

1 H.536

2 Introduced by Representatives Stone of Burlington, Arsenault of Williston,  
3 and Graning of Jericho

4 Referred to Committee on

5 Date:

6 Subject: Health; adulterated food; baby food products; testing; labeling

7 Statement of purpose of bill as introduced: This bill proposes to prohibit the  
8 sale and distribution of a baby food product in the State that contains a toxic  
9 heavy metal. It further proposes labeling requirements and regular laboratory  
10 testing of baby food products for toxic heavy metals.

11 An act relating to toxic heavy metals in baby food products

12 It is hereby enacted by the General Assembly of the State of Vermont:

13 Sec. 1. 18 V.S.A. chapter 82 is amended to read:

14 CHAPTER 82. LABELING OF FOOD, DRUGS, COSMETICS, AND  
15 HAZARDOUS SUBSTANCES

16 Subchapter 1. ~~Labeling and Marketing for Sale~~ General Provisions

17 \* \* \*

18 Subchapter 3. Testing and Labeling of Certain Products

19 § 4091. BABY FOOD PRODUCTS

20 (a) As used in this section:

1           (1) “Baby food product” means any food manufactured, packaged, and  
2           labeled in a jar, pouch, tub, or box sold specifically for babies and children  
3           younger than two years of age. “Baby food product” does not include infant  
4           formula.

5           (2) “Commissioner” means the Commissioner of Health.

6           (3) “Final baby food product” means the finished baby food product and  
7           not the constituent ingredients.

8           (4) “Infant formula” means a commercially available milk-based or soy-  
9           based powder, concentrated liquid, or ready-to-feed substitute for human breast  
10          milk that is intended for infant consumption.

11          (5) “Production aggregate” means a quantity of product that is intended  
12          to have uniform composition, character, and quality and is produced according  
13          to a master manufacturing order.

14          (6) “Proficient laboratory” means a laboratory that:

15               (A) is accredited under the standards of the International  
16               Organization for Standardization or the International Electrotechnical  
17               Commission pursuant to standard ISO/IEC 17025:2017;

18               (B) uses an analytical method as sensitive as the analytical method  
19               described in the U.S. FDA’s Elemental Analysis Manual for Food Related  
20               Products; and

1           (C) demonstrates proficiency in quantifying each toxic element to at  
2           least six micrograms of the toxic element to kilogram of food through an  
3           independent proficiency test by achieving a z-score that is less than or equal to  
4           plus or minus two.

5           (7) “QR code” means a two-dimension matrix barcode consisting of  
6           blocks arranged in a grid that can be read by an imaging device.

7           (8) “Representative sample” means a sample that consists of a number  
8           of units that are drawn based on rational criteria, including random sampling,  
9           and intended to ensure that the sample accurately portrays the material being  
10          sampled.

11          (9) “Toxic heavy metal” means arsenic, cadmium, lead, and mercury.

12          (10) “URL” means a uniform resource locator.

13          (11) “U.S. FDA” means the U.S. Food and Drug Administration.

14          (b) A person shall not sell, distribute, or offer for sale any baby food  
15          product that contains a toxic heavy metal that exceeds the limits established by  
16          the U.S. FDA. The provisions of this subsection shall not restrict the  
17          continued sale of inventory in stock before January 1, 2026.

18          (c) A manufacturer of a baby food shall test a representative sample of each  
19          production aggregate of the manufacturer’s final baby food product for toxic  
20          heavy metals. Testing of a baby food product shall be conducted by a  
21          proficient laboratory at least once a month. A manufacturer of baby food may

1 test the final baby food product before packaging individual units for sale or  
2 distribution. Upon request of the Commissioner, a manufacturer shall provide  
3 the results of the test conducted pursuant to this subsection.

4 (d) A manufacturer of baby food shall make publicly available on its  
5 website for the duration of the product shelf life of a final baby food product,  
6 plus one month, for each baby food product sold, manufactured, delivered,  
7 held, or offered for sale in the State:

8 (1) the name and level of each toxic heavy metal in the final baby food  
9 product as determined by the testing conducted pursuant to subsection (c) of  
10 this section;

11 (2) sufficient information, including the product name, universal product  
12 code, or lot or batch number, to enable consumers to identify the final baby  
13 food product; and

14 (3) a link to the U.S. FDA's website that provides the most recent U.S.  
15 FDA guidance and information about the health effects of toxic heavy metals  
16 on children.

17 (e) If a baby food product is tested for a toxic heavy metal subject to an  
18 action level, regulatory limit, or tolerance established by the U.S. FDA under  
19 21 C.F.R. § 109, the manufacturer shall display on the baby food product:

1           (1) a label stating in a clear, legible, conspicuous manner that more  
2           information about toxic element testing on the product is available by scanning  
3           the QR code; and

4           (2) a QR code or other machine-readable code that directs the  
5           consumers to the manufacturer's website or the baby food product information  
6           page providing:

7                   (A) the test results for the toxic heavy metal; and

8                   (B) a URL to the web page on the U.S. FDA's website that includes  
9           the most recent guidance and information about the health effects of toxic  
10          heavy metals in children.

11          (f) If a consumer reasonably believes, based on the information provided  
12          on the baby food product, that the baby food product is being sold in the State  
13          in violation of this section, the consumer shall report the baby food product to  
14          the Commissioner.

15          (g) A violation of this section shall be deemed a violation of the Consumer  
16          Protection Act, 9 V.S.A. chapter 63. The Attorney General has the same  
17          authority to make rules, conduct civil investigations, enter into assurances of  
18          discontinuance, and bring civil actions, and private parties have the same rights  
19          and remedies, as provided under 9 V.S.A. chapter 63, subchapter 1.

20          Sec. 2. EFFECTIVE DATE

21          This act shall take effect on July 1, 2026.