# Senate Bill 295

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## SUMMARY

The following summary is not prepared by the sponsors of the measure and is not a part of the body thereof subject to consideration by the Legislative Assembly. It is an editor's brief statement of the essential features of the measure **as introduced.** The statement includes a measure digest written in compliance with applicable readability standards.

Digest: The Act lets a pharmacist diagnose and treat some health conditions. (Flesch Readability Score: 64.9).

Authorizes a pharmacist to assess, diagnose and treat certain health conditions. Includes the assessment, diagnosis and treatment of certain conditions in the practice of pharmacy. Authorizes a pharmacist to prescribe and dispense certain drugs and devices. Directs the Public Health and Pharmacy Formulary Advisory Committee to develop a list of health conditions that a pharmacist may assess, diagnose and treat.

Takes effect on the 91st day following adjournment sine die.

## A BILL FOR AN ACT

Relating to pharmacy; creating new provisions; amending ORS 689.005, 689.645 and 689.649; and prescribing an effective date.

**Be It Enacted by the People of the State of Oregon:**

**SECTION 1. Section 2 of this 2025 Act is added to and made a part of ORS chapter 689.**

**SECTION 2. A pharmacist may assess, diagnose and provide patient care services, including treatment, for health conditions:**

**(1) That may be treated by nonprescription drugs;**

**(2) That are minor or generally self-limiting; or**

**(3) For which the Public Health and Pharmacy Formulary Advisory Committee determines that assessment, diagnosis and treatment by a pharmacist is in the best interest of the public.**

**SECTION 3.** ORS 689.005, as amended by section 5, chapter 17, Oregon Laws 2024, and section 9, chapter 70, Oregon Laws 2024, is amended to read:

689.005. As used in this chapter:

(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 by:

(a) A practitioner or the practitioner's authorized agent; or

(b) The patient or research subject at the direction of the practitioner.

(2) "Approved continuing pharmacy education program" means those seminars, classes, meetings, workshops and other educational programs on the subject of pharmacy approved by the State Board of Pharmacy.

(3) "Clinical pharmacy agreement" means an agreement between a pharmacist or pharmacy and a health care organization or a physician as defined in ORS 677.010 or a naturopathic physician as defined in ORS 685.010 that permits the pharmacist to engage in the practice of clinical pharmacy for the benefit of the patients of the health care organization, physician or naturopathic physician.

**NOTE:** Matter in **boldfaced** type in an amended section is new; matter [*italic and bracketed*] is existing law to be omitted. New sections are in **boldfaced** type.

(4) "Continuing pharmacy education" means:

(a) Professional, pharmaceutical post-graduate education in the general areas of socio-economic and legal aspects of health care;

(b) The properties and actions of drugs and dosage forms; and

(c) The etiology, characteristics and therapeutics of the disease state.

(5) "Continuing pharmacy education unit" means the unit of measurement of credits for approved continuing education courses and programs.

(6) "Deliver" or "delivery" means the actual, constructive or attempted transfer of a drug or device other than by administration from one person to another, whether or not for a consideration.

(7) "Device" means an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent or other similar or related article, including any component part or accessory, which is required under federal or state law to be prescribed by a practitioner and dispensed by a pharmacist.

(8) "Dispense" or "dispensing" means the preparation and delivery of a prescription drug pursuant to a lawful order of a practitioner in a suitable container appropriately labeled for subsequent administration to or use by a patient or other individual entitled to receive the prescription drug.

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

(10) "Drug" means:

(a) Articles recognized as drugs in the official United States Pharmacopoeia, official National Formulary, official Homeopathic Pharmacopoeia, other drug compendium or any supplement to any of them;

(b) Articles intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in a human or other animal;

(c) Articles, other than food, intended to affect the structure or any function of the body of humans or other animals; and

(d) Articles intended for use as a component of any articles specified in paragraph (a), (b) or (c) of this subsection.

(11) "Drug order" means a written order, in a hospital or other inpatient care facility, for an ultimate user of any drug or device issued and signed by a practitioner, or an order transmitted by other means of communication from a practitioner, that is immediately reduced to writing by a pharmacist, licensed nurse or other practitioner.

(12) "Drug outlet" means a pharmacy, nursing home, shelter home, convalescent home, extended care facility, drug abuse treatment center, penal institution, hospital, family planning clinic, student health center, retail store, wholesaler, manufacturer, mail-order vendor or other establishment with facilities located within or out of this state that is engaged in dispensing, delivery or distribution of drugs within this state.

(13) "Drug room" means a secure and lockable location within an inpatient care facility that does not have a licensed pharmacy.

(14) "Electronically transmitted" or "electronic transmission" means a communication sent or received through technological apparatuses, including computer terminals or other equipment or mechanisms linked by telephone or microwave relays, or similar apparatus having electrical, digital, magnetic, wireless, optical, electromagnetic or similar capabilities.

(15) "Injectable hormonal contraceptive" means a drug composed of a hormone or a combination of hormones that is approved by the United States Food and Drug Administration to prevent pregnancy and that a health care practitioner administers to the patient by injection.

(16) "Institutional drug outlet" means hospitals and inpatient care facilities where medications

are dispensed to another health care professional for administration to patients served by the hospitals or facilities.

(17) "Intern" means a person who is enrolled in or has completed a course of study at a school or college of pharmacy approved by the board and who is licensed with the board as an intern.

(18) "Internship" means a professional experiential program approved by the board under the supervision of a licensed pharmacist registered with the board as a preceptor.

(19) "Labeling" means the process of preparing and affixing of a label to any drug container exclusive, however, of the labeling by a manufacturer, packer or distributor of a nonprescription drug or commercially packaged legend drug or device.

(20) "Manufacture" means the production, preparation, propagation, compounding, conversion or processing of a device or a drug, either directly or indirectly by extraction from substances of natural origin or independently by means of chemical synthesis or by a combination of extraction and chemical synthesis and includes any packaging or repackaging of the substances or labeling or relabeling of its container, except that this term does not include the preparation or compounding of a drug by an individual for their own use or the preparation, compounding, packaging or labeling of a drug:

(a) By a practitioner as an incident to administering or dispensing of a drug in the course of professional practice; or

(b) By a practitioner or by the practitioner's authorization under supervision of the practitioner for the purpose of or as an incident to research, teaching or chemical analysis and not for sale.

(21) "Manufacturer" means a person engaged in the manufacture of drugs.

(22) "Nonprescription drug outlet" means a business or other establishment that is open to the general public for the sale or nonprofit distribution of nonprescription drugs and is registered under ORS 689.305.

(23) "Nonprescription drugs" means drugs that may be sold without a prescription and that are prepackaged for use by the consumer and labeled in accordance with the requirements of the statutes and regulations of this state and the federal government.

(24) "Person" means an individual, corporation, partnership, association or other legal entity.

(25) "Pharmacist" means an individual licensed by this state to engage in the practice of pharmacy or to engage in the practice of clinical pharmacy.

(26) "Pharmacy" means a place that meets the requirements of rules of the board, is licensed and approved by the board where the practice of pharmacy may lawfully occur and includes apothecaries, drug stores, dispensaries, hospital outpatient pharmacies, pharmacy departments and prescription laboratories but does not include a place used by a manufacturer or wholesaler.

(27) "Pharmacy technician" means a person licensed by the board who assists in the practice of pharmacy pursuant to rules of the board.

(28) "Practice of clinical pharmacy" means:

(a) The health science discipline in which, in conjunction with the patient's other practitioners, a pharmacist provides patient care to optimize medication therapy and to promote disease prevention and the patient's health and wellness;

(b) The provision of patient care services, including but not limited to post-diagnostic disease state management services; and

(c) The practice of pharmacy by a pharmacist pursuant to a clinical pharmacy agreement.

(29) "Practice of pharmacy" means:

(a) The interpretation and evaluation of prescription orders;

(b) The compounding, dispensing and labeling of drugs and devices, except labeling by a manufacturer, packer or distributor of nonprescription drugs and commercially packaged legend drugs and devices;

(c) The prescribing and administering of vaccines and immunizations and the providing of patient care services pursuant to ORS 689.645;

(d) The administering of drugs and devices to the extent permitted under ORS 689.655;

(e) The participation in drug selection and drug utilization reviews;

(f) The proper and safe storage of drugs and devices and the maintenance of proper records regarding the safe storage of drugs and devices;

(g) The responsibility for advising, where necessary or where regulated, of therapeutic values, content, hazards and use of drugs and devices;

(h) The monitoring of therapeutic response or adverse effect to drug therapy;

(i) The optimizing of drug therapy through the practice of clinical pharmacy;

(j) Patient care services, including medication therapy management and comprehensive medication review;

(k) The offering or performing of those acts, services, operations or transactions necessary in the conduct, operation, management and control of pharmacy;

(L) The prescribing and administering of injectable hormonal contraceptives and the prescribing and dispensing of self-administered hormonal contraceptives pursuant to ORS 689.689;

(m) The prescribing and dispensing of emergency refills of insulin and associated insulin-related devices and supplies pursuant to ORS 689.696;

(n) The prescribing, dispensing and administering of preexposure prophylactic antiretroviral therapies and post-exposure prophylactic antiretroviral therapies, pursuant to ORS 689.704 and rules adopted by the board under ORS 689.645 and 689.704;

(o) The delegation of tasks to other health care providers who are appropriately trained and authorized to perform the delegated tasks;

(p) The prescribing and dispensing of early refills of medication for the treatment of opioid use disorder pursuant to section 7, chapter 70, Oregon Laws 2024; [*and*]

(q) The testing for severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) and the prescribing, dispensing and administering of treatment for SARS-CoV-2 pursuant to section 4, chapter 17, Oregon Laws 2024, and rules adopted by the board pursuant to section 4, chapter 17, Oregon Laws 2024[.]**; and**

**(r) The assessment, diagnosis and provision of patient care services of health conditions pursuant to section 2 of this 2025 Act.**

(30) "Practitioner" means a person licensed and operating within the scope of such license to prescribe, dispense, conduct research with respect to or administer drugs in the course of professional practice or research:

(a) In this state; or

(b) In another state or territory of the United States if the person does not reside in Oregon and is registered under the federal Controlled Substances Act.

(31) "Preceptor" means a pharmacist or a person licensed by the board to supervise the internship training of a licensed intern.

(32) "Prescription drug" or "legend drug" means a drug that is:

(a) Required by federal law, prior to being dispensed or delivered, to be labeled with either of the following statements:

(A) "Caution: Federal law prohibits dispensing without prescription"; or

(B) "Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian"; or

(b) Required by any applicable federal or state law or regulation to be dispensed on prescription only or is restricted to use by practitioners only.

(33) "Prescription" or "prescription drug order" means a written, oral or electronically transmitted direction, given by a practitioner authorized to prescribe drugs, for the preparation and use of a drug. When the context requires, "prescription" also means the drug prepared under such written, oral or electronically transmitted direction.

(34) "Retail drug outlet" means a place used for the conduct of the retail sale, administering or dispensing or compounding of drugs or chemicals or for the administering or dispensing of prescriptions and licensed by the board as a place where the practice of pharmacy may lawfully occur.

(35) "Self-administered hormonal contraceptive" means a drug composed of a hormone or a combination of hormones that is approved by the United States Food and Drug Administration to prevent pregnancy and that the patient to whom the drug is prescribed may administer to oneself. "Self-administered hormonal contraceptive" includes, but is not limited to, hormonal contraceptive patches and hormonal contraceptive pills.

(36) "Third-party logistics provider" means an entity that:

(a) Provides or coordinates warehousing of, or other logistics services for, a product in interstate commerce on behalf of a manufacturer, wholesale distributor or dispenser of the product; and

(b) Does not take ownership of, or have responsibility to direct the sale or disposition of, the product.

(37) "Unit dose" means a sealed single-unit container so designed that the contents are administered to the patient as a single dose, direct from the container. Each unit dose container must bear a separate label, be labeled with the name and strength of the medication, the name of the manufacturer or distributor, an identifying lot number and, if applicable, the expiration date of the medication.

(38) "Wholesale distributor drug outlet" means a person, other than a manufacturer, manufacturer's colicensed partner, third-party logistics provider or repackager, as defined in 21 U.S.C. 360eee(16), that is engaged in wholesale distribution, as defined in 21 U.S.C. 353(e)(4).

**SECTION 4.** ORS 689.005, as amended by sections 5 and 6, chapter 17, Oregon Laws 2024, and section 9, chapter 70, Oregon Laws 2024, is amended to read:

689.005. As used in this chapter:

(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 by:

(a) A practitioner or the practitioner's authorized agent; or

(b) The patient or research subject at the direction of the practitioner.

(2) "Approved continuing pharmacy education program" means those seminars, classes, meetings, workshops and other educational programs on the subject of pharmacy approved by the State Board of Pharmacy.

(3) "Clinical pharmacy agreement" means an agreement between a pharmacist or pharmacy and a health care organization or a physician as defined in ORS 677.010 or a naturopathic physician as defined in ORS 685.010 that permits the pharmacist to engage in the practice of clinical pharmacy for the benefit of the patients of the health care organization, physician or naturopathic physician.

(4) "Continuing pharmacy education" means:

(a) Professional, pharmaceutical post-graduate education in the general areas of socio-economic and legal aspects of health care;

(b) The properties and actions of drugs and dosage forms; and

(c) The etiology, characteristics and therapeutics of the disease state.

(5) "Continuing pharmacy education unit" means the unit of measurement of credits for approved continuing education courses and programs.

(6) "Deliver" or "delivery" means the actual, constructive or attempted transfer of a drug or device other than by administration from one person to another, whether or not for a consideration.

(7) "Device" means an instrument, apparatus, implement, machine, contrivance, implant, in vitro reagent or other similar or related article, including any component part or accessory, which is required under federal or state law to be prescribed by a practitioner and dispensed by a pharmacist.

(8) "Dispense" or "dispensing" means the preparation and delivery of a prescription drug pursuant to a lawful order of a practitioner in a suitable container appropriately labeled for subsequent administration to or use by a patient or other individual entitled to receive the prescription drug.

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

(10) "Drug" means:

(a) Articles recognized as drugs in the official United States Pharmacopoeia, official National Formulary, official Homeopathic Pharmacopoeia, other drug compendium or any supplement to any of them;

(b) Articles intended for use in the diagnosis, cure, mitigation, treatment or prevention of disease in a human or other animal;

(c) Articles, other than food, intended to affect the structure or any function of the body of humans or other animals; and

(d) Articles intended for use as a component of any articles specified in paragraph (a), (b) or (c) of this subsection.

(11) "Drug order" means a written order, in a hospital or other inpatient care facility, for an ultimate user of any drug or device issued and signed by a practitioner, or an order transmitted by other means of communication from a practitioner, that is immediately reduced to writing by a pharmacist, licensed nurse or other practitioner.

(12) "Drug outlet" means a pharmacy, nursing home, shelter home, convalescent home, extended care facility, drug abuse treatment center, penal institution, hospital, family planning clinic, student health center, retail store, wholesaler, manufacturer, mail-order vendor or other establishment with facilities located within or out of this state that is engaged in dispensing, delivery or distribution of drugs within this state.

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(16) "Institutional drug outlet" means hospitals and inpatient care facilities where medications are dispensed to another health care professional for administration to patients served by the hos-

pitals or facilities.

(17) "Intern" means a person who is enrolled in or has completed a course of study at a school or college of pharmacy approved by the board and who is licensed with the board as an intern.

(18) "Internship" means a professional experiential program approved by the board under the supervision of a licensed pharmacist registered with the board as a preceptor.

(19) "Labeling" means the process of preparing and affixing of a label to any drug container exclusive, however, of the labeling by a manufacturer, packer or distributor of a nonprescription drug or commercially packaged legend drug or device.

(20) "Manufacture" means the production, preparation, propagation, compounding, conversion or processing of a device or a drug, either directly or indirectly by extraction from substances of natural origin or independently by means of chemical synthesis or by a combination of extraction and chemical synthesis and includes any packaging or repackaging of the substances or labeling or relabeling of its container, except that this term does not include the preparation or compounding of a drug by an individual for their own use or the preparation, compounding, packaging or labeling of a drug:

(a) By a practitioner as an incident to administering or dispensing of a drug in the course of professional practice; or

(b) By a practitioner or by the practitioner's authorization under supervision of the practitioner for the purpose of or as an incident to research, teaching or chemical analysis and not for sale.

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(26) "Pharmacy" means a place that meets the requirements of rules of the board, is licensed and approved by the board where the practice of pharmacy may lawfully occur and includes apothecaries, drug stores, dispensaries, hospital outpatient pharmacies, pharmacy departments and prescription laboratories but does not include a place used by a manufacturer or wholesaler.

(27) "Pharmacy technician" means a person licensed by the board who assists in the practice of pharmacy pursuant to rules of the board.

(28) "Practice of clinical pharmacy" means:

(a) The health science discipline in which, in conjunction with the patient's other practitioners, a pharmacist provides patient care to optimize medication therapy and to promote disease prevention and the patient's health and wellness;

(b) The provision of patient care services, including but not limited to post-diagnostic disease state management services; and

(c) The practice of pharmacy by a pharmacist pursuant to a clinical pharmacy agreement.

(29) "Practice of pharmacy" means:

(a) The interpretation and evaluation of prescription orders;

(b) The compounding, dispensing and labeling of drugs and devices, except labeling by a man-

ufacturer, packer or distributor of nonprescription drugs and commercially packaged legend drugs and devices;

(c) The prescribing and administering of vaccines and immunizations and the providing of patient care services pursuant to ORS 689.645;

(d) The administering of drugs and devices to the extent permitted under ORS 689.655;

(e) The participation in drug selection and drug utilization reviews;

(f) The proper and safe storage of drugs and devices and the maintenance of proper records regarding the safe storage of drugs and devices;

(g) The responsibility for advising, where necessary or where regulated, of therapeutic values, content, hazards and use of drugs and devices;

(h) The monitoring of therapeutic response or adverse effect to drug therapy;

(i) The optimizing of drug therapy through the practice of clinical pharmacy;

(j) Patient care services, including medication therapy management and comprehensive medication review;

(k) The offering or performing of those acts, services, operations or transactions necessary in the conduct, operation, management and control of pharmacy;

(L) The prescribing and administering of injectable hormonal contraceptives and the prescribing and dispensing of self-administered hormonal contraceptives pursuant to ORS 689.689;

(m) The prescribing and dispensing of emergency refills of insulin and associated insulin-related devices and supplies pursuant to ORS 689.696;

(n) The prescribing, dispensing and administering of preexposure prophylactic antiretroviral therapies and post-exposure prophylactic antiretroviral therapies, pursuant to ORS 689.704 and rules adopted by the board under ORS 689.645 and 689.704;

(o) The delegation of tasks to other health care providers who are appropriately trained and authorized to perform the delegated tasks; [*and*]

(p) The prescribing and dispensing of early refills of medication for the treatment of opioid use disorder pursuant to section 7, chapter 70, Oregon Laws 2024[.]**; and**

**(q) The assessment, diagnosis and provision of patient care services of health conditions pursuant to section 2 of this 2025 Act.**

(30) "Practitioner" means a person licensed and operating within the scope of such license to prescribe, dispense, conduct research with respect to or administer drugs in the course of professional practice or research:

(a) In this state; or

(b) In another state or territory of the United States if the person does not reside in Oregon and is registered under the federal Controlled Substances Act.

(31) "Preceptor" means a pharmacist or a person licensed by the board to supervise the internship training of a licensed intern.

(32) "Prescription drug" or "legend drug" means a drug that is:

(a) Required by federal law, prior to being dispensed or delivered, to be labeled with either of the following statements:

(A) "Caution: Federal law prohibits dispensing without prescription"; or

(B) "Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian"; or

(b) Required by any applicable federal or state law or regulation to be dispensed on prescription only or is restricted to use by practitioners only.

(33) "Prescription" or "prescription drug order" means a written, oral or electronically transmitted direction, given by a practitioner authorized to prescribe drugs, for the preparation and use of a drug. When the context requires, "prescription" also means the drug prepared under such written, oral or electronically transmitted direction.

(34) "Retail drug outlet" means a place used for the conduct of the retail sale, administering or dispensing or compounding of drugs or chemicals or for the administering or dispensing of prescriptions and licensed by the board as a place where the practice of pharmacy may lawfully occur.

(35) "Self-administered hormonal contraceptive" means a drug composed of a hormone or a combination of hormones that is approved by the United States Food and Drug Administration to prevent pregnancy and that the patient to whom the drug is prescribed may administer to oneself. "Self-administered hormonal contraceptive" includes, but is not limited to, hormonal contraceptive patches and hormonal contraceptive pills.

(36) "Third-party logistics provider" means an entity that:

(a) Provides or coordinates warehousing of, or other logistics services for, a product in interstate commerce on behalf of a manufacturer, wholesale distributor or dispenser of the product; and

(b) Does not take ownership of, or have responsibility to direct the sale or disposition of, the product.

(37) "Unit dose" means a sealed single-unit container so designed that the contents are administered to the patient as a single dose, direct from the container. Each unit dose container must bear a separate label, be labeled with the name and strength of the medication, the name of the manufacturer or distributor, an identifying lot number and, if applicable, the expiration date of the medication.

(38) "Wholesale distributor drug outlet" means a person, other than a manufacturer, manufacturer's colicensed partner, third-party logistics provider or repackager, as defined in 21 U.S.C. 360eee(16), that is engaged in wholesale distribution, as defined in 21 U.S.C. 353(e)(4).

**SECTION 5.** ORS 689.645 is amended to read:

689.645. (1) In accordance with rules adopted by the State Board of Pharmacy under ORS 689.205:

(a) A pharmacist, or a pharmacy technician under the supervision of a pharmacist, may administer vaccines:

(A) To persons who are seven years of age or older;

(B) If authorized by the Governor or the Director of the Oregon Department of Emergency Management under ORS 433.441 or the Public Health Director under ORS 433.443 or 433.444, to a person three years of age or older; or

(C) To persons who are six months of age or older if the vaccine administered is an influenza vaccine.

(b) A pharmacist may, pursuant to a statewide drug therapy management protocol developed by the Public Health and Pharmacy Formulary Advisory Committee convened under ORS 689.649 and adopted by rule of the board, provide approved patient care services including smoking cessation therapy and travel health services.

(c) A pharmacist may, using a form prescribed by the board, submit a concept for the development of a protocol, other than the protocols pharmacists may establish under subsection (5) of this section, to the committee for consideration by the committee and recommendation to the board for adoption by rule of the board.

(d) A pharmacist may prescribe and dispense a drug or device included on the formulary estab-

lished under subsection (6) of this section if the prescription and dispensation is**:**

**(A)** Pursuant to a diagnosis by a health care practitioner who has prescriptive authority and is qualified to make the diagnosis[.]**; or**

**(B) Made within the pharmacist's scope of practice.**

(2) The board may adopt rules allowing a pharmacist to prescribe vaccines, provide patient care services and submit protocol concepts under subsection (1) of this section. The rules related to the prescription of vaccines may be only as broad as necessary to enable pharmacists to enroll and participate in the Vaccines for Children Program administered by the Centers for Disease Control and Prevention.

(3) The board is authorized to issue, to pharmacists and pharmacy technicians who have completed training accredited by the Centers for Disease Control and Prevention, the Accreditation Council for Pharmacy Education or a similar health authority or professional body, certificates of special competency in the prescription or administration of vaccines.

(4) The board shall adopt rules relating to the reporting of the prescription and administration of vaccines to a patient's primary health care provider and to the Oregon Health Authority.

(5) The board shall adopt rules requiring pharmacists to establish protocols for the prescription and administration of vaccines and the provision of patient care services under subsection (1) of this section.

(6)(a) The board shall establish by rule a formulary of drugs and devices, as recommended by the committee, that a pharmacist may prescribe and dispense to a patient [*pursuant to a diagnosis by a health care practitioner who has prescriptive authority and who is qualified to make the diagnosis*] **under subsection (1)(d) of this section**.

(b) The formulary may include post-diagnostic drugs and devices such as diabetic testing supplies, emergency refills of insulin, albuterol inhalers, epinephrine autoinjectors, smoking cessation aids, discharge medications for transitions of care, rapid strep tests and spacers.

**SECTION 6.** ORS 689.649 is amended to read:

689.649. (1) The State Board of Pharmacy shall convene a Public Health and Pharmacy Formulary Advisory Committee consisting of seven members, appointed by the Governor, for the purpose of advising the board in promulgating rules under ORS 689.645. The committee shall consist of:

(a) Two physicians licensed to practice medicine under ORS 677.100 to 677.228;

(b) Two advanced practice registered nurses who have prescriptive authority and who are licensed by the Oregon State Board of Nursing; and

(c) Three pharmacists licensed by the State Board of Pharmacy, at least one of whom is employed as a community pharmacist and one of whom is employed as a health system pharmacist.

(2) The Oregon Medical Board, the Oregon State Board of Nursing and the State Board of Pharmacy may each submit to the Governor a list of up to three names of individuals to be considered for membership for each of the vacancies required to be filled by licensees of each board.

(3) The term of each member of the committee is two years. A member whose term has expired shall continue to serve until a successor is appointed. If a vacancy occurs, a person who is a representative of the same state agency as the departing member shall serve for the remainder of the term.

(4) The committee shall elect one of its members to serve as chairperson.

(5) Members of the committee are entitled to compensation and expenses as provided in ORS 292.495, to be paid by the State Board of Pharmacy.

(6) A member of the committee who fails to attend two consecutive meetings of the committee shall be removed from the committee unless the failure to attend was because of a serious health condition of the member or a family member of the member.

(7) The committee shall recommend to the State Board of Pharmacy for adoption by rule of the board a formulary of drugs and devices that a pharmacist may prescribe and dispense to a patient pursuant to a diagnosis by a health care practitioner qualified to make the diagnosis. The committee shall periodically review the formulary and recommend the revisions to the board for adoption by rule.

(8) A pharmacist may request that the committee add a drug or device to the formulary by submitting to the committee a request form prescribed by the State Board of Pharmacy. The addition to the formulary of a drug or device under this subsection shall be considered a revision to the formulary that the committee may recommend to the board for adoption by rule.

**(9) The committee may develop and maintain a list of health conditions that the committee determines to be within the scope of practice of a pharmacist to assess, diagnose and treat. The conditions included on the list under this subsection must be minor or generally self-limiting, or within the best interests of the public to be assessed, diagnosed and treated by a pharmacist. The committee may review and revise the list as the committee determines necessary and shall make the list publicly available.**

**SECTION 7. (1) Section 2 of this 2025 Act and the amendments to ORS 689.005, 689.645 and 689.649 by sections 3 to 6 of this 2025 Act become operative on January 1, 2026.**

**(2) The State Board of Pharmacy and the Public Health and Pharmacy Formulary Advisory Committee may take any action before the operative date specified in subsection (1) of this section that is necessary to enable the board and the committee to exercise, on and after the operative date specified in subsection (1) of this section, all of the duties, functions and powers conferred on the board and the committee by section 2 of this 2025 Act and the amendments to ORS 689.005, 689.645 and 689.649 by sections 3 to 6 of this 2025 Act.**

**SECTION 8. This 2025 Act takes effect on the 91st day after the date on which the 2025 regular session of the Eighty-third Legislative Assembly adjourns sine die.**