

112TH CONGRESS
1ST SESSION

S. 136

To establish requirements with respect to bisphenol A.

IN THE SENATE OF THE UNITED STATES

JANUARY 25 (legislative day, JANUARY 5), 2011

Mr. REID (for Mrs. FEINSTEIN (for herself, Mr. SCHUMER, Mr. KERRY, Mr. SANDERS, and Mr. FRANKEN)) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

A BILL

To establish requirements with respect to bisphenol A.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

3 **SECTION 1. SHORT TITLE.**

4 This Act may be cited as the “Ban Poisonous Addi-
5 tives Act of 2011”.

6 **SEC. 2. REQUIREMENTS WITH RESPECT TO BISPHENOL A.**

7 (a) BAN ON USE OF BISPHENOL A IN FOOD AND
8 BEVERAGE CONTAINERS FOR CHILDREN.—

9 (1) BABY FOOD; UNFILLED BABY BOTTLES AND
10 CUPS.—Section 402 of the Federal Food, Drug, and

1 Cosmetic Act (21 U.S.C. 342) is amended by adding
2 at the end the following:

3 “(j)(1) If it is a food intended for children 3 years
4 of age or younger, the container of which (including the
5 lining of such container) is composed, in whole or in part,
6 of bisphenol A.

7 “(2) If it is a baby bottle or cup that is composed,
8 in whole or in part, of bisphenol A.”.

9 (2) DEFINITION.—Section 201 of the Federal
10 Food, Drug, and Cosmetic Act (21 U.S.C. 321) is
11 amended by adding at the end the following:

12 “(rr) BABY BOTTLE OR CUP.—For purposes of sec-
13 tion 402(j), the term ‘baby bottle or cup’ means a bottle
14 or cup that—

15 “(1) is intended to aid in the feeding or pro-
16 viding of drink to children 3 years of age or young-
17 er; and

18 “(2) does not contain a food when such bottle
19 or cup is sold or distributed at retail.”.

20 (3) EFFECTIVE DATES.—

21 (A) BABY FOOD.—Section 402(j)(1) of the
22 Federal Food, Drug, and Cosmetic Act, as
23 added by paragraph (1), shall take effect 1 year
24 after the date of enactment of this Act.

1 (B) UNFILLED BABY BOTTLES AND
2 CUPS.—Section 402(j)(2) of the Federal Food,
3 Drug, and Cosmetic Act, as added by para-
4 graph (1), shall take effect 180 days after the
5 date of enactment of this Act.

6 (b) BAN ON USE OF BISPHENOL A IN INFANT FOR-
7 MULA CONTAINERS.—

8 (1) IN GENERAL.—Section 412(a) of the Fed-
9 eral Food, Drug, and Cosmetic Act (21 U.S.C.
10 350a(a)) is amended—

11 (A) in paragraph (2), by striking “, or”
12 and inserting “,”;

13 (B) in paragraph (3), by striking the pe-
14 riod at the end and inserting “, or”; and

15 (C) by adding at the end the following:

16 “(4) the container of such infant formula (in-
17 cluding the lining of such container and, in the case
18 of infant formula powder, excluding packaging on
19 the outside of the container that does not come into
20 contact with the infant formula powder) is com-
21 posed, in whole or in part, of bisphenol A.”.

22 (2) EFFECTIVE DATE.—The amendments made
23 by paragraph (1) shall take effect 18 months after
24 the date of enactment of this Act.

1 (c) REGULATION OF OTHER CONTAINERS COMPOSED
2 OF BISPHENOL A.—

3 (1) SAFETY ASSESSMENT OF PRODUCTS COM-
4 POSED OF BPA.—Not later than December 1, 2012,
5 the Secretary of Health and Human Services (re-
6 ferred to in this Act as the “Secretary”) shall issue
7 a revised safety assessment for food containers com-
8 posed, in whole or in part, of bisphenol A, taking
9 into consideration different types of such food con-
10 tainers and the use of such food containers with re-
11 spect to different foods, as appropriate.

12 (2) SAFETY STANDARD.—Through the safety
13 assessment described in paragraph (1), and taking
14 into consideration the requirements of section 409 of
15 the Federal Food, Drug, and Cosmetic Act (21
16 U.S.C. 348) and section 170.3(i) of title 21, Code of
17 Federal Regulations, the Secretary shall determine
18 whether there is a reasonable certainty that no harm
19 will result from aggregate exposure to bisphenol A
20 through food containers or other items composed, in
21 whole or in part, of bisphenol A, taking into consid-
22 eration potential adverse effects from low dose expo-
23 sure, and the effects of exposure on vulnerable popu-
24 lations, including pregnant women, infants, children,

1 the elderly, and populations with high exposure to
2 bisphenol A.

3 (3) APPLICATION OF SAFETY STANDARD TO AL-
4 TERNATIVES.—The Secretary shall use the safety
5 standard described under paragraph (2) to evaluate
6 the proposed uses of alternatives to bisphenol A.

7 (d) SAVINGS PROVISION.—Nothing in this section
8 shall affect the right of a State, political subdivision of
9 a State, or Indian Tribe to adopt or enforce any regula-
10 tion, requirement, liability, or standard of performance
11 that is more stringent than a regulation, requirement, li-
12 ability, or standard of performance under this section or
13 that—

14 (1) applies to a product category not described
15 in this section; or

16 (2) requires the provision of a warning of risk,
17 illness, or injury associated with the use of food con-
18 tainers composed, in whole or in part, of bisphenol
19 A.

20 (e) DEFINITION.—For purposes of this section, the
21 term “container” includes the lining of a container.

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