care aide, as the case may be, has completed the required core nurse delegation training required in chapter 18.88A or 18.88B RCW prior to authorizing delegation.
- (B) Before commencing any specific nursing tasks authorized to be delegated in this section, a home care aide must be certified pursuant to chapter 18.88B RCW and must comply with RCW 18.88B.070.
- (vii) The nurse is accountable for his or her own individual actions in the delegation process. Nurses acting within the protocols of their delegation authority are immune from liability for any action performed in the course of their delegation duties.
- (viii) Nursing task delegation protocols are not intended to regulate the settings in which delegation may occur, but are intended

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to ensure that nursing care services have a consistent standard of practice upon which the public and the profession may rely, and to safeguard the authority of the nurse to make independent professional decisions regarding the delegation of a task.

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- (f) The delegation of nursing care tasks only to registered or certified nursing assistants under chapter 18.88A RCW or to home care aides certified under chapter 18.88B RCW may include glucose monitoring and testing.
- 9 (g) The ((nursing care quality assurance commission)) board may 10 adopt rules to implement this section.
- 11 (4) Only a person licensed as a registered nurse may instruct 12 nurses in technical subjects pertaining to nursing.
- 13 (5) Only a person licensed as a registered nurse may hold herself 14 or himself out to the public or designate herself or himself as a 15 registered nurse.
- 16 **Sec. 13.** RCW 18.79.260 and 2024 c 239 s 15 are each amended to read as follows:
 - (1) A registered nurse under his or her license may perform for compensation nursing care, as that term is usually understood, to individuals with illnesses, injuries, or disabilities.
 - (2) A registered nurse may, at or under the general direction of a licensed physician and surgeon, dentist, osteopathic physician and surgeon, naturopathic physician, optometrist, podiatric physician and surgeon, physician assistant, advanced practice registered nurse, prescribing psychologist, or midwife acting within the scope of his or her license, administer medications, treatments, tests, and inoculations, whether or not the severing or penetrating of tissues is involved and whether or not a degree of independent judgment and skill is required. Such direction must be for acts which are within the scope of registered nursing practice.
- 31 (3) A registered nurse may delegate tasks of nursing care to 32 other individuals where the registered nurse determines that it is in 33 the best interest of the patient.
 - (a) The delegating nurse shall:
- 35 (i) Determine the competency of the individual to perform the 36 tasks;
- 37 (ii) Evaluate the appropriateness of the delegation;
- 38 (iii) Supervise the actions of the person performing the 39 delegated task; and

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(iv) Delegate only those tasks that are within the registered nurse's scope of practice.

- (b) A registered nurse, working for a home health or hospice agency regulated under chapter 70.127 RCW, may delegate the application, instillation, or insertion of medications to a registered or certified nursing assistant under a plan of care.
- (c) Except as authorized in (b) or (e) of this subsection, a registered nurse may not delegate the administration of medications. Except as authorized in (e) or (f) of this subsection, a registered nurse may not delegate acts requiring substantial skill, and may not delegate piercing or severing of tissues. Acts that require nursing judgment shall not be delegated.
- (d) No person may coerce a nurse into compromising patient safety by requiring the nurse to delegate if the nurse determines that it is inappropriate to do so. Nurses shall not be subject to any employer reprisal or disciplinary action by the board for refusing to delegate tasks or refusing to provide the required training for delegation if the nurse determines delegation may compromise patient safety.
- (e) For delegation in community-based care settings or in-home care settings, a registered nurse may delegate nursing care tasks only to registered or certified nursing assistants under chapter 18.88A RCW or home care aides certified under chapter 18.88B RCW. Simple care tasks such as blood pressure monitoring, personal care service, diabetic insulin device set up, verbal verification of insulin dosage for sight-impaired individuals, or other tasks as defined by the board are exempted from this requirement.
- (i) "Community-based care settings" includes: Community residential programs for people with developmental disabilities, certified by the department of social and health services under chapter 71A.12 RCW; adult family homes licensed under chapter 70.128 RCW; and assisted living facilities licensed under chapter 18.20 RCW. Community-based care settings do not include acute care or skilled nursing facilities.
- (ii) "In-home care settings" include an individual's place of temporary or permanent residence, but does not include acute care or skilled nursing facilities, and does not include community-based care settings as defined in (e)(i) of this subsection.
- (iii) Delegation of nursing care tasks in community-based care settings and in-home care settings is only allowed for individuals who have a stable and predictable condition. "Stable and predictable

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condition" means a situation in which the individual's clinical and behavioral status is known and does not require the frequent presence and evaluation of a registered nurse.

- (iv) The determination of the appropriateness of delegation of a nursing task is at the discretion of the registered nurse. Other than delegation of the administration of insulin by injection for the purpose of caring for individuals with diabetes, the administration of medications by injection, sterile procedures, and central line maintenance may never be delegated.
- (v) When delegating insulin injections under this section, the registered nurse delegator must instruct the individual regarding proper injection procedures and the use of insulin, demonstrate proper injection procedures, and must supervise and evaluate the individual performing the delegated task as required by the board by rule. If the registered nurse delegator determines that the individual is competent to perform the injection properly and safely, supervision and evaluation shall occur at an interval determined by the board by rule.
- (vi)(A) The registered nurse shall verify that the nursing assistant or home care aide, as the case may be, has completed the required core nurse delegation training required in chapter 18.88A or 18.88B RCW prior to authorizing delegation.
 - (B) Before commencing any specific nursing tasks authorized to be delegated in this section, a home care aide must be certified pursuant to chapter 18.88B RCW and must comply with RCW 18.88B.070.
 - (vii) The nurse is accountable for his or her own individual actions in the delegation process. Nurses acting within the protocols of their delegation authority are immune from liability for any action performed in the course of their delegation duties.
 - (viii) Nursing task delegation protocols are not intended to regulate the settings in which delegation may occur, but are intended to ensure that nursing care services have a consistent standard of practice upon which the public and the profession may rely, and to safeguard the authority of the nurse to make independent professional decisions regarding the delegation of a task.
- (f) The delegation of nursing care tasks only to registered or certified nursing assistants under chapter 18.88A RCW or to home care aides certified under chapter 18.88B RCW may include glucose monitoring and testing.
 - (g) The board may adopt rules to implement this section.

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- 1 (4) Only a person licensed as a registered nurse may instruct 2 nurses in technical subjects pertaining to nursing.
- 3 (5) Only a person licensed as a registered nurse may hold herself 4 or himself out to the public or designate herself or himself as a 5 registered nurse.
- 6 **Sec. 14.** RCW 69.50.101 and 2024 c 62 s 17 are each amended to 7 read as follows:

8 The definitions in this section apply throughout this chapter 9 unless the context clearly requires otherwise.

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- (1) "Administer" means to apply a controlled substance, whether by injection, inhalation, ingestion, or any other means, directly to the body of a patient or research subject by:
- (a) a practitioner authorized to prescribe (or, by the practitioner's authorized agent); or
- 15 (b) the patient or research subject at the direction and in the 16 presence of the practitioner.
 - (2) "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 warehouseperson, or employee of the carrier or warehouseperson.
 - (3) "Board" means the Washington state liquor and cannabis board.
 - (4) "Cannabis" means all parts of the plant *Cannabis*, whether growing or not, with a THC concentration greater than 0.3 percent on a dry weight basis during the growing cycle through harvest and usable cannabis. "Cannabis" does not include hemp or industrial hemp as defined in RCW 15.140.020, or seeds used for licensed hemp production under chapter 15.140 RCW.
 - (5) "Cannabis concentrates" means products consisting wholly or in part of the resin extracted from any part of the plant *Cannabis* and having a THC concentration greater than ten percent.
 - (6) "Cannabis processor" means a person licensed by the board to process cannabis into cannabis concentrates, useable cannabis, and cannabis-infused products, package and label cannabis concentrates, useable cannabis, and cannabis-infused products for sale in retail outlets, and sell cannabis concentrates, useable cannabis, and cannabis-infused products at wholesale to cannabis retailers.
- 37 (7) "Cannabis producer" means a person licensed by the board to 38 produce and sell cannabis at wholesale to cannabis processors and 39 other cannabis producers.

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(8) (a) "Cannabis products" means useable cannabis, cannabis concentrates, and cannabis-infused products as defined in this section, including any product intended to be consumed or absorbed inside the body by any means including inhalation, ingestion, or insertion, with any detectable amount of THC.

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- 6 (b) "Cannabis products" also means any product containing only 7 THC content.
 - (c) "Cannabis products" does not include cannabis health and beauty aids as defined in RCW 69.50.575 or products approved by the United States food and drug administration.
 - (9) "Cannabis researcher" means a person licensed by the board to produce, process, and possess cannabis for the purposes of conducting research on cannabis and cannabis-derived drug products.
 - (10) "Cannabis retailer" means a person licensed by the board to sell cannabis concentrates, useable cannabis, and cannabis-infused products in a retail outlet.
 - (11) "Cannabis-infused products" means products that contain cannabis or cannabis extracts, are intended for human use, are derived from cannabis as defined in subsection (4) of this section, and have a THC concentration no greater than ten percent. The term "cannabis-infused products" does not include either useable cannabis or cannabis concentrates.
- 23 (12) "CBD concentration" has the meaning provided in RCW 24 69.51A.010.
- 25 (13) "CBD product" means any product containing or consisting of cannabidiol.
- 27 (14) "Commission" means the pharmacy quality assurance 28 commission.
 - (15) "Controlled substance" means a drug, substance, or immediate precursor included in Schedules I through V as set forth in federal or state laws, or federal or commission rules, but does not include hemp or industrial hemp as defined in RCW 15.140.020.
 - (16)(a) "Controlled substance analog" means a substance the chemical structure of which is substantially similar to the chemical structure of a controlled substance in Schedule I or II and:
- (i) that has a stimulant, depressant, or hallucinogenic effect on the central nervous system substantially similar to the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance included in Schedule I or II; or

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- (ii) with respect to a particular individual, that the individual represents or intends to have a stimulant, depressant, or hallucinogenic effect on the central nervous system substantially similar to the stimulant, depressant, or hallucinogenic effect on the central nervous system of a controlled substance included in Schedule I or II.
 - (b) The term does not include:
 - (i) a controlled substance;

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- 9 (ii) a substance for which there is an approved new drug 10 application;
- (iii) a substance with respect to which an exemption is in effect for investigational use by a particular person under Section 505 of the federal food, drug, and cosmetic act, 21 U.S.C. Sec. 355, or chapter 69.77 RCW to the extent conduct with respect to the substance is pursuant to the exemption; or
- 16 (iv) any substance to the extent not intended for human 17 consumption before an exemption takes effect with respect to the 18 substance.
- 19 (17) "Deliver" or "delivery" means the actual or constructive 20 transfer from one person to another of a substance, whether or not 21 there is an agency relationship.
 - (18) "Department" means the department of health.
- 23 (19) "Designated provider" has the meaning provided in RCW 69.51A.010.
- 25 (20) "Dispense" means the interpretation of a prescription or 26 order for a controlled substance and, pursuant to that prescription 27 or order, the proper selection, measuring, compounding, labeling, or 28 packaging necessary to prepare that prescription or order for 29 delivery.
 - (21) "Dispenser" means a practitioner who dispenses.
- 31 (22) "Distribute" means to deliver other than by administering or 32 dispensing a controlled substance.
 - (23) "Distributor" means a person who distributes.
- 34 (24) "Drug" means (a) a controlled substance recognized as a drug 35 in the official United States pharmacopoeia/national formulary or the 36 official homeopathic pharmacopoeia of the United States, or any 37 supplement to them; (b) controlled substances intended for use in the 38 diagnosis, cure, mitigation, treatment, or prevention of disease in 39 individuals or animals; (c) controlled substances (other than food) 40 intended to affect the structure or any function of the body of

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- individuals or animals; and (d) controlled substances intended for use as a component of any article specified in (a), (b), or (c) of this subsection. The term does not include devices or their components, parts, or accessories.
 - (25) "Drug enforcement administration" means the drug enforcement administration in the United States Department of Justice, or its successor agency.
 - (26) "Electronic communication of prescription information" means the transmission of a prescription or refill authorization for a drug of a practitioner using computer systems. The term does not include a prescription or refill authorization verbally transmitted by telephone nor a facsimile manually signed by the practitioner.
- 13 (27) "Immature plant or clone" means a plant or clone that has no 14 flowers, is less than twelve inches in height, and is less than 15 twelve inches in diameter.
 - (28) "Immediate precursor" means a substance:

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- (a) that the commission has found to be and by rule designates as being the principal compound commonly used, or produced primarily for use, in the manufacture of a controlled substance;
- (b) that is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance; and
- 22 (c) the control of which is necessary to prevent, curtail, or 23 limit the manufacture of the controlled substance.
 - (29) "Isomer" means an optical isomer, but in subsection (33) (e) of this section, RCW 69.50.204(1) (l) and (hh), and 69.50.206(2) (d), the term includes any geometrical isomer; in RCW 69.50.204(1) (h) and (pp), and 69.50.210(3)($(\frac{1}{1})$) the term includes any positional isomer; and in RCW 69.50.204(1)(ii), 69.50.204(3), and 69.50.208(1)($(\frac{1}{1})$) the term includes any positional or geometric isomer.
 - (30) "Lot" means a definite quantity of cannabis, cannabis concentrates, useable cannabis, or cannabis-infused product identified by a lot number, every portion or package of which is uniform within recognized tolerances for the factors that appear in the labeling.
 - (31) "Lot number" must identify the licensee by business or trade name and Washington state unified business identifier number, and the date of harvest or processing for each lot of cannabis, cannabis concentrates, useable cannabis, or cannabis-infused product.
- 39 (32) "Manufacture" means the production, preparation, 40 propagation, compounding, conversion, or processing of a controlled

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- substance, either directly or indirectly or 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. The term does not include the preparation, compounding, packaging, repackaging, labeling, or relabeling of a controlled substance:
 - (a) by a practitioner as an incident to the practitioner's administering or dispensing of a controlled substance in the course of the practitioner's professional practice; or
 - (b) by a practitioner, or by the practitioner's authorized agent under the practitioner's supervision, for the purpose of, or as an incident to, research, teaching, or chemical analysis and not for sale.
 - (33) "Narcotic drug" means any of the following, whether produced directly or indirectly by extraction from substances of vegetable origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis:
 - (a) Opium, opium derivative, and any derivative of opium or opium derivative, including their salts, isomers, and salts of isomers, whenever the existence of the salts, isomers, and salts of isomers is possible within the specific chemical designation. The term does not include the isoquinoline alkaloids of opium.
 - (b) Synthetic opiate and any derivative of synthetic opiate, including their isomers, esters, ethers, salts, and salts of isomers, esters, and ethers, whenever the existence of the isomers, esters, ethers, and salts is possible within the specific chemical designation.
 - (c) Poppy straw and concentrate of poppy straw.
- 30 (d) Coca leaves, except coca leaves and extracts of coca leaves 31 from which cocaine, ecgonine, and derivatives or ecgonine or their 32 salts have been removed.
 - (e) Cocaine, or any salt, isomer, or salt of isomer thereof.
 - (f) Cocaine base.

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- 35 (g) Ecgonine, or any derivative, salt, isomer, or salt of isomer 36 thereof.
- 37 (h) Any compound, mixture, or preparation containing any quantity 38 of any substance referred to in (a) through (g) of this subsection.
- 39 (34) "Opiate" means any substance having an addiction-forming or 40 addiction-sustaining liability similar to morphine or being capable

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- of conversion into a drug having addiction-forming or addictionsustaining liability. The term includes opium, substances derived from opium (opium derivatives), and synthetic opiates. The term does not include, unless specifically designated as controlled under RCW 69.50.201, the dextrorotatory isomer of 3-methoxy-n-methylmorphinan and its salts (dextromethorphan). The term includes the racemic and levorotatory forms of dextromethorphan.
- 8 (35) "Opium poppy" means the plant of the species Papaver 9 somniferum L., except its seeds.
- 10 (36) "Package" means a container that has a single unit or group 11 of units.
 - (37) "Person" means individual, corporation, business trust, estate, trust, partnership, association, joint venture, government, governmental subdivision or agency, or any other legal or commercial entity.
 - (38) "Plant" has the meaning provided in RCW 69.51A.010.
- 17 (39) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.
 - (40) "Practitioner" means:

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- (a) A physician under chapter 18.71 RCW; a physician assistant under chapter 18.71A RCW; an osteopathic physician and surgeon under chapter 18.57 RCW; an optometrist licensed under chapter 18.53 RCW who is certified by the optometry board under RCW 18.53.010 subject to any limitations in RCW 18.53.010; a dentist under chapter 18.32 RCW; a podiatric physician and surgeon under chapter 18.22 RCW; a veterinarian under chapter 18.92 RCW; a registered nurse, advanced practice registered nurse ((practitioner)), or licensed practical nurse under chapter 18.79 RCW; a naturopathic physician under chapter 18.36A RCW who is licensed under RCW 18.36A.030 subject to any limitations in RCW 18.36A.040; a psychologist under chapter 18.83 RCW who is certified as a prescribing psychologist under section 3 of this act; a pharmacist under chapter 18.64 RCW or a scientific investigator under this chapter, licensed, registered or otherwise permitted insofar as is consistent with those licensing laws to distribute, dispense, conduct research with respect to or administer a controlled substance in the course of their professional practice or research in this state.
- 38 (b) A pharmacy, hospital or other institution licensed, 39 registered, or otherwise permitted to distribute, dispense, conduct

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research with respect to or to administer a controlled substance in the course of professional practice or research in this state.

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- (c) A physician licensed to practice medicine and surgery, a physician licensed to practice osteopathic medicine and surgery, a dentist licensed to practice dentistry, a podiatric physician and surgeon licensed to practice podiatric medicine and surgery, a licensed physician assistant or a licensed osteopathic physician assistant specifically approved to prescribe controlled substances by his or her state's medical commission or equivalent and his or her participating physician as defined in RCW 18.71A.010, an advanced practice registered nurse ((practitioner)) licensed to prescribe controlled substances, or a veterinarian licensed to practice veterinary medicine in any state of the United States.
- (41) "Prescription" means an order for controlled substances issued by a practitioner duly authorized by law or rule in the state of Washington to prescribe controlled substances within the scope of his or her professional practice for a legitimate medical purpose.
- (42) "Production" includes the manufacturing, planting, cultivating, growing, or harvesting of a controlled substance.
- 20 (43) "Qualifying patient" has the meaning provided in RCW 21 69.51A.010.
- 22 (44) "Recognition card" has the meaning provided in RCW 23 69.51A.010.
 - (45) "Retail outlet" means a location licensed by the board for the retail sale of cannabis concentrates, useable cannabis, and cannabis-infused products.
- 27 (46) "Secretary" means the secretary of health or the secretary's designee.
 - (47) "Social equity plan" means a plan that addresses at least some of the elements outlined in this subsection (47), along with any additional plan components or requirements approved by the board following consultation with the task force created in RCW 69.50.336. The plan may include:
- 34 (a) A statement that indicates how the cannabis licensee will work to promote social equity goals in their community;
- 36 (b) A description of how the cannabis licensee will meet social as equity goals as defined in RCW 69.50.335;
- 38 (c) The composition of the workforce the licensee has employed or 39 intends to hire; and

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- 1 (d) Business plans involving partnerships or assistance to 2 organizations or residents with connections to populations with a 3 history of high rates of enforcement of cannabis prohibition.
 - (48) "State," unless the context otherwise requires, means a state of the United States, the District of Columbia, the Commonwealth of Puerto Rico, or a territory or insular possession subject to the jurisdiction of the United States.
 - (49) "THC concentration" means percent of tetrahydrocannabinol content of any part of the plant *Cannabis*, or per volume or weight of cannabis product, or the combined percent of tetrahydrocannabinol and tetrahydrocannabinolic acid in any part of the plant *Cannabis* regardless of moisture content.
 - (50) "Ultimate user" means an individual who lawfully possesses a controlled substance for the individual's own use or for the use of a member of the individual's household or for administering to an animal owned by the individual or by a member of the individual's household.
 - (51) "Unit" means an individual consumable item within a package of one or more consumable items in solid, liquid, gas, or any form intended for human consumption.
- 21 (52) "Useable cannabis" means dried cannabis flowers. The term 22 "useable cannabis" does not include either cannabis-infused products 23 or cannabis concentrates.
 - (53) "Youth access" means the level of interest persons under the age of twenty-one may have in a vapor product, as well as the degree to which the product is available or appealing to such persons, and the likelihood of initiation, use, or addiction by adolescents and young adults.
- 29 **Sec. 15.** RCW 69.41.010 and 2024 c 102 s 1 are each amended to 30 read as follows:
- As used in this chapter, the following terms have the meanings indicated unless the context clearly requires otherwise:
- 33 (1) "Administer" means the direct application of a legend drug 34 whether by injection, inhalation, ingestion, or any other means, to 35 the body of a patient or research subject by:
 - (a) A practitioner; or

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- 37 (b) The patient or research subject at the direction of the 38 practitioner.
 - (2) "Commission" means the pharmacy quality assurance commission.

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- 1 (3) "Community-based care settings" include: Community residential programs for persons with developmental disabilities, certified by the department of social and health services under chapter 71A.12 RCW; adult family homes licensed under chapter 70.128 RCW; and assisted living facilities licensed under chapter 18.20 RCW. Community-based care settings do not include acute care or skilled nursing facilities.
 - (4) "Deliver" or "delivery" means the actual, constructive, or attempted transfer from one person to another of a legend drug, whether or not there is an agency relationship.
 - (5) "Department" means the department of health.
 - (6) "Dispense" means the interpretation of a prescription or order for a legend drug and, pursuant to that prescription or order, the proper selection, measuring, compounding, labeling, or packaging necessary to prepare that prescription or order for delivery.
 - (7) "Dispenser" means a practitioner who dispenses.
- 17 (8) "Distribute" means to deliver other than by administering or 18 dispensing a legend drug.
 - (9) "Distributor" means a person who distributes.
- 20 (10) "Drug" means:

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- (a) Substances recognized as drugs in the official United States pharmacopoeia, official homeopathic pharmacopoeia of the United States, or official national formulary, or any supplement to any of them;
- 25 (b) Substances intended for use in the diagnosis, cure, 26 mitigation, treatment, or prevention of disease in human beings or 27 animals;
 - (c) Substances (other than food, minerals or vitamins) intended to affect the structure or any function of the body of human beings or animals; and
- 31 (d) Substances intended for use as a component of any article 32 specified in (a), (b), or (c) of this subsection. It does not include 33 devices or their components, parts, or accessories.
 - (11) "Electronic communication of prescription information" means the transmission of a prescription or refill authorization for a drug of a practitioner using computer systems. The term does not include a prescription or refill authorization transmitted verbally by telephone nor a facsimile manually signed by the practitioner.
- 39 (12) "In-home care settings" include an individual's place of 40 temporary and permanent residence, but does not include acute care or

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- skilled nursing facilities, and does not include community-based care settings.
 - (13) "Legend drugs" means any drugs which are required by state law or regulation of the pharmacy quality assurance commission to be dispensed on prescription only or are restricted to use by practitioners only.
 - (14) "Legible prescription" means a prescription or medication order issued by a practitioner that is capable of being read and understood by the pharmacist filling the prescription or the nurse or other practitioner implementing the medication order. A prescription must be hand printed, typewritten, or electronically generated.
 - (15) "Medication assistance" means assistance rendered by a nonpractitioner to an individual residing in a community-based care setting or in-home care setting to facilitate the individual's self-administration of a legend drug or controlled substance. It includes reminding or coaching the individual, handing the medication container to the individual, opening the individual's medication container, using an enabler, or placing the medication in the individual's hand, and such other means of medication assistance as defined by rule adopted by the department. A nonpractitioner may help in the preparation of legend drugs or controlled substances for self-administration where a practitioner has determined and communicated orally or by written direction that such medication preparation assistance is necessary and appropriate. Medication assistance shall not include assistance with intravenous medications or injectable medications, except prefilled insulin syringes.
 - (16) "Person" means individual, corporation, government or governmental subdivision or agency, business trust, estate, trust, partnership or association, or any other legal entity.
 - (17) "Practitioner" means:

(a) A physician under chapter 18.71 RCW, an osteopathic physician or an osteopathic physician and surgeon under chapter 18.57 RCW, a dentist under chapter 18.32 RCW, a podiatric physician and surgeon under chapter 18.22 RCW, an acupuncturist or acupuncture and Eastern medicine practitioner to the extent authorized under chapter 18.06 RCW and the rules adopted under RCW 18.06.010(1)(m), a veterinarian under chapter 18.92 RCW, a registered nurse, advanced practice registered nurse ((practitioner)), or licensed practical nurse under chapter 18.79 RCW, an optometrist under chapter 18.53 RCW who is certified by the optometry board under RCW 18.53.010, a physician

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- assistant under chapter 18.71A RCW, a naturopath licensed under 1 chapter 18.36A RCW, a licensed athletic trainer to the extent 2 authorized under chapter 18.250 RCW, a pharmacist under chapter 18.64 3 RCW, a psychologist under chapter 18.83 RCW who is certified as a 4 prescribing psychologist under section 3 of this act, when acting 5 6 under the required supervision of a dentist licensed under chapter 18.32 RCW, a dental hygienist licensed under chapter 18.29 RCW, a 7 licensed dental therapist to the extent authorized under chapter 8 18.265 RCW, or a licensed midwife to the extent authorized under 9 chapter 18.50 RCW; 10
 - (b) A pharmacy, hospital, or other institution licensed, registered, or otherwise permitted to distribute, dispense, conduct research with respect to, or to administer a legend drug in the course of professional practice or research in this state; and

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- 15 (c) A physician licensed to practice medicine and surgery or a 16 physician licensed to practice osteopathic medicine and surgery in 17 any state, or province of Canada, which shares a common border with 18 the state of Washington.
- 19 (18) "Secretary" means the secretary of health or the secretary's designee.
- 21 **Sec. 16.** RCW 69.41.030 and 2024 c 102 s 2 are each amended to 22 read as follows:
 - (1) It shall be unlawful for any person to sell or deliver any legend drug, or knowingly possess any legend drug, or knowingly use any legend drug in a public place, except upon the order or prescription of a physician under chapter 18.71 RCW, an osteopathic physician and surgeon under chapter 18.57 RCW, an optometrist licensed under chapter 18.53 RCW who is certified by the optometry board under RCW 18.53.010, a dentist under chapter 18.32 RCW, a podiatric physician and surgeon under chapter 18.22 RCW, a licensed midwife to the extent authorized under chapter 18.50 RCW, a veterinarian under chapter 18.92 RCW, a commissioned medical or dental officer in the United States armed forces or public health service in the discharge of his or her official duties, a duly licensed physician or dentist employed by the veterans administration in the discharge of his or her official duties, a registered nurse or advanced practice registered nurse ((practitioner)) under chapter 18.79 RCW when authorized by the board of nursing, a pharmacist licensed under chapter 18.64 RCW to the extent permitted by drug

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1 therapy guidelines or protocols established under RCW 18.64.011 and authorized by the commission and approved by a practitioner 2 3 authorized to prescribe drugs, a physician assistant under chapter 18.71A RCW when authorized by the Washington medical commission, \underline{a} 4 psychologist under chapter 18.83 RCW who is certified as a 5 6 prescribing psychologist under section 3 of this act, or any of the 7 following professionals in any province of Canada that shares a common border with the state of Washington or in any state of the 8 United States: A physician licensed to practice medicine and surgery 9 or a physician licensed to practice osteopathic medicine and surgery, 10 a dentist licensed to practice dentistry, a podiatric physician and 11 12 surgeon licensed to practice podiatric medicine and surgery, licensed advanced practice registered nurse ((practitioner)), a 13 licensed physician assistant, or a veterinarian licensed to practice 14 veterinary medicine: PROVIDED, HOWEVER, That the above provisions 15 16 shall not apply to sale, delivery, or possession by drug wholesalers 17 or drug manufacturers, or their agents or employees, or to any 18 practitioner acting within the scope of his or her license, or to a 19 common or contract carrier or warehouse operator, or any employee thereof, whose possession of any legend drug is in the usual course 20 of business or employment: PROVIDED FURTHER, That nothing in this 21 chapter or chapter 18.64 RCW shall prevent a family planning clinic 22 23 that is under contract with the health care authority from selling, delivering, possessing, and dispensing commercially prepackaged oral 24 25 contraceptives prescribed by authorized, licensed health care 26 practitioners: PROVIDED FURTHER, That nothing in this chapter prohibits possession or delivery of legend drugs by an authorized 27 28 collector or other person participating in the operation of a drug 29 take-back program authorized in chapter 69.48 RCW.

(2) (a) A violation of this section involving the sale, delivery, or possession with intent to sell or deliver is a class B felony punishable according to chapter 9A.20 RCW.

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- (b) A violation of this section involving knowing possession is a misdemeanor. The prosecutor is encouraged to divert such cases for assessment, treatment, or other services.
- 36 (c) A violation of this section involving knowing use in a public 37 place is a misdemeanor. The prosecutor is encouraged to divert such 38 cases for assessment, treatment, or other services.

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- 1 (d) No person may be charged with both knowing possession and 2 knowing use in a public place under this section relating to the same 3 course of conduct.
- (e) In lieu of jail booking and referral to the prosecutor for a 4 violation of this section involving knowing possession, or knowing 5 6 use in a public place, law enforcement is encouraged to offer a referral to assessment and services available under RCW 10.31.110 or 7 other program or entity responsible for receiving referrals in lieu 8 of legal system involvement, which may include, but are not limited 9 to, arrest and jail alternative programs established under RCW 10 36.28A.450, law enforcement assisted diversion programs established 11 12 under RCW 71.24.589, and the recovery navigator program established under RCW 71.24.115. 13
- 14 (3) For the purposes of this section, "public place" has the same 15 meaning as defined in RCW 66.04.010, but the exclusions in RCW 16 66.04.011 do not apply.
- 17 (4) For the purposes of this section, "use any legend drug" means 18 to introduce the drug into the human body by injection, inhalation, 19 ingestion, or any other means.
- NEW SECTION. Sec. 17. Sections 6 and 8 of this act expire 21 October 1, 2025.
- NEW SECTION. Sec. 18. Sections 7 and 9 of this act take effect October 1, 2025.
- NEW SECTION. Sec. 19. Section 12 of this act expires June 30, 25 2027.
- NEW SECTION. Sec. 20. Section 13 of this act takes effect June 30, 2027.

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