E2SHB 1472 - H AMD 565

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By Representative Fitzgibbon

- 1 Strike everything after the enacting clause and insert the 2 following:
- Sec. 1. The definitions in this section apply 3 "NEW SECTION. this chapter unless the context clearly requires 4 throughout otherwise. 5
 - (1) "Alternatives assessment" means a process for identifying and comparing chemical and nonchemical alternatives currently existence that can be practicably and economically used to replace the use of a chemical or, if a safer alternative is not identified, to reduce the amount of or exposure to that chemical.
- 11 "Biomonitoring" means assessment of human exposures 12 chemicals by measuring the chemicals or their metabolites in human 13 tissues or specimens, such as blood, breast milk, and urine.
- 14 "Chemical" means a substance, including metals, with a (3) 15 distinct molecular composition or a group of structurally related substances, and includes the breakdown products of the substance or 16 17 substances that form through decomposition, degradation, or 18 metabolism.
- 19 (4) "Chemical action plan" means a report that identifies, 20 characterizes, and evaluates current and legacy uses and releases of a specific chemical or group of chemicals and identifies actions 21 22 needed to protect human health and the environment.
 - (5) "Department" means the department of ecology.
- 24 (6) "Director" means the director of the department of ecology or the director's designee. 25
 - (7) "Distributor" means:
- (a) An importer or a domestic distributor of a product sold or 27 28 offered for sale in or into the state.
 - (b) "Distributor" does not include:
- (i) Small businesses as defined in RCW 19.85.020; or 30

- 1 (ii) For purposes of section 5 of this act only, a person or entity that demonstrates that it does not exercise control over the 2 process by which a product was formulated; or 3
 - (iii) For the purposes of section 3 of this act only, a person or entity that provides documentation to the department that the person or entity is unable to provide any of the information under section 3(2) of this act, consistent with section 3(7) of this act.
 - (8) "Documentation" means:

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- (a) Written or electronic communications between the distributor 9 and manufacturer demonstrating that the distributor has requested 10 11 information on the chemical used in products;
 - (b) A description of the efforts the distributor has undertaken to respond to the information request made by the department under section 3 of this act, including the reason that the distributor is unable to provide the requested information; and
 - signature of an authorized representative the distributor and an accompanying dated statement that declares that the information provided to the department under (b) of this subsection is accurate.
- (9) "Emerging chemicals" means chemicals that meet the criteria 20 21 of a high priority chemical as defined in RCW 70.240.010 and either:
- (a) Meet the criteria for a high priority chemical of high 22 concern for children as described in RCW 70.240.030(1) (a) through 23 24 (c); or
- (b) Have been shown through environmental monitoring studies to 25 be present in fish, wildlife, air, water, soil, or sediment. 26
 - (10) "Manufacturer" means:
- (a) Any person who manufactures a product sold or offered for 28 29 sale in or into the state.
- (b) "Manufacturer" does not include small businesses as defined 30 in RCW 19.85.020. 31
- (11)(a) "Product" means any item sold in or into the state for 32 residential or commercial use, including any component or product 33 34 packaging.
 - (b) "Product" does not include the following items:
- 36 (i) Food or beverage and food or beverage packaging, regulated by the United States food and drug administration or the United States 37 department of agriculture; 38
- 39 (ii) Tobacco products;

- 1 (iii) Drug or biological products and packaging regulated by the 2 United States food and drug administration;
- 3 (iv) Products and components produced under military 4 specifications;
- 5 (v) Products and components regulated by the federal aviation 6 administration;
- 7 (vi) Products registered for distribution in the state under 8 chapter 15.54 or 15.58 RCW;
- 9 (vii) Shipping pallets; and
- 10 (viii) Any previously owned product sold in casual or isolated 11 sales as defined in RCW 82.04.040 or products sold by nonprofit 12 organizations.
- 13 (12) "Product component" means a uniquely identifiable part, 14 material, or coating that is included as a part of a finished 15 product.
- 16 (13) "Safer alternative" means an alternative, demonstrated by an 17 alternatives assessment, that meets improved hazard and exposure 18 considerations, exhibits lower risk, and can be practicably and 19 economically substituted for the original chemical.
- 20 (14) "Unsuitable alternative" means an alternative identified 21 through the alternatives assessment process that does not meet the 22 hazard, exposure, cost, performance, and availability criteria of a 23 safer alternative.
- 24 (15) This section expires June 30, 2025.
- NEW SECTION. Sec. 2. (1) Beginning January 1, 2016, and every two years thereafter, the department, in consultation with the department of health, must select up to two chemicals for the development of chemical action plans, as provided in section 4 of this act, from the following:
- 30 (a) Chemicals regulated by the department under human health 31 criteria for water quality standards in the proposed rule published 32 by the department on February 4, 2015, in the Washington State 33 Register, as WSR 15-03-015;
- 34 (b) Chemicals that affect water quality and are persistent 35 bioaccumulative toxins as defined in chapter 173-333 WAC, as of the 36 effective date of this section; or
 - (c) Emerging chemicals.

38 (2) The department may conduct environmental monitoring or, 39 subject to the availability of amounts appropriated for this specific

- 1 purpose, may request the department of health to conduct
- 2 biomonitoring of a chemical, to verify the chemical is present in the
- 3 state's environment or population or to better understand
- 4 environmental or human exposure in the state. Environmental
- 5 monitoring and biomonitoring conducted pursuant to this chapter must
- 6 be of a minimum scope necessary to adequately inform a chemical
- 7 action plan.
- 8 (3)(a) The first five chemicals selected for chemical action
- 9 plans must be chosen from the chemicals identified in subsection
 - (1)(a) or (b) of this section.
- 11 (b) The director shall notify the public of the selection of a
- 12 chemical for the development of a chemical action plan. The notice
- 13 must state the basis for the selection and include a draft schedule
- 14 for completing the chemical action plan. The notice must be published
- 15 in the Washington State Register. The department shall provide an
- 16 opportunity for public review and comment before finalizing the
- 17 schedule.

- 18 (c) When selecting chemicals for the development of chemical
- 19 action plans, the department must consider:
- 20 (i) Opportunities for reducing or phasing out uses, production,
- 21 or releases of a chemical;
- 22 (ii) Current scientific evidence on the extent and impact of the
- 23 chemical exposure to humans and biota;
- 24 (iii) Current scientific evidence on the combined effects of
- 25 exposure to a chemical and other substances commonly present in the
- 26 Washington environment;
- 27 (iv) Current scientific evidence on the susceptibility o
- 28 sensitive population groups and environmental media from exposure to
- 29 a chemical, as well as cumulative effects of multiple exposures;
- 30 (v) The relative ranking assigned to a chemical by the department
- 31 based on information applicable to Washington about a chemical's
- 32 properties, uses of a chemical, releases of a chemical, and levels of
- 33 a chemical present in the environment and in residents;
- 34 (vi) Whether a chemical has been determined to impact Washington
- 35 state waters through identification under section 303(d) of the
- 36 federal clean water act;
- 37 (vii) The potential for an emerging chemical to impair water
- 38 quality; and
- 39 (viii) Existing plans or regulatory requirements to reduce or
- 40 phase out the uses and releases of a chemical.

- 1 (d) The department must identify the sources of information it 2 relied upon in selecting chemicals for the development of chemical 3 action plans under this section, including peer-reviewed science.
 - (4) This section expires June 30, 2025.

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- 5 <u>NEW SECTION.</u> **Sec. 3.** (1) The department may request information from manufacturers and distributors about products 6 or product components that contain a chemical selected for a chemical action 7 plan under section 2 of this act. Prior to requesting information 8 from a manufacturer or a distributor under this subsection, the 9 10 department must consult with a chemical action plan external advisory 11 committee, if one has been formed, to evaluate the chemical that is the subject of the information request. If a chemical action plan 12 external advisory committee has not yet been formed at the time of an 13 information request under this section, the department must seek 14 15 from representatives of the organizations and 16 identified in section 4(1) of this act prior to making the information request. Requests for information must be reasonable in 17 scope and frequency, and focused on: 18
 - (a) The most common and prevalent uses of the chemical or products or product components that contain the chemical, based on the department's existing knowledge about the chemical;
 - (b) Areas where there is an identified gap in public or department knowledge about the chemical; or
 - (c) Uses of the chemical, products, or product components that the department has reason to believe are likely to be responsible for or associated with a significant portion of chemical releases into the environment or public health exposures.
- 28 (2) Within twelve months of a request by the department, 29 manufacturers or distributors shall report the following:
- 30 (a) The name and address of the manufacturer or distributor and 31 the contact information of the person responsible for responding to 32 the department's requests on behalf of the manufacturer or 33 distributor;
- 34 (b) The chemical abstracts service registry number of the 35 chemical for which information is being requested;
- (c) A brief description of the manufacturer's or distributor'sproduct or product component categories containing the chemical;
- (d) A description of the function or functions of the chemical in the manufacturer's or distributor's products or product components;

The amount of the chemical used in each unit of the manufacturer's or distributor's products or product components, which may be reported in ranges, rather than the exact amount;

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- An estimate of the average daily, weekly, or monthly commercial consumption of the chemical by businesses or the public that is associated with the commercial consumption of manufacturer's or distributor's product; and
- (g) Any other information the manufacturer or distributor deems relevant to the appropriate use of the chemical, products, or product components.
- (3) In response to an information request from the department under this section, a manufacturer or distributor may extrapolate amounts and estimates from data sources that are not specific to Washington state. A manufacturer or distributor may also submit or rely on sources of data compiled by federal agencies, other states, nations, or other sources of data that meet the requirements of the addition, information request. In multiple manufacturers, distributors, or a business association, may collaborate and submit a single response on behalf of multiple manufacturers or distributors for a chemical found in similar products or product components in response to the department's request for information. All submissions in response to the department's information request must include the information required in subsection (2)(a) of this section for each manufacturer or distributor. However, if multiple manufacturers, distributors, or a business association collaborate to submit a single response on behalf of multiple manufacturers or distributors, the information required in subsection (2)(b) through (f) of this section is not required to be provided in a manner that attributes product or chemical use data or information to individual manufacturers or distributors.
- The department shall specify the required format for submission of the information required under subsection (2) of this section. The format should be generally consistent with the format specified by other states or federal agencies with substantially similar reporting requirements.
- (5) Where information submitted by a manufacturer or distributor under chapter 70.240 RCW is the same as the information required to be submitted by the manufacturer or distributor in subsection (2) of this section, the manufacturer or distributor is not required to submit the same information more than once.

(6) The department may, by order, require a manufacturer distributor subject to the reporting requirement in subsection (2) of this section to provide additional information that is relevant to the development of a chemical action plan under section 4 of this act. Prior to an order under this subsection, the department must consult with the external advisory committee formed for the chemical action plan, if one has been formed. If a chemical action plan external advisory committee has not yet been formed at the time of an order pursuant to this subsection, the department must seek input from representatives of the organizations and entities identified in section 4(1) of this act prior to making the information order. An order by the department must also meet the reasonableness criteria of subsection (1) of this section.

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- (7)(a) For purposes of making information requests and orders under this section, the department must use the following hierarchy to determine whether the department will hold the manufacturer or distributor of a product primarily responsible for the response:
- (i) The department must hold a manufacturer primarily responsible for the response to the information request, unless the manufacturer does not have a presence in the United States.
- (ii) If the manufacturer of the product does not have a presence the United States, the department may hold the product's distributor primarily responsible for responding to the information request or order. If the distributor provides documentation to the department that it does not reasonably have access to the full set of information required to respond to the order, the department may only require the distributor to submit information for the categories under subsection (2) of this section for which it has information. The department may not penalize a distributor that provides documentation that it does not reasonably have access to the full set of information required to respond to the order under this section, as long as the distributor's response to the information request includes all requested information that the distributor is reasonably able to access.
- (b) Nothing in this subsection (7) limits the authority of the department to request information from a distributor of a product about the persons or entities who are manufacturers of the product.
- A manufacturer or distributor may request submitted under this section be held confidential as provided under section 7 of this act.

- 1 (9) Nothing in this section may be construed to make a distributor or a manufacturer an agent of the state. 2
- (10) This section expires June 30, 2025. 3

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- **Sec. 4.** (1) When developing a chemical action NEW SECTION. 4 5 plan, the department shall convene an external advisory committee to and provide input, expertise, 6 hold meetings and additional information. All external advisory committee meetings must be open to 7 the public. The department must invite representatives from, 8 minimum, the following organizations and entities to serve 9 10 external advisory committee members: Large and small business 11 sectors; a statewide business association with over one thousand total members and that represents multiple business sectors; 12 community, environmental, and public health advocacy groups; local 13 governments; affected and interested businesses; and public health 14 15 agencies. State agencies and technical experts may be requested to 16 participate.
 - (2) All chemical action plans must include the following:
- (a) The chemical name and a description of its properties, uses, 18 and products or product components in which the chemical is found; 19
- 20 (b) An analysis of the available information on the production, unintentional production, current and legacy uses, and disposal of 21 22 the chemical;
 - (c) Information on the known, potential, or proven impacts on human health and the environment associated with the use and release of the chemical;
 - (d) An evaluation of regulatory and nonregulatory activities that influence production, uses, releases, and management of the chemical;
- (e) Identification of actions that, if needed, would eliminate, 28 reduce, or manage exposures to the chemical; 29
- 30 (f) A prioritization of sources of exposures and releases of the chemical into the environment. The prioritization must address 31 impacts to human health and the environment, potential to affect 32 water quality, and feasibility and cost of actions that could be 33 taken to address exposures or releases; 34
- (g) A determination if an alternatives assessment is recommended, 35 which must address the prioritization required under (f) of this 36 subsection; and 37
- 38 (h) A determination of the persons or entities responsible for completing the alternatives assessments, if recommended. 39

- 1 (3) All recommendations in a chemical action plan may be included 2 only after consideration of the following criteria:
- 3 (a) Opportunity for environmental and human health benefits in 4 the state of Washington;
 - (b) Economic and social impacts;
- 6 (c) Feasibility;

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- 7 (d) Availability and effectiveness of safer alternatives, if 8 known; and
- 9 (e) Consistency with existing federal and state regulatory 10 requirements.
- 11 (4) The department must include in the chemical action plan a 12 summary of all views, including dissenting views, held by external 13 advisory committee members regarding the recommendations contained in 14 the plan.
- 15 (5) The department must identify the sources of information it 16 relied upon in completing a chemical action plan under this section, 17 including peer-reviewed science.
- 18 (6) This section expires June 30, 2025.
 - NEW SECTION. Sec. 5. (1) The department may require manufacturers or distributors to conduct two assessments, each of which must be consistent with recommendations from chemical action plans completed for the chemicals that are the subject of the alternatives assessments and their prioritization under section 4 of this act. However, the department may not require manufacturers to conduct more than one alternatives assessment for a chemical under section 2(1)(b) of this act.
 - (a) If the department orders manufacturers or distributors to conduct an alternatives assessment for a chemical under section 2(1) (a) or (b) of this act, the department may not require the alternatives assessment to be conducted for a greater breadth of uses, products, or manufacturers or distributors than is necessary to address sources of environmental or human exposure to the chemical.
- 33 (b) If the department orders manufacturers or distributors to 34 conduct an alternatives assessment for a chemical under section 35 2(1)(c) of this act, the alternatives assessment must be limited to a 36 single type of use of a chemical or a single type of product or 37 product component in which the chemical is found.
- 38 (2) In addition to the authority granted in subsection (1) of 39 this section, the department may require, by order, manufacturers or Official Print - 9 1472-S2.E AMH FITZ H2863.6

1 distributors to conduct an alternatives assessment for 2 polychlorinated biphenyls in pigments and dyes.

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- (3)(a) When ordered by the department to conduct an alternatives assessment, a manufacturer or distributor must submit:
- (i) An alternatives assessment as required under subsection (6) of this section to the department for each use of the chemical specified by the department; or
- (ii) A peer-reviewed alternatives assessment completed by an authoritative entity, including the United States environmental protection agency, the federal food and drug administration, or other nations or states, that meets the requirements of subsection (6) of this section.
- (b) A manufacturer or distributor must submit the alternatives assessment to the department within eighteen months from the date the alternatives assessment is ordered. However, the department may grant an extension on a case-by-case basis for good cause if the manufacturer or distributor shows that additional time is necessary to complete the alternatives assessment or to substantially improve the quality of the alternatives assessment.
- (c) A manufacturer or distributor may meet its obligation under this section by collaborating with other manufacturers, distributors, or business associations of similar products to conduct and complete an alternatives assessment. A manufacturer or distributor complying with subsection (3)(c) of this section is not in violation of this chapter.
- (4) A manufacturer or distributor may not be required to submit an alternatives assessment when the manufacturer or distributor provides to the department within thirty days of receipt of an order to conduct an alternatives assessment a certificate of compliance.
 - (a) A certificate of compliance must include the following:
- 31 (i) Chemical names and chemical abstracts service registry 32 numbers for all chemicals that currently contribute to the specific 33 function previously served by the chemical that is subject to the 34 alternatives assessment order;
- (ii) How the manufacturer is meeting the function of the chemical that is subject to the alternatives assessment order with a safer alternative;
- 38 (iii) Documents or other evidence demonstrating that the 39 manufacturer or distributor has: (A) Ceased using the chemical or 40 importing or distributing products containing the chemical for which

- 1 it would be required to conduct an alternatives assessment; or (B)
- 2 committed resources in pursuit of a plan to phase out, within a
- reasonable time, the chemical for which the manufacturer or 3
- distributor would be required to conduct an alternatives assessment; 4
- and 5

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- 6 (iv) The signature of an authorized official of the manufacturer 7 or distributor.
- (b) A manufacturer or distributor that is not required to conduct 8 an alternatives assessment under subsection (3) of this section is 9 not in violation of this chapter. 10
 - (5)(a) The department, in consultation with the chemical action plan external advisory committee, may contract with an independent qualified third party to conduct an alternatives assessment when:
 - (i) A manufacturer or distributor is not required to conduct an alternatives assessment under this section; or
 - (ii) The department determines that an alternatives assessment submitted by a manufacturer or distributor does not meet definition or required objectives of an alternatives assessment.
 - The department must ensure an alternatives assessment (b) completed by a qualified third party is peer-reviewed and meets the requirements under subsection (6) of this section.
 - (c) The department may require a manufacturer or distributor, consistent with recommendations in a chemical action plan, to provide additional information that is relevant to the development of a department-conducted alternatives assessment.
 - (6) An alternatives assessment must:
 - (a) Meet the objective of assessing less toxic chemicals and nonchemical alternatives to replace the use of a chemical or, if a safer alternative is not identified, to reduce the amount of or exposure to chemicals in products and product components, and to avoid the unintended consequence of switching to a substitute that presents an equivalent or greater concern;
 - (b) Follow the guidelines issued by the interstate chemicals clearinghouse, the national academy of sciences, or equivalent methodology; and
- 36 (c) Include, at a minimum: (i) An evaluation of chemical hazard, exposure, performance, consumer acceptance, cost, and availability; 37 (ii) equivalent information, as required under (c)(i) of this 38 39 subsection, for each alternative considered; and (iii) the 40 identification of alternatives and unsuitable alternatives.

- 1 (7) If the department determines, based on an alternatives 2 assessment, that a safer alternative exists, the department may 3 submit agency request legislation recommending the prohibition of 4 certain uses of a chemical or other actions determined appropriate, 5 including restrictions on the use of unsuitable alternatives.
 - (8) This section expires June 30, 2025.

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- NEW SECTION. Sec. 6. (1) A manufacturer or distributor violating this chapter is subject to a civil penalty not to exceed five thousand dollars for each violation in the case of a first offense. Manufacturers or distributors who are repeat violators are subject to a civil penalty not to exceed ten thousand dollars for each repeat offense.
- 13 (2) Any penalty provided for in this section, and any order 14 issued by the department under this chapter, may be appealed to the 15 pollution control hearings board.
- 16 (3) All penalties collected under this chapter shall be deposited 17 in the state toxics control account created in RCW 70.105D.070.
- 18 (4) This section expires June 30, 2025.
- NEW SECTION. Sec. 7. (1) Manufacturers or distributors submitting information or records to the department may request that the information or records be made available only for the confidential use of the director, the department, or the appropriate division of the department.
 - (2)(a) A manufacturer or distributor requesting confidentiality for information submitted under this chapter must demonstrate to the department how the records relate to processes of production unique to the manufacturer or distributor or how releasing the records to the public may adversely affect the competitive position of the manufacturer or distributor.
 - (b)(i) The director shall give consideration to the request for confidentiality and if such action would not be detrimental to the public interest and is otherwise within accord with the policies and purposes of chapter 43.21A RCW, the director must grant the request for the information to remain confidential as authorized in RCW 43.21A.160.
- 36 (ii) The department must respond to a manufacturer's or 37 distributor's request within fourteen days of receipt of the request. 38 The department must inform the manufacturer or distributor regarding

- 1 its determination of whether the submitted information should be kept confidential under this section and RCW 43.21A.160 and its reasons 2 for the determination. 3
- (iii) The department must keep confidential any records furnished 4 by a manufacturer or distributor under this chapter that relate to 5 6 proprietary manufacturing processes or chemical formulations used in 7 products or manufacturing processes.
 - (3) If the director denies the request of a manufacturer or distributor to keep submitted information or records confidential under this section, the manufacturer or distributor may appeal the denial to a court of competent jurisdiction. In a review of whether the submitted information or records meet the criteria of RCW 43.21A.160 and this section, a court must examine submitted information or records in camera.
- (4) This section expires June 30, 2025. 15

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- Sec. 8. (1) The department may adopt rules as 16 NEW SECTION. necessary for the purpose of implementing, administering, and 17 enforcing this chapter. Rules adopted to require manufacturers or 18 distributors to conduct alternatives assessments must be consistent 19 with section 5 of this act. 20
- (2) This section expires June 30, 2025. 21
- NEW SECTION. 22 Sec. 9. A new section is added to chapter 39.26 23 RCW to read as follows:
- (1) The department shall establish purchasing and procurement 24 policies that provide a preference for products and products in 25 packaging that do not contain: 26
- 27 (a) Persistent bioaccumulative toxins, as identified by the department of ecology by rule; and 28
 - (b) Chemicals that have been addressed by a completed chemical action plan that has included a recommendation that the state adopt a purchasing and procurement policy for products and products packaging that do not contain the chemical.
- 33 (2) No agency may knowingly purchase products or products in packaging containing chemicals identified in subsection (1) of this 34 section unless there is no cost-effective and technologically 35 feasible alternative. When all available products contain a chemical 36 identified in subsection (1) of this section, a preference must be 37

given to alternative products that contain lesser amounts 1 of chemicals identified in subsection (1) of this section. 2

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- (3) Nothing in this section requires the department or any other state agency to breach an existing contract or dispose of stock that has been ordered or is in the possession of the department or other state agency as of the effective date of this section.
- (4) This section does not require the department or any other agency to test every product procured.
- 9 (5) The department or any other agency may request suppliers of products to provide testing data from an accredited laboratory or 10 11 testing facility documenting levels of a chemical identified in subsection (1) of this section in products or product packaging. 12 Requested or voluntarily received testing data from businesses, 13 manufacturers, distributors, organizations, and individuals must be 14 submitted for review to the department of ecology. 15
- 16 Sec. 10. RCW 43.21B.110 and 2013 c 291 s 33 are each amended to 17 read as follows:
- (1) The hearings board shall only have jurisdiction to hear and 18 decide appeals from the following decisions of the department, the 19 20 director, local conservation districts, the air pollution control boards or authorities as established pursuant to chapter 70.94 RCW, 21 local health departments, the department of natural resources, the 22 department of fish and wildlife, the parks and recreation commission, 23 24 and authorized public entities described in chapter 79.100 RCW:
- 25 (a) Civil penalties imposed pursuant to RCW 18.104.155, 70.94.431, 70.105.080, 70.107.050, 76.09.170, 77.55.291, 78.44.250, 26 27 88.46.090, 90.03.600, 90.46.270, 90.48.144, 90.56.310, 90.56.330, and 90.64.102. 28
- 29 (b) Orders issued pursuant to RCW 18.104.043, 18.104.060, 30 43.27A.190, 70.94.211, 70.94.332, 70.105.095, 86.16.020, 88.46.070, 90.14.130, 90.46.250, 90.48.120, and 90.56.330. 31
- (c) A final decision by the department or director made under 32 chapter 183, Laws of 2009. 33
- 34 Except as provided in RCW 90.03.210(2), the issuance, 35 modification, or termination of any permit, certificate, or license by the department or any air authority in the exercise of its 36 jurisdiction, including the issuance or termination of a waste 37 disposal permit, the denial of an application for a waste disposal 38 permit, the modification of the conditions or the terms of a waste 39

- 1 disposal permit, or a decision to approve or deny an application for a solid waste permit exemption under RCW 70.95.300. 2
- (e) Decisions of local health departments regarding the grant or 3 denial of solid waste permits pursuant to chapter 70.95 RCW. 4
- (f) Decisions of local health departments regarding the issuance 5 and enforcement of permits to use or dispose of biosolids under RCW 6 7 70.95J.080.
- (q) Decisions of the department 8 regarding waste-derived fertilizer or micronutrient fertilizer under RCW 15.54.820, and 9 decisions of the department regarding waste-derived soil amendments 10 11 under RCW 70.95.205.

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- (h) Decisions of local conservation districts related to the denial of approval or denial of certification of a dairy nutrient management plan; conditions contained in a plan; application of any dairy nutrient management practices, standards, methods, and technologies to a particular dairy farm; and failure to adhere to the plan review and approval timelines in RCW 90.64.026.
- (i) Any other decision by the department or an air authority which pursuant to law must be decided as an adjudicative proceeding under chapter 34.05 RCW.
- (j) Decisions of the department of natural resources, the department of fish and wildlife, and the department that reviewable under chapter 76.09 RCW, and the department of natural resources' appeals of county, city, or town objections under RCW 76.09.050(7).
- 26 (k) Forest health hazard orders issued by the commissioner of 27 public lands under RCW 76.06.180.
- (1) Decisions of the department of fish and wildlife to issue, 28 29 deny, condition, or modify a hydraulic project approval permit under chapter 77.55 RCW. 30
- 31 (m) Decisions of the department of natural resources that are reviewable under RCW 78.44.270. 32
- (n) Decisions of an authorized public entity under RCW 79.100.010 33 to take temporary possession or custody of a vessel or to contest the 34 35 amount of reimbursement owed that are reviewable by the hearings board under RCW 79.100.120. 36
- (o) Decisions regarding a restriction, order, or penalty issued 37 under chapter 70.--- RCW (the new chapter created in section 18 of 38 39 this act).

- 1 (2) The following hearings shall not be conducted by the hearings 2 board:
- (a) Hearings required by law to be conducted by the shorelines 3 hearings board pursuant to chapter 90.58 RCW. 4
- (b) Hearings conducted by the department pursuant to RCW 5 6 70.94.332, 70.94.390, 70.94.395, 70.94.400, 70.94.405, 70.94.410, and 7 90.44.180.
- (c) Appeals of decisions by the department under RCW 90.03.110 8 9 and 90.44.220.
- (d) Hearings conducted by the department to adopt, modify, or 10 11 repeal rules.
- (3) Review of rules and regulations adopted by the hearings board 12 13 shall be subject to review in accordance with the provisions of the 14 administrative procedure act, chapter 34.05 RCW.
- 15 **Sec. 11.** RCW 43.21B.110 and 2013 c 291 s 34 are each amended to read as follows: 16
- (1) The hearings board shall only have jurisdiction to hear and 17 decide appeals from the following decisions of the department, the 18 19 director, local conservation districts, the air pollution control 20 boards or authorities as established pursuant to chapter 70.94 RCW, local health departments, the department of natural resources, the 21 department of fish and wildlife, the parks and recreation commission, 22 23 and authorized public entities described in chapter 79.100 RCW:
- 24 (a) Civil penalties imposed pursuant to RCW 18.104.155, 70.94.431, 70.105.080, 70.107.050, 76.09.170, 77.55.291, 78.44.250, 25 88.46.090, 90.03.600, 90.46.270, 90.48.144, 90.56.310, 90.56.330, and 26 27 90.64.102.
- (b) Orders issued pursuant to RCW 18.104.043, 18.104.060, 28 43.27A.190, 70.94.211, 70.94.332, 70.105.095, 86.16.020, 88.46.070, 29 30 90.14.130, 90.46.250, 90.48.120, and 90.56.330.
- (c) Except as provided in RCW 90.03.210(2), the issuance, 31 modification, or termination of any permit, certificate, or license 32 by the department or any air authority in the exercise of its 33 jurisdiction, including the issuance or termination of a waste 34 35 disposal permit, the denial of an application for a waste disposal permit, the modification of the conditions or the terms of a waste 36 37 disposal permit, or a decision to approve or deny an application for a solid waste permit exemption under RCW 70.95.300. 38

1 (d) Decisions of local health departments regarding the grant or 2 denial of solid waste permits pursuant to chapter 70.95 RCW.

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- (e) Decisions of local health departments regarding the issuance and enforcement of permits to use or dispose of biosolids under RCW 70.95J.080.
- 6 (f) Decisions of the department regarding waste-derived 7 fertilizer or micronutrient fertilizer under RCW 15.54.820, and decisions of the department regarding waste-derived soil amendments 8 9 under RCW 70.95.205.
 - (g) Decisions of local conservation districts related to the denial of approval or denial of certification of a dairy nutrient management plan; conditions contained in a plan; application of any dairy nutrient management practices, standards, methods, technologies to a particular dairy farm; and failure to adhere to the plan review and approval timelines in RCW 90.64.026.
- (h) Any other decision by the department or an air authority 16 17 which pursuant to law must be decided as an adjudicative proceeding 18 under chapter 34.05 RCW.
 - (i) Decisions of the department of natural resources, the department of fish and wildlife, and the department that are reviewable under chapter 76.09 RCW, and the department of natural resources' appeals of county, city, or town objections under RCW 76.09.050(7).
- (j) Forest health hazard orders issued by the commissioner of 24 public lands under RCW 76.06.180. 25
 - (k) Decisions of the department of fish and wildlife to issue, deny, condition, or modify a hydraulic project approval permit under chapter 77.55 RCW.
- 29 (1) Decisions of the department of natural resources that are reviewable under RCW 78.44.270. 30
- 31 (m) Decisions of an authorized public entity under RCW 79.100.010 to take temporary possession or custody of a vessel or to contest the 32 amount of reimbursement owed that are reviewable by the hearings 33 board under RCW 79.100.120. 34
- (n) Decisions regarding a restriction, order, or penalty issued 35 under chapter 70.--- RCW (the new chapter created in section 18 of 36 37 this act).
- (2) The following hearings shall not be conducted by the hearings 38 board: 39

- 1 (a) Hearings required by law to be conducted by the shorelines 2 hearings board pursuant to chapter 90.58 RCW.
- (b) Hearings conducted by the department pursuant to RCW 3 70.94.332, 70.94.390, 70.94.395, 70.94.400, 70.94.405, 70.94.410, and 4 90.44.180. 5
- 6 (c) Appeals of decisions by the department under RCW 90.03.110 7 and 90.44.220.
- (d) Hearings conducted by the department to adopt, modify, or 8 9 repeal rules.
- (3) Review of rules and regulations adopted by the hearings board 10 11 shall be subject to review in accordance with the provisions of the administrative procedure act, chapter 34.05 RCW. 12
- NEW SECTION. Sec. 12. (1) By June 30, 2024, the department must 13 provide a report to the appropriate committees of the legislature to 14 review and evaluate the processes for chemical action plans and 15 alternatives assessments provided in this act. 16
- 17 (2) The report must include recommendations for changes to developing chemical action plans and alternatives assessments; 18 necessary legislative actions to improve the processes; and whether 19 20 the department should continue developing chemical actions plans and 21 alternatives assessments.
- 22 (3) This section expires June 30, 2025.

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<u>NEW SECTION.</u> **Sec. 13.** The legislature finds that: 23

- (1) The restrictions on the use of flame retardants in sections 14 and 15 of this act represent a logical outcome of the types of evaluative processes that are to be newly formally codified in chapter 70.--- RCW (the new chapter created in section 18 of this act).
- 29 (2) In 2006, the department of ecology completed a chemical action plan on the class of flame retardants known as polybrominated 30 diphenyl ethers, or PDBEs. This 2006 chemical action plan on the use 31 of flame retardant PBDEs led to a 2007 act, now codified in chapter 32 70.76 RCW, that restricted the use of PBDEs in many types of consumer 33 34 products, and also motivated an assessment by the departments of health and ecology to evaluate whether safer alternatives existed 35 that could be used to replace the function of PBDE chemicals in 36 certain commercial products, including residential upholstered 37 furniture. 38

- (3) A subsequent report, completed in 2015 by the department of ecology at the direction of the legislature, further evaluated publicly available completed assessments of uses of chlorinated and brominated flame retardants. This report included a recommendation that the use of many of these flame retardants should be restricted in certain commercial products.
- 7 (4) The public health and environmental risks associated with the types of flame retardants restricted in sections 14 and 15 of this 8 act have been, and continue to be, thoroughly evaluated by the 9 departments of health and ecology and considered by the legislature 10 11 in processes that began with the 2006 chemical action plan on PBDEs. The restrictions of certain uses of these flame retardants in 12 sections 14 and 15 of this act, therefore, represent an appropriate 13 and carefully tailored outcome of the conjoined evaluations that have 14 been undertaken by the legislative and executive branches of 15 Washington state government. 16
- 17 NEW SECTION. Sec. 14. A new section is added to chapter 70.240 RCW to read as follows: 18
- Beginning July 1, 2016, no manufacturer, wholesaler, or retailer 19 20 may manufacture, knowingly sell, offer for sale, distribute for sale, distribute for use in this state children's 21 residential upholstered furniture, as defined in RCW 70.76.010, 22 23 containing any of the following flame retardants in amounts greater 24 than one thousand parts per million in any product component:
- (1) TDCPP; 25
- 26 (2) TCEP;

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- (3) Decabromodiphenyl ether; 27
- (4) HBCD; 28
- (5) Additive TBBPA. 29
- NEW SECTION. Sec. 15. A new section is added to chapter 70.240 30 RCW to read as follows: 31
- (1)(a) By rule, the secretary of the department of health may 32 33 the manufacture, sale, distribution for sale, 34 distribution for use in this state of residential upholstered furniture, as defined in RCW 70.76.010, or children's products 35 containing a flame retardant in any product component in amounts 36 greater than one thousand parts per million that the department has 37

1 identified as a high priority chemical of high concern for children 2 pursuant to this chapter.

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- (b) After January 1, 2015, a rule that identifies a flame retardant as a high priority chemical of high concern for children must be adopted prior to December 1st of any year, and the restrictions under (a) of this subsection do not take effect before the end of the regular legislative session in the next year.
- (2) Before the department of health may adopt a rule to restrict a flame retardant that is a high priority chemical of high concern for children, the department of health must submit a report to the legislature no later than at the time of publication of the notice of a rule-making hearing required under RCW 34.05.320. The report to the legislature must contain:
- (a) A determination whether children or vulnerable populations are likely to be exposed to the chemical directly or indirectly from its use in products. The determination of the department must be made after an evaluation of available information on:
 - (i) Levels of the flame retardants in consumer products;
- (ii) Migration of the flame retardants out of products during and 19 20 after use; and
- 21 (iii) Levels of the flame retardants in humans environment, including but not limited to the home environment; 22
 - (b) A review of available toxicity data to evaluate the health concerns for children or vulnerable populations; and
 - (c) A determination of whether a safer alternative, as defined in section 1 of this act, has been identified to meet applicable fire safety standards for residential furniture and children's products by evaluating existing chemical action plans and assessments of safer alternatives that have been completed for flame retardant chemicals.
- The department of health must identify the sources of 30 31 information it relied upon in making the determination required in subsection (2) of this section, including peer-reviewed science. 32
 - (4) This section expires December 31, 2021.
- Sec. 16. RCW 70.240.010 and 2008 c 288 s 2 are each amended to 34 35 read as follows:
- The definitions in this section apply throughout this chapter 36 unless the context clearly requires otherwise. 37
- (1) "Children's cosmetics" means cosmetics that are made for, 38 marketed for use by, or marketed to children under the age of twelve. 39

- 1 "Children's cosmetics" includes cosmetics that meet any of the 2 following conditions:
- (a) Represented in its packaging, display, or advertising as 3 appropriate for use by children; 4
 - (b) Sold in conjunction with, attached to, or packaged together with other products that are packaged, displayed, or advertised as appropriate for use by children; or
 - (c) Sold in any of the following:
- (i) Retail store, catalogue, or online web site, in which a 9 person exclusively offers for sale products that are packaged, 10 11 displayed, or advertised as appropriate for use by children; or
 - (ii) A discrete portion of a retail store, catalogue, or online web site, in which a person offers for sale products that are packaged, displayed, or advertised as appropriate for use by children.
- (2) "Children's jewelry" means jewelry that is made for, marketed 16 17 for use by, or marketed to children under the age of twelve. "Children's jewelry" includes jewelry that meets any of the following 18 conditions: 19
- 20 (a) Represented in its packaging, display, or advertising as 21 appropriate for use by children under the age of twelve;
 - (b) Sold in conjunction with, attached to, or packaged together with other products that are packaged, displayed, or advertised as appropriate for use by children;
 - (c) Sized for children and not intended for use by adults; or
 - (d) Sold in any of the following:
 - (i) A vending machine;
- (ii) Retail store, catalogue, or online web site, in which a 28 person exclusively offers for sale products that are packaged, 29 displayed, or advertised as appropriate for use by children; or 30
- 31 (iii) A discrete portion of a retail store, catalogue, or online web site, in which a person offers for sale products that are 32 packaged, displayed, or advertised as appropriate for use by 33 34 children.
- 35 (3)(a) "Children's product" includes any of the following:
- 36 (i) Toys;

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- (ii) Children's cosmetics; 37
- (iii) Children's jewelry; 38

- (iv) A product designed or intended by the manufacturer to help a 1 child with sucking or teething, to facilitate sleep, relaxation, or 2 the feeding of a child, or to be worn as clothing by children; or 3
- (v) ((Child car seats)) A portable infant or child safety seat 4 designed to attach to an automobile seat. 5
 - (b) "Children's product" does not include the following:
- 7 (i) Batteries;

- (ii) Slings and catapults; 8
- (iii) Sets of darts with metallic points; 9
- (iv) Toy steam engines; 10
- 11 (v) Bicycles and tricycles;
- (vi) Video toys that can be connected to a video screen and are 12 13 operated at a nominal voltage exceeding twenty-four volts;
- (vii) Chemistry sets; 14
- (viii) Consumer electronic products, including but not limited to 15 personal computers, audio and video equipment, calculators, wireless 16 phones, game consoles, and hand-held devices incorporating a video 17 screen, used to access interactive software and their associated 18 19 peripherals;
- 20 Interactive software, intended for leisure and 21 entertainment, such as computer games, and their storage media, such 22 as compact disks;
- (x) BB guns, pellet guns, and air rifles; 23
- 24 (xi) Snow sporting equipment, including skis, poles, boots, snow boards, sleds, and bindings; 25
- 26 (xii) Sporting equipment, including, but not limited to bats, 27 balls, gloves, sticks, pucks, and pads;
- (xiii) Roller skates; 28
- 29 (xiv) Scooters;
- (xv) Model rockets; 30
- (xvi) Athletic shoes with cleats or spikes; and 31
- (xvii) Pocket knives and multitools. 32
- (4) "Cosmetics" includes articles intended to be rubbed, poured, 33 sprinkled, or sprayed on, introduced into, or otherwise applied to 34 35 the human body or any part thereof for cleansing, beautifying, 36 promoting attractiveness, or altering the appearance, and articles intended for use as a component of such an article. "Cosmetics" does 37 not include soap, dietary supplements, or food and drugs approved by 38 the United States food and drug administration. 39
- (5) "Department" means the department of ecology. 40

- 1 (6) "High priority chemical" means a chemical identified by a 2 state agency, federal agency, or accredited research university, or 3 other scientific evidence deemed authoritative by the department on 4 the basis of credible scientific evidence as known to do one or more 5 of the following:
- 6 (a) Harm the normal development of a fetus or child or cause other developmental toxicity;
 - (b) Cause cancer, genetic damage, or reproductive harm;
 - (c) Disrupt the endocrine system;

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- 10 (d) Damage the nervous system, immune system, or organs or cause 11 other systemic toxicity;
 - (e) Be persistent, bioaccumulative, and toxic; or
- 13 (f) Be very persistent and very bioaccumulative.
- (7) "Manufacturer" includes any person, firm, association, partnership, corporation, governmental entity, organization, or joint venture that produces a ((children's)) product or an importer or domestic distributor of a ((children's)) product. For the purposes of this subsection, "importer" means the owner of the ((children's)) product.
- 20 (8) "Phthalates" means di-(2-ethylhexyl) phthalate (DEHP), 21 dibutyl phthalate (DBP), benzyl butyl phthalate (BBP), diisonoyl 22 phthalate (DINP), diisodecyl phthalate (DIDP), or di-n-octyl 23 phthalate (DnOP).
- 24 (9) "Toy" means a product designed or intended by the 25 manufacturer to be used by a child at play.
 - (10) "Trade association" means a membership organization of persons engaging in a similar or related line of commerce, organized to promote and improve business conditions in that line of commerce and not to engage in a regular business of a kind ordinarily carried on for profit.
- 31 (11) "Very bioaccumulative" means having a bioconcentration 32 factor or bioaccumulation factor greater than or equal to five 33 thousand, or if neither are available, having a log Kow greater than 34 5.0.
- 35 (12) "Very persistent" means having a half-life greater than or 36 equal to one of the following:
- 37 (a) A half-life in soil or sediment of greater than one hundred 38 eighty days;
- 39 (b) A half-life greater than or equal to sixty days in water or 40 evidence of long-range transport.

- 1 (13) "TCEP" means the chemical (tris(2-chloroethyl)phosphate); 2 chemical abstracts service number 115-96-8, as of the effective date 3 of this section.
- 4 (14) "TDCPP" means the chemical (tris(1,3-dichloro-2-5 propyl)phosphate); chemical abstracts service number 13674-87-8, as 6 of the effective date of this section.
- 7 (15) "Decabromodiphenyl ether" means the chemical 8 decabromodiphenyl ether, chemical abstracts service number 1163-19-5, 9 as of the effective date of this section.
- 10 (16) "HBCD" means the chemical hexabromocyclododecane, chemical
 11 abstracts service number 25637-99-4, as of the effective date of this
 12 section.
- 13 (17) "Additive TBBPA" means the chemical tetrabromobisphenol A,
 14 chemical abstracts service number 79-94-7 as of the effective date of
 15 the section, in a form that has not undergone a reactive process and
 16 is not covalently bonded to a polymer in a product or product
 17 component.
- 18 **Sec. 17.** RCW 70.240.050 and 2008 c 288 s 7 are each amended to 19 read as follows:

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- (1) A manufacturer of products that are restricted under this chapter must notify persons that sell the manufacturer's products in this state about the provisions of this chapter no less than ninety days prior to the effective date of the restrictions.
- (2) A manufacturer that produces, sells, or distributes a product prohibited from manufacture, sale, or distribution in this state under this chapter shall recall the product and reimburse the retailer or any other purchaser for the product.
- (3) A manufacturer of ((children's)) products in violation of this chapter is subject to a civil penalty not to exceed five thousand dollars for each violation in the case of a first offense. Manufacturers who are repeat violators are subject to a civil penalty not to exceed ten thousand dollars for each repeat offense. Penalties collected under this section must be deposited in the state toxics control account created in RCW 70.105D.070.
- 35 (4) Retailers who unknowingly sell products that are restricted 36 from sale under this chapter are not liable under this chapter.
- 37 <u>(5) The sale or purchase of any previously owned products</u> 38 <u>containing a chemical restricted under this chapter made in casual or</u>

- isolated sales as defined in RCW 82.04.040, or by a nonprofit 1
- 2 organization, is exempt from this chapter.
- 3 NEW SECTION. Sec. 18. Sections 1 through 8 and 12 of this act
- constitute a new chapter in Title 70 RCW. 4
- 5 NEW SECTION. Sec. 19. This act may be known and cited as the
- 6 toxics reduction act.
- 7 NEW SECTION. Sec. 20. (1) Section 10 of this act expires June
- 8 30, 2019.
- 9 (2) Section 11 of this act expires June 30, 2025.
- NEW SECTION. Sec. 21. Section 11 of this act takes effect June 10
- 30, 2019. 11
- 12 NEW SECTION. Sec. 22. If specific funding for the purposes of
- this act, referencing this act by bill or chapter number, is not 13
- provided by June 30, 2015, in the omnibus appropriations act, this 14
- act is null and void. 15
- Sec. 23. If any provision of this act or its 16 NEW SECTION.
- 17 application to any person or circumstance is held invalid, the
- 18 remainder of the act or the application of the provision to other
- persons or circumstances is not affected." 19
- 20 Correct the title.

EFFECT: Makes the following changes to the Engrossed Second Substitute House Bill:

- Distinguishes between product "distributors" "manufacturers" by defining "distributors" as entities that import or distribute a product and by defining "manufacturers" as entities that manufacture a product, that control the manufacturing process, or that specify chemical composition of a product;
- (2) Exempts product distributors that do not control product formulation from department of ecology orders requiring distributors and manufacturers to assess alternatives to the use of a chemical in a product;
- (3) Exempts federally regulated food and beverage packaging, drug biological product packaging, components of FAA-regulated products, products produced under military specifications, and shipping pallets from the types of products that require a product manufacturer or distributor to respond to information requests and complete an alternatives assessment at the direction of the department of ecology;

- (4) Limits the number of chemicals that the department of ecology may select for chemical action plans to up to two every two years;
- (5) Requires that the first five chemicals for which the department of ecology initiates chemical action plans must be: (a) Chemicals regulated under the clean water act as human health criteria chemicals; or (b) chemicals that are currently identified by department rule as persistent, bioaccumulative toxics (PBTs);
- (6) Requires the department of ecology, when selecting chemicals for a chemical action plan, to consider the water quality impacts of the chemical, the impacts and extent of exposure to a chemical by humans and environmental biota, and to rank chemicals relative to one another based on information about the use, release, and exposure to the chemical;
- (7) Extends the period of time from six months to one year for manufacturers and distributors to respond to requests from the department of ecology for information about the use of targeted chemicals in their products;
- (8) Requires the department of ecology to consult with stakeholders if a chemical action plan external advisory committee has not yet been formed at the time of a request for information about chemical use to a manufacturer or distributor;
- (9) Authorizes manufacturers and distributors to use information that is not specific to Washington state and third-party data in responding to a request for information about a chemical by the department of ecology;
- (10) Requires the department of ecology to hold a product's manufacturer primarily responsible for responding to information requests if the manufacturer has a presence in the United States, and authorizes the department of ecology to hold a product distributor responsible for responding to the information request if the manufacturer does not have a presence in the United States;
- (11) Prohibits the department of ecology from penalizing a distributor of a product for failing to fulfill an information request if the distributor provides documentation that the distributor does not reasonably have access to the full set of information requested;
- (12) Specifies that a distributor's documentation that it does not reasonably have access to requested information must include copies of documents or written communications that demonstrate the distributor has requested relevant information from a manufacturer, a description of steps undertaken by the distributor to respond to the information request, and the signature of a representative attesting to the accuracy of the response;
- (13) Requires the department of ecology include in chemical action plans a prioritization of chemical exposure sources and releases based on human health, water quality, environmental impacts, and the cost and feasibility of actions to address sources and releases, and directs the department of ecology to consider this prioritization in making alternative assessment recommendations;
- (14) Limits the alternatives assessment authority of the department of ecology to two alternatives assessments recommended in chemical action plans, only one of which may be an alternatives assessment on a PBT chemical, plus an alternatives assessment for the use of polychlorinated biphenyls (PCBs) in pigments and dyes;
- (15) Limits the scope of the department of ecology's authority to require alternatives assessments on emerging chemicals that impact human health and the environment to an alternatives assessment for a single type of use of a chemical or a single type of product or product component containing the chemical, while limiting the scope of alternatives assessments on PBTs and water quality chemicals to no

greater breadth of uses, products, or manufacturers or distributors than is necessary to address sources of environmental or human exposures to the chemical;

- (16) Eliminates the requirement that the department of ecology complete a summary report at the end of the alternatives assessment process for a chemical;
- (17) Authorizes, rather than requires, the department of ecology to submit agency request legislation if an alternatives assessment identifies safer alternatives to the use of a chemical;
- (18) Expands the scope of the state purchasing preference for chemicals that do not contain PBTs to encompass any chemicals that the department of ecology identifies by rule as PBTs, rather than chemicals that the department of ecology has identified as PBTs by rule as of the act's 2015 effective date;
- (19) Restricts the use of the following five flame retardant chemicals in children's products and residential upholstered furniture in amounts greater than 1,000 parts per million, beginning July 1, 2016: TDCPP, TCEP, Decabromodiphenyl ether, HBCD, and additive TBBPA that has not covalently bonded with a polymer;
- (20) Authorizes the department of health to restrict, by rule, flame retardants that the department of ecology identifies as a chemical of high concern for children, in amounts greater than 1,000 parts per million in children's products and residential upholstered furniture;
- (21) Requires the department of ecology to complete the adoption of a rule to identify a flame retardant as a high priority chemical of high concern for children by December 1st of a given year, and makes the effective date of any department of health rule to restrict the flame retardant effective no sooner than the end of the regular legislative session the following year;
- (22) Requires the department of health, prior to adopting a rule to restrict a flame retardant, to submit a report to the legislature regarding the human health impacts and exposure pathways for the flame retardant; and
- (23) Adds an intent section noting the previous evaluations and legislative actions that the state has taken with respect to certain flame retardant chemicals.

--- END ---