SHB 1047 - H AMD **817**
By Representatives Peterson, Schmick

ADOPTED 02/09/2018

Strike everything after the enacting clause and insert the following:

"NEW SECTION. **Sec. 1.** LEGISLATIVE FINDINGS. (1) Abuse, fatal overdoses, and poisonings from prescription and over-the-counter medicines used in the home have emerged as an epidemic in recent years. Poisoning is the leading cause of unintentional injury-related death in Washington, and more than ninety percent of poisoning deaths are due to drug overdoses. Poisoning by prescription and over-the-counter medicines is also one of the most common means of suicide and suicide attempts, with poisonings involved in more than twenty-eight thousand suicide attempts between 2004 and 2013.

(2) Home medicine cabinets are the most common source of prescription drugs that are diverted and misused. Studies find about seventy percent of those who abuse prescription medicines obtain the drugs from family members or friends, usually for free. People who are addicted to heroin often first abused prescription opiate medicines. Unused, unwanted, and expired medicines that accumulate in homes increase risks of drug abuse, overdoses, and preventable poisonings.

(3) A safe system for the collection and disposal of unused, unwanted, and expired medicines is a key element of a comprehensive strategy to prevent prescription drug abuse, but disposing of medicines by flushing them down the toilet or placing them in the garbage can contaminate groundwater and other bodies of water, contributing to long-term harm to the environment and animal life.

(4) The legislature therefore finds that it is in the interest of public health to establish a single, uniform, statewide system of regulation for safe and secure collection and disposal of medicines through a uniform drug "take-back" program operated and funded by drug manufacturers.

NEW SECTION. **Sec. 2.** DEFINITIONS. The definitions in this section apply throughout this chapter unless the context clearly requires otherwise.

(1) "Administer" means the direct application of a legend drug whether by injection, inhalation, ingestion, or any other means, to the body of the patient or research subject by:

(a) A practitioner; or

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

(2) "Authorized collector" means any of the following persons or entities that have entered into an agreement with a program operator to collect covered drugs:

(a) A person or entity that is registered with the United States drug enforcement administration and that qualifies under federal law to modify its registration to collect controlled substances for the purpose of destruction;

(b) A law enforcement agency; or

(c) An entity authorized by the department to provide an alternative collection mechanism for certain covered drugs that are not controlled substances, as defined in RCW 69.50.101.

(3) "Collection site" means the location where an authorized collector operates a secure collection receptacle for collecting covered drugs.

(4)(a) "Covered drug" means a drug from a covered entity that the covered entity no longer wants and that the covered entity has abandoned or discarded or intends to abandon or discard. "Covered drug" includes legend drugs and nonlegend drugs, brand name and generic drugs, drugs for veterinary use for household pets, and drugs in medical devices and combination products.

(b) "Covered drug" does not include:

(i) Vitamins, minerals, or supplements;

(ii) Herbal-based remedies and homeopathic drugs, products, or remedies;

(iii) Controlled substances contained in schedule I of the uniform controlled substances act, chapter 69.50 RCW;

(iv) Cosmetics, shampoos, sunscreens, lip balm, toothpaste, antiperspirants, or other personal care products that are regulated as both cosmetics and nonprescription drugs under the federal food, drug, and cosmetic act, 21 U.S.C. Sec. 301 et seq.;

(v) Drugs for which manufacturers provide a pharmaceutical product stewardship or drug take-back program as part of a federal food and drug administration managed risk evaluation and mitigation strategy under 21 U.S.C. Sec. 355-1;

(vi) Biological drug products, as defined by 21 C.F.R. 600.3 (h) as it exists on the effective date of this section, for which manufacturers provide a pharmaceutical product stewardship or drug take-back program and who provide the department with a report describing the program, including how the drug product is collected and safely disposed and how patients are made aware of the drug take-back program, and who updates the department on changes that substantially alter their drug take-back program;

(vii) Drugs that are administered in a clinical setting;

(viii) Emptied injector products or emptied medical devices and their component parts or accessories;

(ix) Exposed needles or sharps, or used drug products that are medical wastes; or

(x) Pet pesticide products contained in pet collars, powders, shampoos, topical applications, or other forms.

(5) "Covered entity" means a state resident or other nonbusiness entity and includes an ultimate user, as defined by regulations adopted by the United States drug enforcement administration. "Covered entity" does not include a business generator of pharmaceutical waste, such as a hospital, clinic, health care provider's office, veterinary clinic, pharmacy, or law enforcement agency.

(6) "Covered manufacturer" means a person, corporation, or other entity engaged in the manufacture of covered drugs sold in or into Washington state. "Covered manufacturer" does not include a retail pharmacy that sells a drug under the retail pharmacy's store label if the manufacturer of the drug is identified under section 4 of this act.

(7) "Department" means the department of health.

(8)(a) "Drug" means:

(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;

(b) Substances intended for use in the diagnosis, cure, mitigation, treatment, or prevention of disease in human beings or animals;

(c) Substances other than food, minerals, or vitamins that are intended to affect the structure or any function of the body of human beings or animals; and

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

(9) "Drug take-back organization" means an organization designated by a manufacturer or group of manufacturers to act as an agent on behalf of each manufacturer to develop and implement a drug take-back program.

(10) "Drug take-back program" or "program" means a program implemented by a program operator for the collection, transportation, and disposal of covered drugs.

(11) "Drug wholesaler" means an entity licensed as a wholesaler under chapter 18.64 RCW.

(12) "Generic drug" means a drug that is chemically identical or bioequivalent to a brand name drug in dosage form, safety, strength, route of administration, quality, performance characteristics, and intended use. The inactive ingredients in a generic drug need not be identical to the inactive ingredients in the chemically identical or bioequivalent brand name drug.

(13) "Legend drug" means a drug, including a controlled substance under chapter 69.50 RCW, that is required by any applicable federal or state law or regulation to be dispensed by prescription only or that is restricted to use by practitioners only.

(14) "Mail-back distribution location" means a facility, such as a town hall or library, that offers prepaid, preaddressed mailing envelopes to covered entities.

(15) "Mail-back program" means a method of collecting covered drugs from covered entities by using prepaid, preaddressed mailing envelopes.

(16) "Manufacture" has the same meaning as in RCW 18.64.011.

(17) "Nonlegend drug" means a drug that may be lawfully sold without a prescription.

(18) "Pharmacy" means a place licensed as a pharmacy under chapter 18.64 RCW.

(19) "Program operator" means a drug take-back organization, covered manufacturer, or group of covered manufacturers that

implements or intends to implement a drug take-back program approved by the department.

(20) "Retail pharmacy" means a place licensed as a pharmacy under chapter 18.64 RCW for the retail sale and dispensing of drugs.

(21) "Secretary" means the secretary of health.

NEW SECTION. **Sec. 3.** REQUIREMENT TO PARTICIPATE IN A DRUG TAKE-BACK PROGRAM. A covered manufacturer must establish and implement a drug take-back program that complies with the requirements of this chapter. A manufacturer that becomes a covered manufacturer after the effective date of this section must, no later than six months after the date on which the manufacturer became a covered manufacturer, participate in an approved drug take-back program or establish and implement a drug take-back program that complies with the requirements of this chapter. A covered manufacturer may establish and implement a drug take-back program independently, as part of a group of covered manufacturers, or through membership in a drug take-back organization.

NEW SECTION. **Sec. 4.** IDENTIFICATION OF COVERED MANUFACTURERS. (1) No later than ninety days after the effective date of this section, a drug wholesaler that sells a drug in or into Washington must provide a list of drug manufacturers to the department in a form agreed upon with the department. A drug wholesaler must provide an updated list to the department on January 15th of each year.

(2) No later than ninety days after the effective date of this section, a retail pharmacy must provide written notification to the department identifying the drug manufacturer from which the retail pharmacy obtains a drug that the retail pharmacy sells under its store label.

(3) A person or entity that receives a letter of inquiry from the department regarding whether or not it is a covered manufacturer under this chapter shall respond in writing no later than sixty days after receipt of the letter. If the person or entity does not believe it is a covered manufacturer for purposes of this chapter, it shall: (a) State the basis for the belief; (b) provide a list of any drugs it sells, distributes, repackages, or otherwise offers for sale within the state; and (c) identify the name and contact information of the manufacturer of the drugs identified under (b) of this subsection.

NEW SECTION. **Sec. 5.** DRUG TAKE-BACK PROGRAM APPROVAL. (1) By July 1, 2019, a program operator must submit a proposal for the establishment and implementation of a drug take-back program to the department for approval. The department shall approve a proposed program if the applicant submits a completed application, the proposed program meets the requirements of subsection (2) of this section, and the applicant pays the appropriate fee established by the department under section 12 of this act.

(2) To be approved by the department, a proposed drug take-back program must:

(a) Identify and provide contact information for the program operator and each participating covered manufacturer;

(b) Identify and provide contact information for the authorized collectors for the proposed program, as well as the reasons for excluding any potential authorized collectors from participation in the program;

(c) Provide for a collection system that complies with section 6 of this act;

(d) Provide for a handling and disposal system that complies with section 8 of this act;

(e) Identify any transporters and waste disposal facilities that the program will use;

(f) Adopt policies and procedures to be followed by persons handling covered drugs collected under the program to ensure safety, security, and compliance with regulations adopted by the United States drug enforcement administration, as well as any applicable laws;

(g) Ensure the security of patient information on drug packaging during collection, transportation, recycling, and disposal;

(h) Promote the program by providing consumers, pharmacies, and other entities with educational and informational materials as required by section 7 of this act;

(i) Demonstrate adequate funding for all administrative and operational costs of the drug take-back program, with costs apportioned among participating covered manufacturers;

(j) Set long-term and short-term goals with respect to collection amounts and public awareness; and

(k) Consider: (i) The use of existing providers of pharmaceutical waste transportation and disposal services; (ii) separation of

covered drugs from packaging to reduce transportation and disposal costs; and (iii) recycling of drug packaging.

(3)(a) No later than one hundred twenty days after receipt of a drug take-back program proposal, the department shall either approve or reject the proposal in writing to the applicant. The department may extend the deadline for approval or rejection of a proposal for good cause. If the department rejects the proposal, it shall provide the reason for rejection.

(b) No later than ninety days after receipt of a notice of rejection under (a) of this subsection, the applicant shall submit a revised proposal to the department. The department shall either approve or reject the revised proposal in writing to the applicant within ninety days after receipt of the revised proposal, including the reason for rejection, if applicable.

(c) If the department rejects a revised proposal, the department may:

(i) Require the program operator to submit a further revised proposal;

(ii) Develop and impose changes to some or all of the revised proposal to address deficiencies;

(iii) Require the covered manufacturer or covered manufacturers that proposed the rejected revised proposal to participate in a previously approved drug take-back program; or

(iv) Find the covered manufacturer out of compliance with the requirements of this chapter and take enforcement action as provided in section 11 of this act.

(4) The program operator must initiate operation of an approved drug take-back program no later than one hundred eighty days after approval of the proposal by the department.

(5)(a) Proposed changes to an approved drug take-back program that substantially alter program operations must have prior written approval of the department. A program operator must submit to the department such a proposed change in writing at least fifteen days before the change is scheduled to occur. Changes requiring prior approval of the department include changes to participating covered manufacturers, collection methods, achievement of the service convenience goal described in section 6 of this act, policies and procedures for handling covered drugs, education and promotion methods, and selection of disposal facilities.

(b) For changes to a drug take-back program that do not substantially alter program operations, a program operator must notify the department at least seven days before implementing the change. Changes that do not substantially alter program operations include changes to collection site locations, methods for scheduling and locating periodic collection events, and methods for distributing prepaid, preaddressed mailers.

(c) A program operator must notify the department of any changes to the official point of contact for the program no later than fifteen days after the change. A program operator must notify the department of any changes in ownership or contact information for participating covered manufacturers no later than ninety days after such change.

(6) No later than four years after a drug take-back program initiates operations, and every four years thereafter, the program operator must submit an updated proposal to the department describing any substantive changes to program elements described in subsection (2) of this section. The department shall approve or reject the updated proposal using the process described in subsection (3) of this section.

(7) The department shall make all proposals submitted under this section available to the public and shall provide an opportunity for written public comment on each proposal.

NEW SECTION. **Sec. 6.** COLLECTION SYSTEM. (1)(a) At least one hundred twenty days prior to submitting a proposal under section 5 of this act, a program operator must notify potential authorized collectors of the opportunity to serve as an authorized collector for the proposed drug take-back program. A program operator must commence good faith negotiations with a potential authorized collector no later than thirty days after the potential authorized collector expresses interest in participating in a proposed program.

(b) A person or entity may serve as an authorized collector for a drug take-back program voluntarily or in exchange for compensation, but nothing in this chapter requires a person or entity to serve as an authorized collector.

(c) A drug take-back program must include as an authorized collector any retail pharmacy, hospital or clinic with an on-site pharmacy, or law enforcement agency that offers to participate in the program without compensation and meets the requirements of subsection

(2) of this section. Such a pharmacy, hospital, clinic, or law enforcement agency must be included as an authorized collector in the program no later than ninety days after receiving the offer to participate.

(d) A drug take-back program may also locate collection sites at:

(i) A long-term care facility where a pharmacy, or a hospital or clinic with an on-site pharmacy, operates a secure collection receptacle;

(ii) A substance use disorder treatment program, as defined in RCW 71.24.025; or

(iii) Any other authorized collector willing to participate as a collection site and able to meet the requirements of subsection (2) of this section.

(2)(a) A collection site must accept all covered drugs from covered entities during the hours that the authorized collector is normally open for business with the public.

(b) A collection site located at a long-term care facility may only accept covered drugs that are in the possession of individuals who reside or have resided at the facility.

(c) A collection site must use secure collection receptacles in compliance with state and federal law, including any applicable on-site storage and collection standards adopted by rule pursuant to chapter 70.95 or 70.105 RCW and United States drug enforcement administration regulations. The program operator must provide a service schedule that meets the needs of each collection site to ensure that each secure collection receptacle is serviced as often as necessary to avoid reaching capacity and that collected covered drugs are transported to final disposal in a timely manner, including a process for additional prompt collection service upon notification from the collection site. Secure collection receptacle signage must prominently display a toll-free telephone number and web site for the program so that members of the public may provide feedback on collection activities.

(d) An authorized collector must comply with applicable provisions of chapters 70.95 and 70.105 RCW, including rules adopted pursuant to those chapters that establish collection and transportation standards, and federal laws and regulations governing the handling of covered drugs, including United States drug enforcement administration regulations.

(3)(a) A drug take-back program's collection system must be safe, secure, and convenient on an ongoing, year-round basis and must provide equitable and reasonably convenient access for residents across the state.

(b) In establishing and operating a collection system, a program operator must give preference to locating collection sites at retail pharmacies, hospitals or clinics with on-site pharmacies, and law enforcement agencies.

(c)(i) Each population center must have a minimum of one collection site, plus one additional collection site for every fifty thousand residents of the city or town located within the population center. Collection sites must be geographically distributed to provide reasonably convenient and equitable access to all residents of the population center.

(ii) On islands and in areas outside of population centers, a collection site must be located at the site of each potential authorized collector that is regularly open to the public, unless the program operator demonstrates to the satisfaction of the department that a potential authorized collector is unqualified or unwilling to participate in the drug take-back program, in accordance with the requirements of subsection (1) of this section.

(iii) For purposes of this section, "population center" means a city or town and the unincorporated area within a ten-mile radius from the center of the city or town.

(d) A program operator must establish mail-back distribution locations or hold periodic collection events to supplement service to any area of the state that is underserved by collection sites, as determined by the department, in consultation with the local health jurisdiction. The program operator, in consultation with the department, local law enforcement, the local health jurisdiction, and the local community, must determine the number and locations of mail-back distribution locations or the frequency and location of these collections events, to be held at least twice a year, unless otherwise determined through consultation with the local community. The program must arrange any periodic collection events in advance with local law enforcement agencies and conduct periodic collection events in compliance with United States drug enforcement administration regulations and protocols and applicable state laws.

(e) Upon request, a drug take-back program must provide a mail-back program free of charge to covered entities and to retail

pharmacies that offer to distribute prepaid, preaddressed mailing envelopes for the drug take-back program. A drug take-back program must permit covered entities to request prepaid, preaddressed mailing envelopes through the program's web site, the program's toll-free telephone number, and a request to a pharmacist at a retail pharmacy distributing the program's mailing envelopes.

(f) The program operator must provide alternative collection methods for any covered drugs, other than controlled substances, that cannot be accepted or commingled with other covered drugs in secure collection receptacles, through a mail-back program, or at periodic collection events, to the extent permissible under applicable state and federal laws. The department shall review and approve of any alternative collection methods prior to their implementation.

NEW SECTION. **Sec. 7.** DRUG TAKE-BACK PROGRAM PROMOTION. (1) A drug take-back program must develop and provide a system of promotion, education, and public outreach about the safe storage and secure collection of covered drugs. This system may include signage, written materials to be provided at the time of purchase or delivery of covered drugs, and advertising or other promotional materials. At a minimum, each program must:

(a) Promote the safe storage of legend drugs and nonlegend drugs by residents before secure disposal through a drug take-back program;

(b) Discourage residents from disposing of covered drugs in solid waste collection, sewer, or septic systems;

(c) Promote the use of the drug take-back program so that where and how to return covered drugs is widely understood by residents, pharmacists, retail pharmacies, health care facilities and providers, veterinarians, and veterinary hospitals;

(d) Establish a toll-free telephone number and web site publicizing collection options and collection sites and discouraging improper disposal practices for covered drugs, such as flushing them or placing them in the garbage;

(e) Prepare educational and outreach materials that: Promote safe storage of covered drugs; discourage the disposal of covered drugs in solid waste collection, sewer, or septic systems; and describe how to return covered drugs to the drug take-back program. The materials must use plain language and explanatory images to make collection services and discouraged disposal practices readily understandable to all residents, including residents with limited English proficiency;

(f) Disseminate the educational and outreach materials described in (e) of this subsection to pharmacies, health care facilities, and other interested parties for dissemination to covered entities;

(g) Work with authorized collectors to develop a readily recognizable, consistent design of collection receptacles, as well as clear, standardized instructions for covered entities on the use of collection receptacles. The department may provide guidance to program operators on the development of the instructions and design; and

(h) Annually report on its promotion, outreach, and public education activities in its annual report required by section 10 of this act.

(2) If more than one drug take-back program is approved by the department, the programs must coordinate their promotional activities to ensure that all state residents can easily identify, understand, and access the collection services provided by any drug take-back program. Coordination efforts must include providing residents with a single toll-free telephone number and single web site to access information about collection services for every approved program.

(3) Pharmacies and other entities that sell medication in the state are encouraged to promote secure disposal of covered drugs through the use of one or more approved drug take-back programs. Upon request, a pharmacy must provide materials explaining the use of approved drug take-back programs to its customers. The program operator must provide pharmacies with these materials upon request and at no cost to the pharmacy.

(4) The department, the health care authority, the department of social and health services, the department of ecology, and any other state agency that is responsible for health, solid waste management, and wastewater treatment shall, through their standard educational methods, promote safe storage of prescription and nonprescription drugs by covered entities, secure disposal of covered drugs through a drug take-back program, and the toll-free telephone number and web site for approved drug take-back programs. Local health jurisdictions and local government agencies are encouraged to promote approved drug take-back programs.

(5) The department:

(a) Shall conduct a survey of covered entities and a survey of pharmacists, health care providers, and veterinarians who interact with covered entities on the use of medicines after the first full

year of operation of the drug take-back program, and again every two years thereafter. Survey questions must: Measure consumer awareness of the drug take-back program; assess the extent to which collection sites and other collection methods are convenient and easy to use; assess knowledge and attitudes about risks of abuse, poisonings, and overdoses from drugs used in the home; and assess covered entities' practices with respect to unused, unwanted, or expired drugs, both currently and prior to implementation of the drug take-back program; and

(b) May, upon review of results of public awareness surveys, direct a program operator for an approved drug take-back program to modify the program's promotion and outreach activities to better achieve widespread awareness among Washington state residents and health care professionals about where and how to return covered drugs to the drug take-back program.

NEW SECTION. **Sec. 8.** DISPOSAL AND HANDLING OF COVERED DRUGS. (1) Covered drugs collected under a drug take-back program must be disposed of at a permitted hazardous waste disposal facility that meets the requirements of 40 C.F.R. parts 264 and 265, as they exist on the effective date of this section.

(2) If use of a hazardous waste disposal facility described in subsection (1) of this section is unfeasible based on cost, logistics, or other considerations, the department, in consultation with the department of ecology, may grant approval for a program operator to dispose of some or all collected covered drugs at a permitted large municipal waste combustor facility that meets the requirements of 40 C.F.R. parts 60 and 62, as they exist on the effective date of this section.

(3) A program operator may petition the department for approval to use final disposal technologies or processes that provide superior environmental and human health protection than that provided by the technologies described in subsections (1) and (2) of this section, or equivalent protection at less cost. In reviewing a petition under this subsection, the department shall take into consideration regulations or guidance issued by the United States environmental protection agency on the disposal of pharmaceutical waste. The department, in consultation with the department of ecology, shall approve a disposal petition under this section if the disposal

technology or processes described in the petition provides equivalent or superior protection in each of the following areas:

(a) Monitoring of any emissions or waste;

(b) Worker health and safety;

(c) Air, water, or land emissions contributing to persistent, bioaccumulative, and toxic pollution; and

(d) Overall impact to the environment and human health.

(4) If a drug take-back program encounters a safety or security problem during collection, transportation, or disposal of covered drugs, the program operator must notify the department as soon as practicable after encountering the problem.

NEW SECTION. **Sec. 9.** PROGRAM FUNDING. (1) A covered manufacturer or group of covered manufacturers must pay all administrative and operational costs associated with establishing and implementing the drug take-back program in which they participate. Such administrative and operational costs include, but are not limited to: Collection and transportation supplies for each collection site; purchase of secure collection receptacles for each collection site; ongoing maintenance or replacement of secure collection receptacles when requested by authorized collectors; prepaid, preaddressed mailers; compensation of authorized collectors, if applicable; operation of periodic collection events, including the cost of law enforcement staff time; transportation of all collected covered drugs to final disposal; environmentally sound disposal of all collected covered drugs in compliance with section 8 of this act; and program promotion and outreach.

(2) A program operator, covered manufacturer, authorized collector, or other person may not charge:

(a) A specific point-of-sale fee to consumers to recoup the costs of a drug take-back program; or

(b) A specific point-of-collection fee at the time covered drugs are collected from covered entities.

NEW SECTION. **Sec. 10.** ANNUAL PROGRAM REPORT. (1) By July 1st after the first full year of implementation, and each July 1st thereafter, a program operator must submit to the department a report describing implementation of the drug take-back program during the previous calendar year. The report must include:

(a) A list of covered manufacturers participating in the drug take-back program;

(b) The amount, by weight, of covered drugs collected, including the amount by weight from each collection method used;

(c) The following details regarding the program's collection system: A list of collection sites with addresses; the number of mailers provided; locations where mailers were provided, if applicable; dates and locations of collection events held, if applicable; and the transporters and disposal facility or facilities used;

(d) Whether any safety or security problems occurred during collection, transportation, or disposal of covered drugs, and if so, completed and anticipated changes to policies, procedures, or tracking mechanisms to address the problem and improve safety and security;

(e) A description of the public education, outreach, and evaluation activities implemented;

(f) A description of how collected packaging was recycled to the extent feasible;

(g) A summary of the program's goals for collection amounts and public awareness, the degree of success in meeting those goals, and if any goals have not been met, what effort will be made to achieve those goals the following year; and

(h) The program's annual expenditures, itemized by program category.

(2) Within thirty days after each annual period of operation of an approved drug take-back program, the program operator shall submit an annual collection amount report to the department that provides the total amount, by weight, of covered drugs collected from each collection site during the prior year.

(3) The department shall make reports submitted under this section available to the public through the internet.

NEW SECTION. **Sec. 11.** ENFORCEMENT AND PENALTIES. (1) The department may audit or inspect the activities and records of a drug take-back program to determine compliance with this chapter or investigate a complaint.

(2)(a) The department shall send a written notice to a covered manufacturer that fails to participate in a drug take-back program as

required by this chapter. The notice must provide a warning regarding the penalties for violation of this chapter.

(b) A covered manufacturer that receives a notice under this subsection (2) may be assessed a penalty if, sixty days after receipt of the notice, the covered manufacturer continues to sell a covered drug in or into the state without participating in a drug take-back program approved under this chapter.

(3)(a) The department may send a program operator a written notice warning of the penalties for noncompliance with this chapter if it determines that the program operator's drug take-back program is in violation of this chapter or does not conform to the proposal approved by the department. The department may assess a penalty on the program operator and participating covered manufacturers if the program does not come into compliance by thirty days after receipt of the notice.

(b) The department may immediately suspend operation of a drug take-back program and assess a penalty if it determines that the program is in violation of this chapter and the violation creates a condition that, in the judgment of the department, constitutes an immediate hazard to the public or the environment.

(4)(a) The department shall send a written notice to a drug wholesaler or a retail pharmacy that fails to provide a list of drug manufacturers to the department as required by section 4 of this act. The notice must provide a warning regarding the penalties for violation of this chapter.

(b) A drug wholesaler or retail pharmacy that receives a notice under this subsection may be assessed a penalty if, sixty days after receipt of the notice, the drug wholesaler or retail pharmacy fails to provide a list of drug manufacturers.

(5) In enforcing the requirements of this chapter, the department:

(a) May require an informal administrative conference;

(b) May require a person or entity to engage in or refrain from engaging in certain activities pertaining to this chapter;

(c) May, in accordance with RCW 43.70.095, assess a civil fine of up to two thousand dollars. Each day upon which a violation occurs or is permitted to continue constitutes a separate violation. In determining the appropriate amount of the fine, the department shall consider the extent of harm caused by the violation, the nature and persistence of the violation, the frequency of past violations, any

action taken to mitigate the violation, and the financial burden to the entity in violation; and

(d) May not prohibit a covered manufacturer from selling a drug in or into the state of Washington.

NEW SECTION. **Sec. 12.** DEPARTMENT FEE. (1)(a) By July 1, 2019, the department shall: Determine its costs for the administration, oversight, and enforcement of the requirements of this chapter, including the survey required under section 20 of this act; pursuant to RCW 43.70.250, set fees at a level sufficient to recover the costs associated with administration, oversight, and enforcement; and adopt rules establishing requirements for program operator proposals.

(b) The department shall not impose any fees in excess of its actual administrative, oversight, and enforcement costs. The fees collected from each program operator in calendar year 2020 and any subsequent year may not exceed ten percent of the program's annual expenditures as reported to the department in the annual report required by section 10 of this act and determined by the department.

(c) Adjustments to the department's fees may be made annually and shall not exceed actual administration, oversight, and enforcement costs. Adjustments for inflation may not exceed the percentage change in the consumer price index for all urban consumers in the United States as calculated by the United States department of labor as averaged by city for the twelve-month period ending with June of the previous year.

(d) The department shall collect fees from each program operator by October 1, 2019, and annually thereafter.

(2) All fees collected under this section must be deposited in the secure drug take-back program account established in section 13 of this act.

NEW SECTION. **Sec. 13.** SECURE DRUG TAKE-BACK PROGRAM ACCOUNT. The secure drug take-back program account is created in the state treasury. All receipts received by the department under this chapter must be deposited in the account. Moneys in the account may be spent only after appropriation. Expenditures from the account may be used by the department only for administering and enforcing this chapter.

NEW SECTION. **Sec. 14.** ANTITRUST IMMUNITY. The activities authorized by this chapter require collaboration among covered

manufacturers. These activities will enable safe and secure collection and disposal of covered drugs in Washington state and are therefore in the best interest of the public. The benefits of collaboration, together with active state supervision, outweigh potential adverse impacts. Therefore, the legislature intends to exempt from state antitrust laws, and provide immunity through the state action doctrine from federal antitrust laws, activities that are undertaken, reviewed, and approved by the department pursuant to this chapter that might otherwise be constrained by such laws. The legislature does not intend and does not authorize any person or entity to engage in activities not provided for by this chapter, and the legislature neither exempts nor provides immunity for such activities.

NEW SECTION. **Sec. 15.** FEDERAL LAW. This chapter is void if a federal law, or a combination of federal laws, takes effect that establishes a national program for the collection of covered drugs that substantially meets the intent of this chapter, including the creation of a funding mechanism for collection, transportation, and proper disposal of all covered drugs in the United States.

NEW SECTION. **Sec. 16.** LOCAL LAWS. (1)(a) For a period of twelve months after a drug take-back program approved under section 5 of this act begins operating, a county may enforce a grandfathered ordinance. During that twelve-month period, if a county determines that a covered manufacturer is in compliance with its grandfathered ordinance, the department shall find the covered manufacturer in compliance with the requirements of this chapter with respect to that county.

(b) In any county enforcing a grandfathered ordinance as described in (a) of this subsection, the program operator of an approved drug take-back program must work with the county and the department to incorporate the local program into the approved drug take-back program on or before the end of the twelve-month period.

(2) After the effective date of this section, a political subdivision may not enact or enforce a local ordinance that requires a retail pharmacy, clinic, hospital, or local law enforcement agency to provide for collection and disposal of covered drugs from covered entities.

(3) At the end of the twelve-month period provided in subsection (1) of this section, this chapter preempts all laws enacted by a county, city, town, or other political subdivision of the state regarding a drug take-back program for the collection, transportation, and disposal of covered drugs, or promotion, education, and public outreach relating to such a program.

(4) For purposes of this section, "grandfathered ordinance" means a pharmaceutical product stewardship or drug take-back ordinance that: (a) Is in effect on the effective date of this section; and (b) the department determines meets or exceeds the requirements of this chapter with respect to safe and secure collection and disposal of unwanted medicines from residents, including the types of drugs covered by the program, the convenience of the collection system for residents, and required promotion of the program.

NEW SECTION. **Sec. 17.** PUBLIC DISCLOSURE. Proprietary information submitted to the department under this chapter is exempt from public disclosure under RCW 42.56.270. The department may use and disclose such information in summary or aggregated form that does not directly or indirectly identify financial, production, or sales data of an individual covered manufacturer or drug take-back organization.

NEW SECTION. **Sec. 18.** RULE MAKING. The department shall adopt any rules necessary to implement and enforce this chapter.

NEW SECTION. **Sec. 19.** REPORT TO LEGISLATURE. (1) No later than thirty days after the department first approves a drug take-back program under section 5 of this act, the department shall submit an update to the legislature describing rules adopted under this chapter and the approved drug take-back program.

(2) By November 15th after the first full year of operation of an approved drug take-back program and biennially thereafter, the department shall submit a report to the legislature. The report must:

(a) Describe the status of approved drug take-back programs;

(b) Evaluate the secure medicine collection and disposal system and the program promotion, education, and public outreach requirements established by this chapter;

(c) Evaluate, in conjunction with an academic institution that is not an agency of the state and is qualified to conduct and evaluate

research relating to prescription and nonprescription drug use and abuse and environmental impact, to the extent feasible, the impact of approved drug take-back programs on: Awareness and compliance of residents with safe storage of medicines in the home and secure disposal of covered drugs; rates of misuse, abuse, overdoses, and poisonings from prescription and nonprescription drugs; and diversions of covered drugs from sewer, solid waste, and septic systems. To conduct this evaluation, the department and the academic institution may rely on available data sources, including the public awareness surveys required under this chapter, and the prescription drug monitoring program and public health surveys such as the Washington state healthy youth survey. The department and the academic institution may also consult with other state and local agencies and interested stakeholders; and

(d) Provide any recommendations for legislation.

NEW SECTION. **Sec. 20.** (1)(a) The department shall contract with the statewide program of poison and drug information services identified in RCW 18.76.030 to conduct a survey of residents to measure whether the secure medicine collection and disposal system and the program promotion, education, and public outreach requirements established in this chapter have led to statistically significant changes in: (i) Resident attitudes and behavior on safe storage and secure disposal of prescription and nonprescription medications used in the home; and (ii) the rates of abuse or misuse of or accidental exposure to prescription and nonprescription drugs.

(b) The survey of residents must include telephone follow-up with users of the program's emergency telephone service. The survey must be conducted before the secure medicine collection and disposal system is implemented and again no earlier than four years after the system is implemented.

(2) The statewide program of poison and drug information services shall report the survey results to the legislature and the department of health within six months of completion of the survey.

(3) This section expires July 1, 2026.

Sec. 21. RCW 42.56.270 and 2017 c 317 s 17 are each amended to read as follows:

The following financial, commercial, and proprietary information is exempt from disclosure under this chapter:

(1) Valuable formulae, designs, drawings, computer source code or object code, and research data obtained by any agency within five years of the request for disclosure when disclosure would produce private gain and public loss;

(2) Financial information supplied by or on behalf of a person, firm, or corporation for the purpose of qualifying to submit a bid or proposal for (a) a ferry system construction or repair contract as required by RCW 47.60.680 through 47.60.750 or (b) highway construction or improvement as required by RCW 47.28.070;

(3) Financial and commercial information and records supplied by private persons pertaining to export services provided under chapters 43.163 and 53.31 RCW, and by persons pertaining to export projects under RCW 43.23.035;

(4) Financial and commercial information and records supplied by businesses or individuals during application for loans or program services provided by chapters 43.325, 43.163, 43.160, 43.330, and 43.168 RCW, or during application for economic development loans or program services provided by any local agency;

(5) Financial information, business plans, examination reports, and any information produced or obtained in evaluating or examining a business and industrial development corporation organized or seeking certification under chapter 31.24 RCW;

(6) Financial and commercial information supplied to the state investment board by any person when the information relates to the investment of public trust or retirement funds and when disclosure would result in loss to such funds or in private loss to the providers of this information;

(7) Financial and valuable trade information under RCW 51.36.120;

(8) Financial, commercial, operations, and technical and research information and data submitted to or obtained by the clean Washington center in applications for, or delivery of, program services under chapter 70.95H RCW;

(9) Financial and commercial information requested by the public stadium authority from any person or organization that leases or uses the stadium and exhibition center as defined in RCW 36.102.010;

(10)(a) Financial information, including but not limited to account numbers and values, and other identification numbers supplied by or on behalf of a person, firm, corporation, limited liability company, partnership, or other entity related to an application for a horse racing license submitted pursuant to RCW 67.16.260(1)(b),

marijuana producer, processor, or retailer license, liquor license, gambling license, or lottery retail license;

(b) Internal control documents, independent auditors' reports and financial statements, and supporting documents: (i) Of house-banked social card game licensees required by the gambling commission pursuant to rules adopted under chapter 9.46 RCW; or (ii) submitted by tribes with an approved tribal/state compact for class III gaming;

(11) Proprietary data, trade secrets, or other information that relates to: (a) A vendor's unique methods of conducting business; (b) data unique to the product or services of the vendor; or (c) determining prices or rates to be charged for services, submitted by any vendor to the department of social and health services for purposes of the development, acquisition, or implementation of state purchased health care as defined in RCW 41.05.011;

(12)(a) When supplied to and in the records of the department of commerce:

(i) Financial and proprietary information collected from any person and provided to the department of commerce pursuant to RCW 43.330.050(8); and

(ii) Financial or proprietary information collected from any person and provided to the department of commerce or the office of the governor in connection with the siting, recruitment, expansion, retention, or relocation of that person's business and until a siting decision is made, identifying information of any person supplying information under this subsection and the locations being considered for siting, relocation, or expansion of a business;

(b) When developed by the department of commerce based on information as described in (a)(i) of this subsection, any work product is not exempt from disclosure;

(c) For the purposes of this subsection, "siting decision" means the decision to acquire or not to acquire a site;

(d) If there is no written contact for a period of sixty days to the department of commerce from a person connected with siting, recruitment, expansion, retention, or relocation of that person's business, information described in (a)(ii) of this subsection will be available to the public under this chapter;

(13) Financial and proprietary information submitted to or obtained by the department of ecology or the authority created under chapter 70.95N RCW to implement chapter 70.95N RCW;

(14) Financial, commercial, operations, and technical and research information and data submitted to or obtained by the life sciences discovery fund authority in applications for, or delivery of, grants under chapter 43.350 RCW, to the extent that such information, if revealed, would reasonably be expected to result in private loss to the providers of this information;

(15) Financial and commercial information provided as evidence to the department of licensing as required by RCW 19.112.110 or 19.112.120, except information disclosed in aggregate form that does not permit the identification of information related to individual fuel licensees;

(16) Any production records, mineral assessments, and trade secrets submitted by a permit holder, mine operator, or landowner to the department of natural resources under RCW 78.44.085;

(17)(a) Farm plans developed by conservation districts, unless permission to release the farm plan is granted by the landowner or operator who requested the plan, or the farm plan is used for the application or issuance of a permit;

(b) Farm plans developed under chapter 90.48 RCW and not under the federal clean water act, 33 U.S.C. Sec. 1251 et seq., are subject to RCW 42.56.610 and 90.64.190;

(18) Financial, commercial, operations, and technical and research information and data submitted to or obtained by a health sciences and services authority in applications for, or delivery of, grants under RCW 35.104.010 through 35.104.060, to the extent that such information, if revealed, would reasonably be expected to result in private loss to providers of this information;

(19) Information gathered under chapter 19.85 RCW or RCW 34.05.328 that can be identified to a particular business;

(20) Financial and commercial information submitted to or obtained by the University of Washington, other than information the university is required to disclose under RCW 28B.20.150, when the information relates to investments in private funds, to the extent that such information, if revealed, would reasonably be expected to result in loss to the University of Washington consolidated endowment fund or to result in private loss to the providers of this information;

(21) Market share data submitted by a manufacturer under RCW 70.95N.190(4);

(22) Financial information supplied to the department of financial institutions or to a portal under RCW 21.20.883, when filed by or on behalf of an issuer of securities for the purpose of obtaining the exemption from state securities registration for small securities offerings provided under RCW 21.20.880 or when filed by or on behalf of an investor for the purpose of purchasing such securities;

(23) Unaggregated or individual notices of a transfer of crude oil that is financial, proprietary, or commercial information, submitted to the department of ecology pursuant to RCW 90.56.565(1)(a), and that is in the possession of the department of ecology or any entity with which the department of ecology has shared the notice pursuant to RCW 90.56.565;

(24) Financial institution and retirement account information, and building security plan information, supplied to the liquor and cannabis board pursuant to RCW 69.50.325, 69.50.331, 69.50.342, and 69.50.345, when filed by or on behalf of a licensee or prospective licensee for the purpose of obtaining, maintaining, or renewing a license to produce, process, transport, or sell marijuana as allowed under chapter 69.50 RCW;

(25) Marijuana transport information, vehicle and driver identification data, and account numbers or unique access identifiers issued to private entities for traceability system access, submitted by an individual or business to the liquor and cannabis board under the requirements of RCW 69.50.325, 69.50.331, 69.50.342, and 69.50.345 for the purpose of marijuana product traceability. Disclosure to local, state, and federal officials is not considered public disclosure for purposes of this section;

(26) Financial and commercial information submitted to or obtained by the retirement board of any city that is responsible for the management of an employees' retirement system pursuant to the authority of chapter 35.39 RCW, when the information relates to investments in private funds, to the extent that such information, if revealed, would reasonably be expected to result in loss to the retirement fund or to result in private loss to the providers of this information except that (a) the names and commitment amounts of the private funds in which retirement funds are invested and (b) the aggregate quarterly performance results for a retirement fund's portfolio of investments in such funds are subject to disclosure;

(27) Proprietary financial, commercial, operations, and technical and research information and data submitted to or obtained by the liquor and cannabis board in applications for marijuana research licenses under RCW 69.50.372, or in reports submitted by marijuana research licensees in accordance with rules adopted by the liquor and cannabis board under RCW 69.50.372; ((~~and~~))

(28) Trade secrets, technology, proprietary information, and financial considerations contained in any agreements or contracts, entered into by a licensed marijuana business under RCW 69.50.395, which may be submitted to or obtained by the state liquor and cannabis board; and

(29) Proprietary information filed with the department of health under chapter 69.--- RCW (the new chapter created in section 25 of this act).

Sec. 22. RCW 69.41.030 and 2016 c 148 s 11 are each amended to read as follows:

(1) It shall be unlawful for any person to sell, deliver, or possess any legend drug 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, 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, an osteopathic physician assistant under chapter 18.57A RCW when authorized by the board of osteopathic medicine and surgery, a physician assistant under chapter 18.71A RCW when authorized by the medical quality assurance 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, a licensed osteopathic 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.--- RCW (the new chapter created in section 25 of this act).

(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 possession is a misdemeanor.

NEW SECTION. **Sec. 23.** A new section is added to chapter 69.50 RCW to read as follows:

It is not a violation of this chapter to possess or deliver a controlled substance in compliance with chapter 69.--- RCW (the new chapter created in section 25 of this act).

NEW SECTION. **Sec. 24.** A new section is added to chapter 70.95 RCW to read as follows:

An authorized collector regulated under chapter 69.--- RCW (the new chapter created in section 25 of this act) is not required to obtain a permit under RCW 70.95.170 unless the authorized collector is required to obtain a permit under RCW 70.95.170 as a consequence

of activities that are not directly associated with the collection facility's activities under chapter 69.--- RCW (the new chapter created in section 25 of this act).

NEW SECTION. **Sec. 25.** Sections 2 through 20 of this act constitute a new chapter in Title 69 RCW.

NEW SECTION. **Sec. 26.** A new section is added to chapter 43.131 RCW to read as follows:

The authorization for drug take-back programs created in this act shall be terminated on January 1, 2029, as provided in section 27 of this act.

NEW SECTION. **Sec. 27.** A new section is added to chapter 43.131 RCW to read as follows:

The following acts or parts of acts, as now existing or hereafter amended, are each repealed, effective January 1, 2030:

(1) RCW 69.--.--- and 2018 c ... s 2 (section 2 of this act);
(2) RCW 69.--.--- and 2018 c ... s 3 (section 3 of this act);
(3) RCW 69.--.--- and 2018 c ... s 4 (section 4 of this act);
(4) RCW 69.--.--- and 2018 c ... s 5 (section 5 of this act);
(5) RCW 69.--.--- and 2018 c ... s 6 (section 6 of this act);
(6) RCW 69.--.--- and 2018 c ... s 7 (section 7 of this act);
(7) RCW 69.--.--- and 2018 c ... s 8 (section 8 of this act);
(8) RCW 69.--.--- and 2018 c ... s 9 (section 9 of this act);
(9) RCW 69.--.--- and 2018 c ... s 10 (section 10 of this act);
(10) RCW 69.--.--- and 2018 c ... s 11 (section 11 of this act);
(11) RCW 69.--.--- and 2018 c ... s 12 (section 12 of this act);
(12) RCW 69.--.--- and 2018 c ... s 13 (section 13 of this act);
(13) RCW 69.--.--- and 2018 c ... s 14 (section 14 of this act);
(14) RCW 69.--.--- and 2018 c ... s 15 (section 15 of this act);
(15) RCW 69.--.--- and 2018 c ... s 16 (section 16 of this act);
(16) RCW 69.--.--- and 2018 c ... s 17(section 17 of this act);
(17) RCW 69.--.--- and 2018 c ... s 18 (section 18 of this act);
(18) RCW 69.--.--- and 2018 c ... s 19 (section 19 of this act); and
(19) RCW 69.--.--- and 2018 c ... s 20 (section 20 of this act)."

Correct the title.

EFFECT: (1) Clarifies that the intent of the legislature is to establish a single, uniform, statewide system of regulation for drug-take back programs.

(2) Defines "administer" and "mail-back distribution location," and limits the definition of "covered drug" to exclude a number of products that were previously excluded from the definition of "drug" and specifically excludes emptied injector products, emptied medical devices, and exposed needles or sharps.

(3) Requires program operators to submit program proposals by July 1, 2019, rather than October 1, 2019.

(4) Allows covered manufacturers in a drug take-back program to apportion the costs as the manufacturers determine, rather than according to sales revenue in Washington state.

(5) Limits the fees the Department of Health (Department) can collect from program operators to the actual costs of administration, oversight, enforcement, and starting in 2020 to 10% of the total program costs. Limits adjustments to the fees for inflation to not exceed the percentage change in the consumer price index for all urban consumers averaged by city, in the previous year.

(6) Increases the time limit for a program operator to initiate operation of an approved drug take-back program to 180 days from 90 days, and provides 90 days for a program operator to submit a revised proposal that was rejected by the Department instead of 60 days.

(7) Requires program operators to notify the Department of substantial changes to the program and 15 days prior to the change, rather than notification of any substantive change 30 days in advance. For changes that are not substantive, the program operator must notify the Department seven days in advance, instead of 15 days. Program operators must notify the department of any changes in ownership or contact information for participating covered manufacturers within 90 days, instead of 30 days.

(8) Requires drug take-back program collection systems to provide reasonably convenient access and that there must be a collection site for each population center plus an additional site for every 50,000 residents instead of every 20,000 residents.

(9) Allows a program operator to establish mail-back distribution locations or hold periodic collection events instead of requiring periodic collection events.

(10) Clarifies that the alternative collection methods of mail-back programs and periodic events may only be conducted to the extent permissible under state and federal laws.

(11) Removes the requirement on program operators to conduct a survey on consumer awareness, the use of the collection methods, and knowledge of the risks of abuse and overdose and places the responsibility on the Department.

(12) Changes the date the annual report must be submitted to July 1st instead of April 1st.

(13) Requires program operators submit within 30 days of the annual period of operation the total amount, by weight, of covered drugs collected from each collection site during the prior year.

(14) Limits the Department's enforcement authority by only allowing the Department to require a person or entity to engage in or refrain from engaging in certain activities that pertain to the chapter.

(15) Reduces the time period in which a county may enforce a grandfathered ordinance after an approved drug take-back program begins operating and a manufacturer in compliance with a grandfathered ordinance is considered in compliance with the state

law for purposes of that county, from 18 months to 12 months. Clarifies that after the 12-month period that all local laws regarding drug take-back programs are preempted.

(16) Requires the Department to utilize an academic institution that is not an agency of the state in evaluation of the impact of the drug take-back program.

(17) Subjects the authorization for drug take-back programs to the sunset review process, terminating the authority on January 1, 2029.

(18) Increases dates in the bill by one year, and makes a technical change to reflect legislation enacted in 2017.

--- END ---