

## SENATE MINORITY REPORT AMENDMENTS TO SENATE BILL 478

By Nonconcurring Members of COMMITTEE ON ENVIRONMENT AND NATURAL  
RESOURCES

April 27

1 On page 1 of the printed bill, line 2, delete “; and declaring an emergency”.

2 Delete lines 4 through 22 and delete pages 2 through 10 and insert:

3  
4 **“DEFINITIONS**

5  
6 **“SECTION 1. As used in sections 1 to 10 of this 2015 Act:**

7 **“(1) ‘Chemical’ means:**

8 **“(a) A substance with a distinct molecular composition and the breakdown products of**  
9 **the substance that form through decomposition, degradation or metabolism.**

10 **“(b) A group of structurally related substances and the breakdown products of the sub-**  
11 **stances that form through decomposition, degradation or metabolism.**

12 **“(2)(a) ‘Children’s cosmetics’ means products that are intended to be rubbed, poured,**  
13 **sprinkled or sprayed on, introduced into or otherwise applied to the human body or any part**  
14 **thereof for cleansing, moisturizing, beautifying, promoting attractiveness or altering the**  
15 **appearance, and articles intended for use as a component of such products.**

16 **“(b) ‘Children’s cosmetics’ does not mean soap, dietary supplements or food and drugs**  
17 **approved by the United States Food and Drug Administration.**

18 **“(3)(a) ‘Children’s product’ means any of the following products that are made for, mar-**  
19 **keted for use by or marketed to children under 12 years of age:**

20 **“(A) Products designed or intended by the manufacturer to facilitate sucking, teething,**  
21 **sleep, relaxation or feeding.**

22 **“(B) Children’s clothing.**

23 **“(C) Car seats.**

24 **“(D) Children’s cosmetics.**

25 **“(E) Children’s jewelry.**

26 **“(F) Toys.**

27 **“(b) ‘Children’s product’ does not mean:**

28 **“(A) Inaccessible components of a product specified in paragraph (a) of this subsection**  
29 **that during reasonably foreseeable use and abuse of the product would not come into direct**  
30 **contact with a child’s skin or mouth.**

31 **“(B) Used products specified in paragraph (a) of this subsection that are sold in second-**  
32 **hand product markets.**

33 **“(C) Athletic shoes with cleats or spikes.**

34 **“(D) Batteries.**

1       **“(E) BB guns, pellet guns and air rifles.**

2       **“(F) Bicycles and tricycles.**

3       **“(G) Chemistry sets.**

4       **“(H) Consumer electronic products, including personal computers, audio and video**  
5 **equipment, calculators, wireless telephones and game consoles, handheld devices that incor-**  
6 **porate a video screen and are used to access interactive software, and the associated pe-**  
7 **ripherals.**

8       **“(I) Interactive software intended for leisure and entertainment, such as computer**  
9 **games, and their storage media, such as compact discs.**

10       **“(J) Model rockets.**

11       **“(K) Pocketknives and multitools.**

12       **“(L) Roller skates.**

13       **“(M) Scooters.**

14       **“(N) Sets of darts with metallic points.**

15       **“(O) Slings and catapults.**

16       **“(P) Snow sporting equipment, including skis, poles, boots, snowboards, sleds and**  
17 **bindings.**

18       **“(Q) Sporting equipment, including bats, balls, gloves, sticks, pucks and pads.**

19       **“(R) Video toys that can be connected to a video screen and are operated at a nominal**  
20 **voltage exceeding 24 volts.**

21       **“(S) Food and beverages, and food and beverage packaging, regulated by the United**  
22 **States Food and Drug Administration or the United States Department of Agriculture.**

23       **“(T)(i) Drug and biologics regulated by the United States Food and Drug Administration**  
24 **that are over-the-counter drugs, prescription drugs, dietary supplements, medical devices or**  
25 **products that are both a cosmetic and a drug; and**

26       **“(ii) The packaging of a drug or biologic described in sub-subparagraph (i) of this sub-**  
27 **paragraph.**

28       **“(U) The packaging in which a product specified in paragraph (a) of this subsection is**  
29 **sold, offered for sale or distributed.**

30       **“(V) Paper and forest products.**

31       **“(4) ‘Component’ means a uniquely identifiable article that is included as a part of a fin-**  
32 **ished product.**

33       **“(5) ‘Contaminant’ means a chemical that is present in a trace amount, that is incidental**  
34 **to manufacturing and serves no intended function in the children’s product or any compo-**  
35 **nent of the children’s product, and that is:**

36       **“(a) An unintended by-product of chemical reactions during the manufacture of the**  
37 **children’s product;**

38       **“(b) A chemical that is unavoidably present in products because of the chemicals’ ubiq-**  
39 **uitous presence in the environment;**

40       **“(c) A trace impurity in feedstock;**

41       **“(d) An incompletely reacted chemical mixture; or**

42       **“(e) A degradation product.**

43       **“(6)(a) ‘Manufacturer’ means:**

44       **“(A) A person that manufactures a children’s product in the form in which the product**  
45 **is sold at retail.**



1 on the basis of the weight of credible, peer-reviewed, scientific evidence, the authority de-  
2 termines that the chemical meets both of the following criteria:

3 “(A) The chemical has been demonstrated by a state or federal agency or an accredited  
4 research university to:

5 “(i) Harm the normal development of a fetus or child or cause other developmental  
6 toxicity;

7 “(ii) Cause cancer, genetic damage or reproductive harm;

8 “(iii) Disrupt the endocrine system such that it causes adverse effects in children;

9 “(iv) Damage the nervous system, immune system or organs or cause other systemic  
10 toxicity; or

11 “(v) Be a very persistent and very bioaccumulative toxic substance; and

12 “(B) There are conditions particular to this state resulting in likely exposure to the  
13 chemical that is expected to cause negative human health impacts, or the chemical has been  
14 found through:

15 “(i) Biomonitoring to be present in human blood, umbilical cord blood, breast milk, urine  
16 or other bodily tissues or fluids;

17 “(ii) Sampling and analysis to be present above 100 parts per million in household dust,  
18 indoor air, drinking water or elsewhere in the home environment; or

19 “(iii) Monitoring to be present above 100 parts per million in fish, wildlife or the natural  
20 environment.

21 “(b) May recommend removing a chemical from the list if the authority determines that  
22 the chemical no longer meets the criteria required for addition to the list as described in  
23 paragraph (a) of this subsection.

24 “(5) A person may petition the authority to consider developing a recommendation to add  
25 or remove a chemical from the list of high priority chemicals by providing the following in-  
26 formation about a chemical to the authority:

27 “(a) The chemical name and the Chemical Abstracts Service Registry Number; and

28 “(b) Information documenting why the chemical meets or fails to meet the criteria re-  
29 quired for addition to the list as described in subsection (4)(a) of this section.

30 “(6) The authority shall present a recommendation to revise the list in a report to the  
31 interim committees of the Legislative Assembly related to environment and natural re-  
32 sources in the manner provided for in ORS 192.245 no later than September 15 of the year  
33 in which the recommendation is proposed. The authority may not adopt a revision to the list  
34 except upon the express consent of the Legislative Assembly.

35 “(7) The authority shall update the list of high priority chemicals on its website within  
36 one year after the date on which a chemical is added to or removed from the list as provided  
37 for in subsection (6) of this section.

38 “(8) This section may not be construed to require the public disclosure by the authority  
39 of any information received from a manufacturer under section 3 or 4 of this 2015 Act that  
40 is a trade secret. If a manufacturer asserts, and can substantiate in a notice submitted un-  
41 der section 3 of this 2015 Act, that the specific identity of a chemical subject to reporting is  
42 a trade secret, the authority shall, in place of the chemical name, publish on the authority’s  
43 website the generic class or category of the chemical, as provided by the manufacturer.

44  
45 “MANUFACTURER DISCLOSURE OF HIGH PRIORITY

CHEMICALS OF CONCERN FOR CHILDREN'S HEALTH

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2  
3       **“SECTION 3. (1) A manufacturer of a children’s product sold or offered for sale in this**  
4 **state that contains a chemical included on the list established and maintained under section**  
5 **2 of this 2015 Act shall provide notice to the Oregon Health Authority as described in this**  
6 **section if the chemical is:**

7           **“(a) Intentionally added in the manufacturing of a children’s product produced by the**  
8 **manufacturer, or a component of the product, is present at a level above the practical**  
9 **quantification limit and serves an intended function in the product; or**

10          **“(b) A contaminant in a children’s product produced by the manufacturer, or a compo-**  
11 **nent of the product, and is present at a concentration above 100 parts per million.**

12          **“(2) Subject to subsection (3) of this section, the authority shall by rule specify the for-**  
13 **mat for the notice required under this section. In adopting rules under this subsection, the**  
14 **authority shall consider, and to the greatest extent practicable develop, a format for the**  
15 **notice that is consistent with the format required by other states with substantially similar**  
16 **reporting requirements.**

17           **“(3)(a) The notice required by this section must contain:**

18           **“(A) The chemical name and Chemical Abstracts Service Registry Number of the chemi-**  
19 **cal contained in the children’s product;**

20           **“(B) A description of the children’s product or product component containing the chem-**  
21 **ical;**

22           **“(C) The amount of the chemical used in each unit of the children’s product reported as**  
23 **a range rather than an exact amount;**

24           **“(D) The name and address of the manufacturer, and the name, address and telephone**  
25 **number of a contact person for the manufacturer;**

26           **“(E) Any other information that the manufacturer deems relevant to the appropriate use**  
27 **of the children’s product; and**

28           **“(F) Any other information determined by the authority by rule to be relevant and es-**  
29 **sential to fulfilling the reporting requirements of this section.**

30          **“(b) The notice required by this section may not be required to contain the disclosure**  
31 **of:**

32           **“(A) Any specific formulation of a chemical or chemicals that is a trade secret; or**

33           **“(B) The name and address of the person responsible for the introduction of the chemical**  
34 **into the children’s product, if that person is a supplier of components, or a person that**  
35 **manufactures components, and is not:**

36           **“(i) The manufacturer required to provide notice under this section; or**

37           **“(ii) Owned or operated by the manufacturer required to provide notice under this sec-**  
38 **tion.**

39          **“(4)(a) A manufacturer required to provide notice under this section may rely on a cer-**  
40 **tificate of compliance, data or other information received from the manufacturer’s suppliers**  
41 **for the purposes of determining reporting obligations under this section.**

42          **“(b) ‘Certificate of compliance,’ for purposes of this subsection and section 4 (2) of this**  
43 **2015 Act, means a certificate provided by a supplier to a manufacturer solely for the purpose**  
44 **of indicating compliance with the provisions of sections 1 to 10 of this 2015 Act.**

45          **“(5)(a) The authority may enter into reciprocal data sharing agreements with other**

1 states in which manufacturers of children’s products are required to disclose information  
2 related to high priority chemicals of concern for children’s health. The authority must use  
3 the GS1 Global Product Classification system to identify and specify product categories sub-  
4 ject to the data sharing agreements. If the authority has entered into a data sharing agree-  
5 ment with another state, and a manufacturer has reported the information required in the  
6 notice under subsections (2) and (3) of this section to that state, the manufacturer may re-  
7 quest that the other state provide the authority with the information in lieu of the  
8 manufacturer’s direct reporting of the information to the authority.

9 “(b) A manufacturer fulfills the notice requirement of subsection (1) of this section when  
10 the authority receives the information from the other state and the authority determines  
11 that the information received satisfies the requirements for the notice under subsections (2)  
12 and (3) of this section.

13 “(6) In lieu of the manufacturer’s providing notice to the authority under subsection (1)  
14 or (5) of this section the authority may require that the notice described in subsections (2)  
15 and (3) of this section be submitted to the Interstate Chemicals Clearinghouse. The authority  
16 by rule shall specify procedures for the provision of such notice by manufacturers to the  
17 Interstate Chemicals Clearinghouse.

18 “(7) A trade association may provide required notices on behalf of its member manufac-  
19 turers under the provisions of this section.

20 “(8) When a manufacturer provides notice to the authority under the provisions of this  
21 section, the manufacturer may submit recommendations to the authority regarding techni-  
22 cal, financial or logistical support deemed necessary for innovation and green chemistry  
23 solutions related to high priority chemicals of concern for children’s health used in children’s  
24 products.

25  
26 “STATEMENTS OF REMOVAL OF CHEMICALS  
27 FROM CHILDREN’S PRODUCTS OR REMOVAL  
28 OF PRODUCTS FROM STATE, EXEMPTIONS  
29

30 “SECTION 4. (1) A manufacturer that is subject to section 3 of this 2015 Act may, at any  
31 time, submit to the Oregon Health Authority a statement that:

32 “(a) The manufacturer has removed from a children’s product sold or offered for sale in  
33 this state the chemical for which the manufacturer is required to submit a notice under  
34 section 3 of this 2015 Act; or

35 “(b) The manufacturer no longer sells, offers for sale or distributes in this state the  
36 children’s product containing the chemical.

37 “(2) A statement submitted under subsection (1)(a) of this section must include relevant  
38 testing results, supplier certificates of compliance or other information received from the  
39 manufacturer’s suppliers demonstrating that the chemical has been removed from the  
40 children’s product.

41 “(3) The authority shall approve or disapprove a statement submitted under subsection  
42 (1) of this section within 30 days after its submittal. Within 30 days after the date that the  
43 authority approves a statement submitted under this section, the authority shall remove  
44 from its website all information related to the children’s product that is the subject of the  
45 statement.



1 Authority may impose a civil penalty on a manufacturer of children’s products for a violation  
2 of any provision of section 3, 4 or 5 of this 2015 Act.

3 “(2) For purposes of assessing civil penalties under this section, a violation consists of a  
4 single course of conduct with regard to an entire children’s product line that is sold or of-  
5 fered for sale in this state.

6 “(3) The authority shall adopt by rule a schedule of civil penalties for violations of  
7 sections 3, 4 and 5 of this 2015 Act. A civil penalty may not exceed \$5,000 for the first vio-  
8 lation. A civil penalty may not exceed \$10,000 for the second and each subsequent violation.

9 “(4) In imposing a penalty under subsection (1) or (5) of this section, the authority shall  
10 consider the following factors:

11 “(a) The past history of the manufacturer in taking all feasible steps or following all  
12 feasible procedures necessary or appropriate to correct any violation.

13 “(b) Any prior violations of statutes, rules, orders or permits pertaining to high priority  
14 chemicals of concern for children’s health used in children’s products.

15 “(c) The gravity and magnitude of the violation.

16 “(d) Whether the violation was a sole event, repeated or continuous.

17 “(e) Whether the violation was a result of an unavoidable accident, negligence or an in-  
18 tentional act.

19 “(f) The manufacturer’s cooperativeness and efforts to correct the violation.

20 “(g) The economic and financial conditions of the manufacturer.

21 “(h) If a manufacturer asserts that a chemical on the list established and maintained  
22 under section 2 of this 2015 Act is present in a children’s product only as a contaminant,  
23 evidence that the manufacturer had in place a reasonable manufacturing control program  
24 for the contaminant and exercised due diligence.

25 “(5)(a) If a manufacturer violates the notice requirement described in section 3 of this  
26 2015 Act, the authority shall inform the manufacturer in writing of the violation and that  
27 the manufacturer may avoid a civil penalty for the violation by providing the notice required  
28 under section 3 of this 2015 Act within 90 days.

29 “(b) If the manufacturer fails to cure the violation within 90 days, the authority may  
30 impose a civil penalty not to exceed \$2,500. For a continuing violation, each 90-day period that  
31 the violation continues after the preceding imposition of a civil penalty is a separate offense  
32 subject to a separate civil penalty not to exceed \$5,000. The authority is not required to  
33 provide the manufacturer with an opportunity to cure the continuing violation before im-  
34 posing a civil penalty for the continuing violation.

35 “(6) If the authority has reason to believe that a children’s product that contains a  
36 chemical on the list established and maintained under section 2 of this 2015 Act is being sold  
37 or offered for sale in this state in violation of section 3, 4 or 5 of this 2015 Act, the authority  
38 may request that the manufacturer provide a statement of compliance on a form provided  
39 by the authority. The manufacturer must submit the statement of compliance within 30 days  
40 after receipt of the request. To prove compliance with sections 3, 4 and 5 of this 2015 Act,  
41 the manufacturer must:

42 “(a) Show that the children’s product does not contain the chemical;

43 “(b) Show that the manufacturer has previously provided the authority with notice as  
44 required by section 3 of this 2015 Act;

45 “(c) Provide the authority with notice as required by section 3 of this 2015 Act; or





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**“CAPTIONS**

**“SECTION 12. The unit captions used in this 2015 Act are provided only for the convenience of the reader and do not become part of the statutory law of this state or express any legislative intent in the enactment of this 2015 Act.**

**“SUNSET**

**“SECTION 13. Sections 1 to 10 of this 2015 Act are repealed on January 2, 2020.”.**

/s/ Alan Olsen  
Senator

/s/ Chuck Thomsen  
Senator

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