S-3858.1

SENATE BILL 6019

State of Washington 68th Legislature 2024 Regular Session

By Senator Muzzall

Prefiled 01/05/24.

AN ACT Relating to expanding prescriptive authority for pharmacists; amending RCW 18.64.011 and 69.41.030; reenacting and amending RCW 69.50.101; adding a new section to chapter 18.64 RCW; and providing an effective date.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF WASHINGTON:

Sec. 1. RCW 18.64.011 and 2021 c 78 s 1 are each 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 does not constitute compounding if prepared pursuant to a 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) "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.

(12) "Distribute" means the delivery of a drug or device other than by administering or dispensing.

(13) "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.

(14) "Drugs" means:

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

(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.

(15) "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.

(16) "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.

(17) "Institutional facility" means any organization whose primary purpose is to provide a physical environment for patients to obtain health care services including, but not limited to, services in a hospital, long-term care facility, hospice program, mental health facility, drug abuse treatment center, residential habilitation center, or a local, state, or federal correction facility.

(18) "Labeling" means the process of preparing and affixing a label to any drug or device container. The label must include all information required by current federal and state law and pharmacy rules.

(19) "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.

(20) "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.

(21) "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

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

(22) "Manufacturer" means a person, corporation, or other entity engaged in the manufacture of drugs or devices.

(23) "Nonlegend" or "nonprescription" drugs means any drugs which may be lawfully sold without a prescription.

(24) "Person" means an individual, corporation, government, governmental subdivision or agency, business trust, estate, trust, partnership or association, or any other legal entity.

(25) "Pharmacist" means a person duly licensed by the commission to engage in the practice of pharmacy.

(26) "Pharmacy" means every place properly licensed by the commission where the practice of pharmacy is conducted.

(27) "Poison" does not include any article or mixture covered by the Washington pesticide control act (chapter 15.58 RCW), as enacted or hereafter amended.

(28) "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 prescribing and ordering of drugs and devices as authorized by the commission in rule; 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.

(29) "Practitioner" means a physician, dentist, veterinarian, nurse, or other person duly authorized by law or rule in the state of Washington to prescribe drugs.

(30) "Prescription" means an order for drugs or devices issued by a practitioner duly authorized by law or rule in the state of Washington to prescribe drugs or devices in the course of his or her professional practice for a legitimate medical purpose.

(31) "Secretary" means the secretary of health or the secretary's designee.

(32) "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.

(33) "Wholesaler" means a corporation, individual, or other entity which buys drugs or devices for resale and distribution to corporations, individuals, or entities other than consumers.

NEW SECTION. **Sec. 2.** A new section is added to chapter 18.64 RCW to read as follows:

By July 1, 2026, the commission shall adopt rules identifying specific drugs and devices, types or classes of drugs and devices, or both, that a pharmacist may prescribe in the absence of written guidelines or protocols previously established and approved for the pharmacist's practice by a practitioner authorized to prescribe drugs. The rules may also establish the types of patients or circumstances in which a pharmacist may or may not prescribe or order drugs or devices and any required education, training, or continuing education that must be completed prior to prescribing or ordering drugs or devices.

Sec. 3. RCW 69.41.030 and 2023 1st sp.s. c 1 s 4 are each amended to 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 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 registered nurse practitioner under chapter 18.79 RCW when authorized by the ((~~nursing care quality assurance commission~~)) board of nursing, a pharmacist licensed under chapter 18.64 RCW to the extent permitted ((~~by drug therapy guidelines or protocols established under RCW 18.64.011 and authorized by the commission and approved by a practitioner authorized to prescribe drugs~~)) under chapter 18.64 RCW or when authorized by the commission, a physician assistant under chapter 18.71A RCW when authorized by the Washington medical commission, or any of the following professionals in any province of Canada that shares a common border with the state of Washington or in any state of the United States: A physician licensed to practice medicine and surgery or 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 advanced registered nurse practitioner, a licensed physician assistant, or a veterinarian licensed to practice veterinary medicine: PROVIDED, HOWEVER, That the above provisions shall not apply to sale, delivery, or possession by drug wholesalers or drug manufacturers, or their agents or employees, or to any practitioner acting within the scope of his or her license, or to a common or contract carrier or warehouse operator, or any employee thereof, whose possession of any legend drug is in the usual course of business or employment: PROVIDED FURTHER, That nothing in this chapter or chapter 18.64 RCW shall prevent a family planning clinic that is under contract with the health care authority from selling, delivering, possessing, and dispensing commercially prepackaged oral contraceptives prescribed by authorized, licensed health care practitioners: PROVIDED FURTHER, That nothing in this chapter prohibits possession or delivery of legend drugs by an authorized collector or other person participating in the operation of a drug 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.

(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.

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

(d) No person may be charged with both knowing possession and knowing use in a public place under this section relating to the same course of conduct.

(e) In lieu of jail booking and referral to the prosecutor for a violation of this section involving knowing possession, or knowing use in a public place, law enforcement is encouraged to offer a referral to assessment and services available under RCW 10.31.110 or other program or entity responsible for receiving referrals in lieu of legal system involvement, which may include, but are not limited to, arrest and jail alternative programs established under RCW 36.28A.450, law enforcement assisted diversion programs established under RCW 71.24.589, and the recovery navigator program established under RCW 71.24.115.

(3) For the purposes of this section, "public place" has the same meaning as defined in RCW 66.04.010, but the exclusions in RCW 66.04.011 do not apply.

(4) For the purposes of this section, "use any legend drug" means to introduce the drug into the human body by injection, inhalation, ingestion, or any other means.

Sec. 4. RCW 69.50.101 and 2023 c 365 s 2 and 2023 c 220 s 6 are each reenacted and amended to read as follows:

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

((~~(a) [(1)]~~)) (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:

((~~(1) [(a)] a~~)) (a) A practitioner authorized to prescribe (or, by the practitioner's authorized agent); or

((~~(2) [(b)] the~~)) (b) The patient or research subject at the direction and in the presence of the practitioner.

((~~(b) [(2)]~~)) (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.

((~~(c) [(3)]~~)) (3) "Board" means the Washington state liquor and cannabis board.

((~~(d) [(4)]~~)) (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.

((~~(e) [(5)]~~)) (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.

((~~(f) [(6)]~~)) (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.

((~~(g) [(7)]~~)) (7) "Cannabis producer" means a person licensed by the board to produce and sell cannabis at wholesale to cannabis processors and other cannabis producers.

((~~(h)(1) [(8)(a)]~~)) (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.

((~~(2) [(b)]~~)) (b) "Cannabis products" also means any product containing only THC content.

((~~(3) [(c)]~~)) (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.

((~~(i) [(9)]~~)) (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.

((~~(j) [(10)]~~)) (10) "Cannabis retailer" means a person licensed by the board to sell cannabis concentrates, useable cannabis, and cannabis-infused products in a retail outlet.

((~~(k) [(11)]~~)) (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 ((~~(d) [(4)]~~)) (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.

((~~(l) [(12)]~~)) (12) "CBD concentration" has the meaning provided in RCW 69.51A.010.

((~~(m) [(13)]~~)) (13) "CBD product" means any product containing or consisting of cannabidiol.

((~~(n) [(14)]~~)) (14) "Commission" means the pharmacy quality assurance commission.

((~~(o) [(15)]~~)) (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.

((~~(p)(1) [(16)(a)]~~)) (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

(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.

((~~(2) [(b)]~~)) (b) The term does not include:

(i) a controlled substance;

(ii) a substance for which there is an approved new drug 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

(iv) any substance to the extent not intended for human consumption before an exemption takes effect with respect to the substance.

((~~(q) [(17)]~~)) (17) "Deliver" or "delivery" means the actual or constructive transfer from one person to another of a substance, whether or not there is an agency relationship.

((~~(r) [(18)]~~)) (18) "Department" means the department of health.

((~~(s) [(19)]~~)) (19) "Designated provider" has the meaning provided in RCW 69.51A.010.

((~~(t) [(20)]~~)) (20) "Dispense" means the interpretation of a prescription or order for a controlled substance and, pursuant to that prescription or order, the proper selection, measuring, compounding, labeling, or packaging necessary to prepare that prescription or order for delivery.

((~~(u) [(21)]~~)) (21) "Dispenser" means a practitioner who dispenses.

((~~(v) [(22)]~~)) (22) "Distribute" means to deliver other than by administering or dispensing a controlled substance.

((~~(w) [(23)]~~)) (23) "Distributor" means a person who distributes.

((~~(x) [(24)]~~)) (24) "Drug" means ((~~(1) [(a)]~~)) (a) a controlled substance recognized as a drug in the official United States pharmacopoeia/national formulary or the official homeopathic pharmacopoeia of the United States, or any supplement to them; ((~~(2) [(b)]~~)) (b) controlled substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in individuals or animals; ((~~(3) [(c)]~~)) (c) controlled substances (other than food) intended to affect the structure or any function of the body of individuals or animals; and ((~~(4) [(d)]~~)) (d) controlled substances intended for use as a component of any article specified in ((~~(1), (2), or (3) [(a), (b), or (c)]~~)) (a), (b), or (c) of this subsection. The term does not include devices or their components, parts, or accessories.

((~~(y) [(25)]~~)) (25) "Drug enforcement administration" means the drug enforcement administration in the United States Department of Justice, or its successor agency.

((~~(z) [(26)]~~)) (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.

((~~(aa) [(27)]~~)) (27) "Immature plant or clone" means a plant or clone that has no flowers, is less than twelve inches in height, and is less than twelve inches in diameter.

((~~(bb) [(28)]~~)) (28) "Immediate precursor" means a substance:

((~~(1) [(a)] that~~)) (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;

((~~(2) [(b)] that~~)) (b) That is an immediate chemical intermediary used or likely to be used in the manufacture of a controlled substance; and

((~~(3) [(c)] the~~)) (c) The control of which is necessary to prevent, curtail, or limit the manufacture of the controlled substance.

((~~(cc) [(29)]~~)) (29) "Isomer" means an optical isomer, but in subsection ((~~(gg)(5) [(33)(e)]~~)) (33)(e) of this section, RCW 69.50.204((~~(a) (12) and (34) [(1) (l) and (hh)]~~)) (1) (l) and (hh), and 69.50.206((~~(b)(4) [(2)(d)]~~)) (2)(d), the term includes any geometrical isomer; in RCW 69.50.204((~~(a) (8) and (42) [(1) (h) and (pp)]~~)) (1) (h) and (pp), and 69.50.210((~~(c) [(3)]~~)) (3) the term includes any positional isomer; and in RCW 69.50.204((~~(a)(35) [(1)(ii)]~~)) (1)(ii), 69.50.204((~~(c) [(3)]~~)) (3), and 69.50.208((~~(a) [(1)]~~)) (1) the term includes any positional or geometric isomer.

((~~(dd) [(30)]~~)) (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.

((~~(ee) [(31)]~~)) (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.

((~~(ff) [(32)]~~)) (32) "Manufacture" means the production, preparation, propagation, compounding, conversion, or processing of a controlled 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:

((~~(1) [(a)] by~~)) (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

((~~(2) [(b)] by~~)) (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.

((~~(gg) [(33)]~~)) (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:

((~~(1) [(a)]~~)) (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.

((~~(2) [(b)]~~)) (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.

((~~(3) [(c)]~~)) (c) Poppy straw and concentrate of poppy straw.

((~~(4) [(d)]~~)) (d) Coca leaves, except coca leaves and extracts of coca leaves from which cocaine, ecgonine, and derivatives or ecgonine or their salts have been removed.

((~~(5) [(e)]~~)) (e) Cocaine, or any salt, isomer, or salt of isomer thereof.

((~~(6) [(f)]~~)) (f) Cocaine base.

((~~(7) [(g)]~~)) (g) Ecgonine, or any derivative, salt, isomer, or salt of isomer thereof.

((~~(8) [(h)]~~)) (h) Any compound, mixture, or preparation containing any quantity of any substance referred to in ((~~(1) [(a)]~~)) (a) through ((~~(7) [(g)]~~)) (g) of this subsection.

((~~(hh) [(34)]~~)) (34) "Opiate" means any substance having an addiction-forming or addiction-sustaining liability similar to morphine or being capable of conversion into a drug having addiction-

forming or addiction-sustaining 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.

((~~(ii) [(35)]~~)) (35) "Opium poppy" means the plant of the species Papaver somniferum L., except its seeds.

((~~(jj) [(36)]~~)) (36) "Package" means a container that has a single unit or group of units.

((~~(kk) [(37)]~~)) (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.

((~~(ll) [(38)]~~)) (38) "Plant" has the meaning provided in RCW 69.51A.010.

((~~(mm) [(39)]~~)) (39) "Poppy straw" means all parts, except the seeds, of the opium poppy, after mowing.

((~~(nn) [(40)]~~)) (40) "Practitioner" means:

((~~(1) [(a)]~~)) (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 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 pharmacist under chapter 18.64 RCW subject to any limitations in RCW 18.64.011, section 2 of this act, and rules adopted by the commission; 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.

((~~(2) [(b)]~~)) (b) A pharmacy, hospital or other institution licensed, registered, or otherwise permitted to distribute, dispense, conduct research with respect to or to administer a controlled

substance in the course of professional practice or research in this state.

((~~(3) [(c)]~~)) (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 supervising physician, an advanced registered nurse practitioner licensed to prescribe controlled substances, or a veterinarian licensed to practice veterinary medicine in any state of the United States.

((~~(oo) [(41)]~~)) (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.

((~~(pp) [(42)]~~)) (42) "Production" includes the manufacturing, planting, cultivating, growing, or harvesting of a controlled substance.

((~~(qq) [(43)]~~)) (43) "Qualifying patient" has the meaning provided in RCW 69.51A.010.

((~~(rr) [(44)]~~)) (44) "Recognition card" has the meaning provided in RCW 69.51A.010.

((~~(ss) [(45)]~~)) (45) "Retail outlet" means a location licensed by the board for the retail sale of cannabis concentrates, useable cannabis, and cannabis-infused products.

((~~(tt) [(46)]~~)) (46) "Secretary" means the secretary of health or the secretary's designee.

((~~(uu) [(47)]~~)) (47) "Social equity plan" means a plan that addresses at least some of the elements outlined in this subsection ((~~(uu) [(47)]~~)) (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:

((~~(1) [(a)]~~)) (a) A statement that indicates how the cannabis licensee will work to promote social equity goals in their community;

((~~(2) [(b)]~~)) (b) A description of how the cannabis licensee will meet social equity goals as defined in RCW 69.50.335;

((~~(3) [(c)]~~)) (c) The composition of the workforce the licensee has employed or intends to hire; and

((~~(4) [(d)]~~)) (d) Business plans involving partnerships or assistance to organizations or residents with connections to populations with a history of high rates of enforcement of cannabis prohibition.

((~~(vv) [(48)]~~)) (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.

((~~(ww) [(49)]~~)) (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.

((~~(xx) [(50)]~~)) (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.

((~~(yy) [(51)]~~)) (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.

((~~(zz) [(52)]~~)) (52) "Useable cannabis" means dried cannabis flowers. The term "useable cannabis" does not include either cannabis-infused products or cannabis concentrates.

((~~(aaa) [(53)]~~)) (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.

NEW SECTION. **Sec. 5.** Sections 1, 3, and 4 of this act take effect July 1, 2026.

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