

2SHB 1472 - H AMD TO H AMD (H-2198.5/15) 265  
By Representative Shea

WITHDRAWN 3/11/2015

1 Beginning on page 1, line 3 of the amendment, strike all material  
2 through page 16 and insert the following:

3 "NEW SECTION. **Sec. 1.** The definitions in this section apply  
4 throughout this chapter unless the context clearly requires  
5 otherwise.

6 (1) "Alternatives assessment" means a process for identifying and  
7 comparing chemical and nonchemical alternatives currently in  
8 existence that can be practicably and economically used to replace  
9 the use of a chemical or to reduce the amount of or exposure to that  
10 chemical. The objective of an alternatives assessment is to assess  
11 less toxic chemicals or nonchemical alternatives to reduce the amount  
12 of or exposure to the chemical in a product and to avoid the  
13 unintended consequence of switching to a substitute that presents an  
14 equivalent or greater concern. An alternatives assessment must follow  
15 the guidelines issued by the interstate chemicals clearinghouse, the  
16 national academy of sciences, or equivalent methodology. At a  
17 minimum, an alternatives assessment includes: An evaluation of  
18 chemical hazard, exposure, performance, consumer acceptance, cost,  
19 and availability; information for each alternative considered; and  
20 the identification of alternatives.

21 (2) "Biomonitoring" means assessment of human exposures to  
22 chemicals by measuring the chemicals or their metabolites in human  
23 tissues or specimens, such as blood, breast milk, and urine.

24 (3) "Chemical" means a substance, including metals, with a  
25 distinct molecular composition or a group of structurally related  
26 substances and includes the breakdown products of the substance or  
27 substances that form through decomposition, degradation, or  
28 metabolism.

29 (4) "Chemical action plan" means a plan that identifies,  
30 characterizes, and evaluates uses and releases of a specific chemical  
31 or group of chemicals and identifies actions needed to protect human  
32 health and the environment.

33 (5) "Department" means the department of ecology.

1 (6) "Director" means the director of the department of ecology or  
2 the director's designee.

3 (7) "Manufacturer" means any person, firm, association,  
4 partnership, corporation, governmental entity, organization, or joint  
5 venture that produces a product sold or offered for sale in or into  
6 the state. "Manufacturer" does not include small businesses as  
7 defined in RCW 19.85.020.

8 (8) "Product" means any item sold for residential or commercial  
9 use including any component or product packaging. "Product" does not  
10 include the following items, but does include their packaging, except  
11 as provided in (a) and (c) of this subsection:

12 (a) Food or beverage, and food and beverage packaging, regulated  
13 by the United States food and drug administration or the United  
14 States department of agriculture;

15 (b) Tobacco products;

16 (c) Drug or biological products regulated by the United States  
17 food and drug administration;

18 (d) Products produced under military specifications;

19 (e) Finished products regulated by the federal aviation  
20 administration;

21 (f) Substances regulated under chapter 15.54 or 15.58 RCW; and

22 (g) Any previously owned product sold in casual or isolated sales  
23 as defined in RCW 82.04.040 or products sold by nonprofit  
24 organizations.

25 (9) "Product component" means a uniquely identifiable material or  
26 coating that is included as a part of a finished product.

27 (10) "Safer alternative" means an alternative that is  
28 demonstrated by an alternatives assessment to meet improved hazard  
29 and exposure considerations that possess lower risk and can be  
30 practicably and economically substituted for the original chemical or  
31 allow use of a reduced amount of or exposure to that chemical than  
32 the existing chemical or chemical process. A safer alternative to a  
33 particular chemical may include a chemical substitute or a change in  
34 materials or design that eliminates the need for a chemical  
35 alternative.

36 (11) "Summary report" means a report prepared by the department  
37 summarizing available alternatives assessments and includes a  
38 determination regarding the existence of a safer alternative. The  
39 summary report also includes a determination of the completeness of

1 the alternatives assessments reviewed and identifies unsuitable  
2 alternatives.

3 (12) "Unsuitable alternative" means an alternative identified  
4 through the alternatives assessment process that does not meet the  
5 hazard, exposure, cost, performance, and availability criteria of a  
6 safer alternative.

7 NEW SECTION. **Sec. 2.** (1) Beginning January 1, 2016, and every  
8 two years thereafter, the department, in consultation with the  
9 department of health, must select up to two chemicals for the  
10 development of chemical action plans as specified in section 4 of  
11 this act from the following:

12 (a) Chemicals regulated by the department as human health  
13 criteria in the draft rule issued by the department on January 8,  
14 2015, and filed as Washington State Register 15-03-15; or

15 (b) Chemicals that are persistent, bioaccumulative, and toxic  
16 chemicals as defined in chapter 173-333 WAC, as of the effective date  
17 of this section, that affect water quality.

18 (2) The department may conduct environmental monitoring or,  
19 subject to the availability of amounts appropriated for this specific  
20 purpose, may request the department of health to conduct  
21 biomonitoring of a chemical to verify the chemical is present in the  
22 state's environment or population or to better understand  
23 environmental or human exposures in the state. Environmental  
24 monitoring and biomonitoring conducted pursuant to this chapter must  
25 be of a minimum scope necessary to adequately inform a chemical  
26 action plan.

27 (3)(a) At least two of the first four chemicals selected for a  
28 chemical action plan must be chosen from the chemicals identified in  
29 subsection (1)(a) of this section.

30 (b) When selecting chemicals for the development of chemical  
31 action plans, the director shall notify the public of the selection,  
32 the basis for the selection, and a draft schedule. The notice must be  
33 published in the Washington State Register. The department shall  
34 provide the public with an opportunity for review and comment before  
35 finalizing the schedule.

36 (c) When selecting chemicals for the development of chemical  
37 action plans, the department must consider:

38 (i) Opportunities for reducing or phasing out uses, production,  
39 or releases of a chemical;

1 (ii) Current scientific evidence on the combined effects of  
2 exposure to the chemical and other substances commonly present in the  
3 Washington environment;

4 (iii) Current scientific evidence on the susceptibility of  
5 sensitive population groups and environmental media from exposure to  
6 the chemical, as well as cumulative effects of multiple exposures;

7 (iv) The relative ranking assigned to a chemical by the  
8 department based on information applicable to Washington state about  
9 chemical characteristics, uses of the chemical, releases of the  
10 chemical, and levels of the chemical present in the environment and  
11 in residents;

12 (v) Whether the chemical has been determined to impact Washington  
13 state waters through identification under section 303(d) of the  
14 federal clean water act; and

15 (vi) Existing plans or regulatory requirements to reduce or phase  
16 out the use and releases of the chemical.

17 (d) The department must identify the sources of information it  
18 relied upon in selecting chemicals for the development of chemical  
19 action plans under this section, including peer-reviewed science.

20 NEW SECTION. **Sec. 3.** (1) The department may require information  
21 from manufacturers of products that contain a chemical selected for a  
22 chemical action plan under section 2 of this act. Prior to requesting  
23 information from a manufacturer under this subsection, the department  
24 must consult with a chemical action plan external advisory committee,  
25 if one has been formed yet, to evaluate the particular chemical that  
26 is the subject of the information request. The department may only  
27 make reasonable requests of manufacturers that are limited in their  
28 scope and frequency and that are focused on:

29 (a) The most common and prevalent uses of the chemicals or  
30 products containing the chemicals, based on the department's existing  
31 knowledge about the chemical;

32 (b) Areas where there is an identified gap in public or  
33 department knowledge about a chemical; and

34 (c) Chemical uses or products that the department has reason to  
35 believe are likely to be responsible for or associated with a  
36 significant portion of releases into the environment or public health  
37 exposures.

38 (2) Within twelve months of a request by the department,  
39 manufacturers shall report the following:

1 (a) The name and address of the manufacturer and the name,  
2 address, and phone number of a contact person for the manufacturer;

3 (b) The name of the chemical used or produced and its chemical  
4 abstracts service registry number;

5 (c) A brief description of the product or product component  
6 categories containing the substance;

7 (d) A description of the function or functions of the chemical in  
8 the product;

9 (e) An estimate of average daily, weekly, or monthly commercial  
10 consumption of the chemical by businesses or the public; and

11 (f) Any other information the manufacturer deems relevant to the  
12 appropriate use of the product.

13 (3) In response to an information request from the department  
14 under this section, a manufacturer may extrapolate amounts and  
15 estimates from national data. The resulting submission must include  
16 the information in subsection (2)(a) of this section for each  
17 manufacturer. However, the information required by subsection (2)(b)  
18 through (f) of this section is not required to be provided in a  
19 manner that identifies individual manufacturers.

20 (4) The department shall specify the required format for  
21 submission of the information required under subsection (2) of this  
22 section. The format should be generally consistent with the format  
23 specified in other states with substantially similar reporting  
24 requirements.

25 (5) Multiple businesses, or a business association, may  
26 collaborate and submit a single submission on a chemical found in  
27 similar products.

28 (6) Where information submitted by a manufacturer under chapter  
29 70.240 RCW is the same as the information required to be submitted by  
30 the manufacturer in subsection (2) of this section, that manufacturer  
31 is not required to submit the same information again.

32 (7) The department may, by order, require a manufacturer subject  
33 to the reporting requirement in subsection (2) of this section to  
34 provide additional information that is relevant to the development of  
35 a chemical action plan under section 4 of this act. Prior to an order  
36 under this subsection, the department must consult with the external  
37 advisory committee formed for the chemical action plan, if one has  
38 been formed yet. An order by the department must also meet the  
39 reasonableness criteria of subsection (1) of this section.

1        NEW SECTION.    **Sec. 4.**    (1) When developing a chemical action  
2 plan, the department shall convene an external advisory committee to  
3 provide stakeholder input, expertise, and additional information. All  
4 advisory committee meetings must be open to the public. The  
5 department must invite representatives from, at minimum, the  
6 following organizations and entities to serve as external advisory  
7 committee members: Large and small business sectors; a representative  
8 of a statewide business association with over one thousand total  
9 members and that represents multiple business sectors; community,  
10 environmental, and public health advocacy groups; local governments;  
11 affected and interested businesses; and public health agencies. State  
12 agencies and technical experts may be requested to participate.

13        (2) All chemical action plans must include the following types of  
14 information, evaluations, and recommendations:

15        (a) Chemical name, properties, uses, and product manufacturers;

16        (b) An analysis of the available information on the production,  
17 unintentional production, uses, and disposal of the chemical;

18        (c) Information on the known or potential and proven impacts on  
19 human health and the environment associated with the use and release  
20 of the chemical; and

21        (d) An evaluation of the regulatory and nonregulatory approaches  
22 that influence production, uses, releases, and management of the  
23 chemical.

24        (3)(a) All chemical action plans must identify actions, if  
25 needed, to eliminate, reduce, or manage exposures and include  
26 recommendations for managing, reducing, or phasing out the uses and  
27 releases of the chemicals identified as primary sources of risk to  
28 human health or the environment in Washington state to minimize  
29 exposure.

30        (b) Recommendations must be based on an evaluation of the  
31 following factors:

32        (i) Opportunity for environmental and human health benefits in  
33 Washington state;

34        (ii) Economic and social impacts;

35        (iii) Feasibility;

36        (iv) Availability and effectiveness of safer substitutes for uses  
37 of the chemical; and

38        (v) Consistency with existing federal and state regulatory  
39 requirements.

1 (4) The department must include in the chemical action plan a  
2 summary of any dissenting views held by external advisory committee  
3 members regarding the recommendations contained in the plan.

4 (5) The department must identify the sources of information it  
5 relied upon in completing a chemical action plan under this section,  
6 including peer-reviewed science.

7 NEW SECTION. **Sec. 5.** (1)(a) Consistent with a recommendation in  
8 a chemical action plan, the department is authorized to require  
9 manufacturers, by order, to conduct alternatives assessments, as  
10 detailed in this section. The department may not require  
11 manufacturers to complete an alternatives assessment for a greater  
12 breadth of uses or products, nor require alternatives assessments to  
13 be completed by a greater number of manufacturers, than is necessary  
14 to address demonstrated statistically significant sources of  
15 environmental or public health hazard and exposures to the chemical.

16 (b) The scope of an alternatives assessment request must be:

17 (i) A single type of use of a chemical in a specific type of  
18 manufacturing process; or

19 (ii) The inclusion of a chemical in a specific type of product.

20 (2)(a) If ordered by the department, a manufacturer of a product  
21 that contains a chemical for which a chemical action plan has been  
22 completed under section 4 of this act or under chapter 173-333 WAC  
23 must submit an alternatives assessment to the department for each use  
24 of the chemical specified by the department.

25 (b) The manufacturer must submit the alternatives assessment to  
26 the department within twenty-four months of receipt of the  
27 department's order; however, the department may grant an extension on  
28 a case-by-case basis for good cause if the manufacturer shows that  
29 additional time is necessary to complete an alternatives assessment  
30 or would substantially improve the quality of the alternatives  
31 assessment. Multiple businesses, or a business association, may  
32 collaborate and submit a single alternatives assessment on a chemical  
33 found in similar products.

34 (c) In lieu of an alternatives assessment, a manufacturer may  
35 submit a certificate of compliance, as described in (d) of this  
36 subsection, if:

37 (i) The manufacturer has ceased using the chemical for which it  
38 would be required to do an alternatives assessment; or

1 (ii) The manufacturer can demonstrate its plans to phase out the  
2 use of the chemical within a time frame that is reasonable based on  
3 the manufacturing process used to produce the product and the use of  
4 the product.

5 (d) A certificate of compliance must include the following:

6 (i) Chemical names and chemical abstracts service registry  
7 numbers for all chemicals that currently contribute to the specific  
8 function previously served by the prohibited chemical;

9 (ii) How the manufacturer is meeting the function of the  
10 prohibited chemical with a safer alternative; and

11 (iii) The signature of an authorized official of the  
12 manufacturer.

13 (3) If the department determines that a submitted alternatives  
14 assessment does not meet the definition or required objectives of an  
15 alternatives assessment, or the department does not identify a  
16 manufacturer that may be required to submit an alternatives  
17 assessment, the department may contract with an independent  
18 scientific organization to conduct an independent alternatives  
19 assessment in consultation with the chemical action plan advisory  
20 committee. Any alternatives assessment conducted by the independent  
21 contractor must include a process to involve interested parties.

22 NEW SECTION. **Sec. 6.** (1)(a) The department, in consultation  
23 with the department of health, shall prepare a summary report of all  
24 reviewed alternatives assessments and other relevant information  
25 assembled under section 5 of this act. The summary report must  
26 include a determination of whether a safer alternative exists and  
27 identify unsuitable alternatives.

28 (b) In making its determination, the department shall evaluate  
29 whether the alternatives assessment submitted by manufacturers  
30 follows the guidelines on alternatives assessment issued by the  
31 interstate chemicals clearinghouse, the national academy of sciences,  
32 or equivalent methodology.

33 (2) If the department determines that the large majority of  
34 alternatives assessments support a safer alternative, the department  
35 may submit a recommendation to prohibit specific uses of the  
36 chemical, in the form of draft legislation, to the appropriate  
37 committees of the house of representatives and senate.

38 (3) If the department determines that a safer alternative does  
39 not exist, then the department may reevaluate information on the



1 availability of safer alternatives not more often than once every  
2 five years.

3 NEW SECTION. **Sec. 7.** (1) A manufacturer violating a requirement  
4 of this chapter, a rule adopted under this chapter, or an order  
5 issued under this chapter, is subject to a civil penalty not to  
6 exceed five thousand dollars for each violation in the case of a  
7 first offense. Manufacturers who are repeat violators are subject to  
8 a civil penalty not to exceed ten thousand dollars for each repeat  
9 offense.

10 (2) Any penalty provided for in this section, and any order  
11 issued by the department under this chapter, maybe appealed to the  
12 pollution control hearings board.

13 (3) All penalties collected under this chapter shall be deposited  
14 in the state toxics control account created in RCW 70.105D.070.

15 NEW SECTION. **Sec. 8.** (1) Manufacturers submitting information  
16 or records to the department may request that the information or  
17 records be made available only for the confidential use of the  
18 director, the department, or the appropriate division of the  
19 department. The director shall give consideration to the request and  
20 if such action would not be detrimental to the public interest and is  
21 otherwise within accord with the policies and purposes of chapter  
22 43.21A RCW, the director must grant the request for the information  
23 to remain confidential as authorized in RCW 43.21A.160. The  
24 department must keep confidential any records furnished by a  
25 manufacturer under this chapter that relate to proprietary  
26 manufacturing processes or chemical formulations used in products or  
27 processes.

28 (2) Within three working days of receipt of a request, the  
29 department must send a letter to a manufacturer that asks for  
30 information demonstrating how the records relate to processes of  
31 production unique to the owner or operator or how releasing the  
32 records to the public may adversely affect the owner's or operator's  
33 competitive position. If the manufacturer does not respond to this  
34 information within fourteen days, the department is no longer  
35 required to treat the submitted information or records as  
36 confidential. If the manufacturer responds within fourteen days with  
37 the requested information, the department must inform the  
38 manufacturer with its determination of whether the submitted

1 information should be kept confidential under this section and RCW  
2 43.21A.160.

3 (3) If the director denies the request of a manufacturer to keep  
4 submitted information or records confidential under this section, the  
5 manufacturer may appeal that denial to a court of competent  
6 jurisdiction. In a review of whether the submitted information or  
7 records meet the criteria of RCW 43.21A.160 and this section, a court  
8 must examine submitted information or records in camera.

9 NEW SECTION. **Sec. 9.** The department may adopt rules as  
10 necessary for the purpose of implementing, administering, and  
11 enforcing this chapter.

12 NEW SECTION. **Sec. 10.** A new section is added to chapter 39.26  
13 RCW to read as follows:

14 (1) The department shall establish purchasing and procurement  
15 policies that provide a preference for products and products in  
16 packaging that do not contain:

17 (a) Persistent, bioaccumulative, and toxic chemicals as defined  
18 in chapter 173-333 WAC as of the effective date of this section; and

19 (b) Chemicals that have been addressed by a completed chemical  
20 action plan that has included a recommendation that the state adopt a  
21 purchasing and procurement policy for products and products in  
22 packaging that do not contain the chemical.

23 (2) No agency may knowingly purchase products or products in  
24 packaging containing chemicals identified in subsection (1) of this  
25 section unless there is no cost-effective and technologically  
26 feasible alternative. When all available products contain a chemical  
27 identified in subsection (1) of this section, a preference must be  
28 given to alternative products that contain lesser amounts of  
29 chemicals identified in subsection (1) of this section.

30 (3) Nothing in this section requires the department or any other  
31 state agency to breach an existing contract or dispose of stock that  
32 has been ordered or is in the possession of the department or other  
33 state agency as of the effective date of this section.

34 (4) This section does not require the department or any other  
35 agency to test every product procured.

36 (5) The department or any other agency may request suppliers of  
37 products to provide testing data from an accredited laboratory or  
38 testing facility documenting levels of a chemical identified in

1 subsection (1) of this section in products or product packaging.  
2 Requested or voluntarily received testing data from businesses,  
3 manufacturers, organizations, and individuals must be submitted for  
4 review to the department of ecology.

5 **Sec. 11.** RCW 43.21B.110 and 2013 c 291 s 33 are each amended to  
6 read as follows:

7 (1) The hearings board shall only have jurisdiction to hear and  
8 decide appeals from the following decisions of the department, the  
9 director, local conservation districts, the air pollution control  
10 boards or authorities as established pursuant to chapter 70.94 RCW,  
11 local health departments, the department of natural resources, the  
12 department of fish and wildlife, the parks and recreation commission,  
13 and authorized public entities described in chapter 79.100 RCW:

14 (a) Civil penalties imposed pursuant to RCW 18.104.155,  
15 70.94.431, 70.105.080, 70.107.050, 76.09.170, 77.55.291, 78.44.250,  
16 88.46.090, 90.03.600, 90.46.270, 90.48.144, 90.56.310, 90.56.330, and  
17 90.64.102.

18 (b) Orders issued pursuant to RCW 18.104.043, 18.104.060,  
19 43.27A.190, 70.94.211, 70.94.332, 70.105.095, 86.16.020, 88.46.070,  
20 90.14.130, 90.46.250, 90.48.120, and 90.56.330.

21 (c) A final decision by the department or director made under  
22 chapter 183, Laws of 2009.

23 (d) Except as provided in RCW 90.03.210(2), the issuance,  
24 modification, or termination of any permit, certificate, or license  
25 by the department or any air authority in the exercise of its  
26 jurisdiction, including the issuance or termination of a waste  
27 disposal permit, the denial of an application for a waste disposal  
28 permit, the modification of the conditions or the terms of a waste  
29 disposal permit, or a decision to approve or deny an application for  
30 a solid waste permit exemption under RCW 70.95.300.

31 (e) Decisions of local health departments regarding the grant or  
32 denial of solid waste permits pursuant to chapter 70.95 RCW.

33 (f) Decisions of local health departments regarding the issuance  
34 and enforcement of permits to use or dispose of biosolids under RCW  
35 70.95J.080.

36 (g) Decisions of the department regarding waste-derived  
37 fertilizer or micronutrient fertilizer under RCW 15.54.820, and  
38 decisions of the department regarding waste-derived soil amendments  
39 under RCW 70.95.205.

1 (h) Decisions of local conservation districts related to the  
2 denial of approval or denial of certification of a dairy nutrient  
3 management plan; conditions contained in a plan; application of any  
4 dairy nutrient management practices, standards, methods, and  
5 technologies to a particular dairy farm; and failure to adhere to the  
6 plan review and approval timelines in RCW 90.64.026.

7 (i) Any other decision by the department or an air authority  
8 which pursuant to law must be decided as an adjudicative proceeding  
9 under chapter 34.05 RCW.

10 (j) Decisions of the department of natural resources, the  
11 department of fish and wildlife, and the department that are  
12 reviewable under chapter 76.09 RCW, and the department of natural  
13 resources' appeals of county, city, or town objections under RCW  
14 76.09.050(7).

15 (k) Forest health hazard orders issued by the commissioner of  
16 public lands under RCW 76.06.180.

17 (l) Decisions of the department of fish and wildlife to issue,  
18 deny, condition, or modify a hydraulic project approval permit under  
19 chapter 77.55 RCW.

20 (m) Decisions of the department of natural resources that are  
21 reviewable under RCW 78.44.270.

22 (n) Decisions of an authorized public entity under RCW 79.100.010  
23 to take temporary possession or custody of a vessel or to contest the  
24 amount of reimbursement owed that are reviewable by the hearings  
25 board under RCW 79.100.120.

26 (o) Decisions regarding a restriction, order, or penalty issued  
27 under chapter 70.--- RCW (the new chapter created in section 14 of  
28 this act).

29 (2) The following hearings shall not be conducted by the hearings  
30 board:

31 (a) Hearings required by law to be conducted by the shorelines  
32 hearings board pursuant to chapter 90.58 RCW.

33 (b) Hearings conducted by the department pursuant to RCW  
34 70.94.332, 70.94.390, 70.94.395, 70.94.400, 70.94.405, 70.94.410, and  
35 90.44.180.

36 (c) Appeals of decisions by the department under RCW 90.03.110  
37 and 90.44.220.

38 (d) Hearings conducted by the department to adopt, modify, or  
39 repeal rules.

1 (3) Review of rules and regulations adopted by the hearings board  
2 shall be subject to review in accordance with the provisions of the  
3 administrative procedure act, chapter 34.05 RCW.

4 **Sec. 12.** RCW 43.21B.110 and 2013 c 291 s 34 are each amended to  
5 read as follows:

6 (1) The hearings board shall only have jurisdiction to hear and  
7 decide appeals from the following decisions of the department, the  
8 director, local conservation districts, the air pollution control  
9 boards or authorities as established pursuant to chapter 70.94 RCW,  
10 local health departments, the department of natural resources, the  
11 department of fish and wildlife, the parks and recreation commission,  
12 and authorized public entities described in chapter 79.100 RCW:

13 (a) Civil penalties imposed pursuant to RCW 18.104.155,  
14 70.94.431, 70.105.080, 70.107.050, 76.09.170, 77.55.291, 78.44.250,  
15 88.46.090, 90.03.600, 90.46.270, 90.48.144, 90.56.310, 90.56.330, and  
16 90.64.102.

17 (b) Orders issued pursuant to RCW 18.104.043, 18.104.060,  
18 43.27A.190, 70.94.211, 70.94.332, 70.105.095, 86.16.020, 88.46.070,  
19 90.14.130, 90.46.250, 90.48.120, and 90.56.330.

20 (c) Except as provided in RCW 90.03.210(2), the issuance,  
21 modification, or termination of any permit, certificate, or license  
22 by the department or any air authority in the exercise of its  
23 jurisdiction, including the issuance or termination of a waste  
24 disposal permit, the denial of an application for a waste disposal  
25 permit, the modification of the conditions or the terms of a waste  
26 disposal permit, or a decision to approve or deny an application for  
27 a solid waste permit exemption under RCW 70.95.300.

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

30 (e) Decisions of local health departments regarding the issuance  
31 and enforcement of permits to use or dispose of biosolids under RCW  
32 70.95J.080.

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34 fertilizer or micronutrient fertilizer under RCW 15.54.820, and  
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38 denial of approval or denial of certification of a dairy nutrient  
39 management plan; conditions contained in a plan; application of any

1 dairy nutrient management practices, standards, methods, and  
2 technologies to a particular dairy farm; and failure to adhere to the  
3 plan review and approval timelines in RCW 90.64.026.

4 (h) Any other decision by the department or an air authority  
5 which pursuant to law must be decided as an adjudicative proceeding  
6 under chapter 34.05 RCW.

7 (i) Decisions of the department of natural resources, the  
8 department of fish and wildlife, and the department that are  
9 reviewable under chapter 76.09 RCW, and the department of natural  
10 resources' appeals of county, city, or town objections under RCW  
11 76.09.050(7).

12 (j) Forest health hazard orders issued by the commissioner of  
13 public lands under RCW 76.06.180.

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15 deny, condition, or modify a hydraulic project approval permit under  
16 chapter 77.55 RCW.

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20 to take temporary possession or custody of a vessel or to contest the  
21 amount of reimbursement owed that are reviewable by the hearings  
22 board under RCW 79.100.120.

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24 under chapter 70.--- RCW (the new chapter created in section 14 of  
25 this act).

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30 (b) Hearings conducted by the department pursuant to RCW  
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36 repeal rules.

37 (3) Review of rules and regulations adopted by the hearings board  
38 shall be subject to review in accordance with the provisions of the  
39 administrative procedure act, chapter 34.05 RCW.

1        NEW SECTION.    **Sec. 13.**    If specific funding for the purposes of  
2 this act, referencing this act by bill or chapter number, is not  
3 provided by June 30, 2015, in the omnibus appropriations act, this  
4 act is null and void.

5        NEW SECTION.    **Sec. 14.**    Sections 1 through 9 of this act  
6 constitute a new chapter in Title 70 RCW.

7        NEW SECTION.    **Sec. 15.**    A new section is added to chapter 43.131  
8 RCW to read as follows:

9        The authority of the department of ecology to do the following  
10 under the authority of chapter 70.--- RCW (the new chapter created in  
11 section 14 of this act) expires June 30, 2025: Require manufacturers  
12 to provide information on chemicals and conduct alternatives  
13 assessments; prepare summary reports on alternatives assessments;  
14 prohibit the use of chemicals and the sale, offer for sale, or  
15 distribution of a product containing a prohibited chemical; and  
16 assess penalties.

17        NEW SECTION.    **Sec. 16.**    A new section is added to chapter 43.131  
18 RCW to read as follows:

19        The following acts or parts of acts, as now existing or hereafter  
20 amended, are each repealed, effective June 30, 2026:

- 21        (1) Section 1 of this act;
- 22        (2) Section 2 of this act;
- 23        (3) Section 3 of this act;
- 24        (4) Section 4 of this act;
- 25        (5) Section 5 of this act;
- 26        (6) Section 6 of this act;
- 27        (7) Section 7 of this act;
- 28        (8) Section 8 of this act; and
- 29        (9) Section 9 of this act.

30        NEW SECTION.    **Sec. 17.**    This act may be known and cited as the  
31 toxics reduction act.

32        NEW SECTION.    **Sec. 18.**    Section 11 of this act expires June 30,  
33 2019.

1        NEW SECTION.    **Sec. 19.**    Section 12 of this act takes effect June  
2    30, 2019.

3        NEW SECTION.    **Sec. 20.**    If any provision of this act or its  
4    application to any person or circumstance is held invalid, the  
5    remainder of the act or the application of the provision to other  
6    persons or circumstances is not affected."

7        Correct the title.

EFFECT: (1) Reduces the maximum number of chemicals subject to chemical action plans from four to two every two years.

(2) Changes the scope of chemicals regulated under the federal clean water act that are subject to chemical action plans to specifically include chemicals regulated as human health criteria under the draft water quality standards rule published by the department of ecology on January 12, 2015.

(3) Eliminates authority for the department of ecology to complete chemical action plans for chemicals present in the human or natural environment and that meet other exposure, hazard, and health effect criteria, and instead authorizes the department to complete chemical action plans for chemicals identified by rule by the department as persistent, bioaccumulative, and toxic.

(4) Redefines the outcome of an alternatives assessment from replacing the use of a chemical to reducing the amount of or exposures to the chemical.

(5) Requires alternatives assessments to consider consumer acceptance of a product among the evaluated criteria.

(6) Exempts importers and domestic distributors from the obligations placed on manufacturers under the act.

(7) Exempts all chemicals regulated under state fertilizer or pesticide laws from the act, rather than exempting chemical products used to produce an agricultural commodity.

(8) Authorizes a manufacturer whose request to keep submitted information confidential is denied by the department of ecology to appeal that denial to a court of competent jurisdiction, and requires *in camera* review by the court of the relevant records and information.

(9) Extends the amount of time for manufacturers to respond to an information request from the department of ecology from six months to twelve months.

(10) Eliminates the requirement that manufacturers, upon request by the department of ecology, report the amount of a chemical in a product or product component.

(11) Requires the department of ecology to consider a chemical's relative risks as compared to other chemicals and whether the chemical impacts state water quality when selecting chemicals that will be subject to chemical action plans.

(12) Authorizes, rather than requires, the department of ecology to recommend, in the form of draft legislation, that the legislature prohibit specific uses of a chemical for which the department has identified a safer alternative.

(13) Requires the department of ecology to consult with a chemical action plan external advisory committee, if formed, prior to making an information request from manufacturers.



(14) Extends the time frame for a manufacturer to respond to an ordered alternatives assessment from one year to twenty-four months.

(15) Exempts federally regulated food, beverage, drug, and biological product packaging from the requirements of the act.

(16) Eliminates the authority for the department of ecology to use equivalent information to an alternatives assessment in order to identify a safer alternative to a chemical.

(17) Requires the department of ecology to determine that a large majority of alternatives assessments support a safer alternative prior to the department recommending to the legislature any restrictions on the use of the chemical.

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