

117TH CONGRESS
2D SESSION

S. _____

To provide for the expedited and duty-free importation of infant formula that may be lawfully marketed in the European Union, Canada, Japan, or the United Kingdom, and for other purposes.

IN THE SENATE OF THE UNITED STATES

Mrs. GILLIBRAND introduced the following bill; which was read twice and referred to the Committee on _____

A BILL

To provide for the expedited and duty-free importation of infant formula that may be lawfully marketed in the European Union, Canada, Japan, or the United Kingdom, and for other purposes.

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 “Emergency Infant For-
5 mula Act”.

6 **SEC. 2. EXPEDITED IMPORTATION OF INFANT FORMULA.**

7 (a) AUTHORIZATION FOR IMPORTATION AND SALE.—

1 (1) DECLARATION OF SHORTAGE.—The Presi-
2 dent may declare, in consultation with the Commis-
3 sioner of Food and Drugs and by issuing an Execu-
4 tive order, that a shortage exists in the United
5 States of infant formula during any period specified
6 in that Executive order.

7 (2) AUTHORIZATION FOR IMPORTATION AND
8 SALE.—

9 (A) IN GENERAL.—The President may au-
10 thorize, in consultation with the Commissioner
11 of Food and Drugs, the importation, distribu-
12 tion, and sale of any covered infant formula,
13 notwithstanding the provisions of the Federal
14 Food, Drug, and Cosmetic Act (21 U.S.C. 201
15 et seq.), if the applicable brand, manufacturer,
16 or manufacturing plant, or the specific infant
17 formula product, is included in the Executive
18 order issued pursuant to paragraph (1).

19 (B) ADDITIONAL REQUIREMENTS.—The
20 Executive order issued pursuant to paragraph
21 (1) may specify, with respect to any covered in-
22 fant formula, specific requirements with respect
23 to labeling or usage guidance to be eligible for
24 importation, distribution, and sale pursuant
25 subparagraph (A).

1 (3) LABELING REQUIREMENTS.—The Commis-
2 sioner of Food and Drugs shall require any retailer
3 of covered infant formula imported pursuant to
4 paragraph (2), including an online retailer, to in-
5 clude in an appropriate and conspicuous place next
6 to the product or description of the product, as ap-
7 plicable, a label that—

8 (A) indicates that the covered infant for-
9 mula may not meet the standards under section
10 412 of the Federal Food, Drug, and Cosmetic
11 Act (21 U.S.C. 350a) for importation, distribu-
12 tion, or sale and is authorized for sale only sub-
13 ject to the provisions of this section; and

14 (B) may additionally indicate the foreign
15 country or countries in which the covered infant
16 formula may be lawfully marketed.

17 (4) TERMINATION OF SHORTAGE.—The Presi-
18 dent may, upon determining that a shortage no
19 longer exists in the United States of infant formula,
20 terminate a declaration issued under paragraph (1).

21 (b) DUTY-FREE TREATMENT.—Notwithstanding any
22 other provision of law, the President may, during any pe-
23 riod in which an infant formula shortage is declared pur-
24 suant to subsection (a)(1), reduce or suspend any duties
25 imposed—

1 (1) with respect to the importation of covered
2 infant formula; or

3 (2) with respect to any other article used in the
4 production of infant formula that the importer cer-
5 tifies is being imported for such production.

6 (c) **PRIORITY HANDLING OF ENTRIES.**—During any
7 period in which an infant formula shortage is declared
8 pursuant to subsection (a)(1), the Commissioner of U.S.
9 Customs and Border Patrol shall give the highest priority
10 and take such steps as may be necessary to expedite the
11 processing of all entries of covered infant formula and ar-
12 ticles used in the production of infant formula (as de-
13 scribed in subsection (b)(2)).

14 (d) **USE OF DEFENSE PRODUCTION ACT AUTHORI-**
15 **TIES.**—During any period in which an infant formula
16 shortage is declared pursuant to subsection (a)(1)—

17 (1) the President may use authorities provided
18 by the Defense Production Act of 1950 (50 U.S.C.
19 4501 et seq.) with respect to the production of in-
20 fant formula; and

21 (2) infant formula shall be deemed to meet the
22 criteria specified in section 101(b) of such Act (50
23 U.S.C. 4511(b)).

24 (e) **DEFINITIONS.**—In this section:

1 (1) COVERED INFANT FORMULA.—The term
2 “covered infant formula” means any infant formula
3 that is lawfully marketed, as of the date of the en-
4 actment of this Act, in the European Union, Can-
5 ada, Japan, or the United Kingdom.

6 (2) INFANT FORMULA.—The term “infant for-
7 mula” has the meaning given that term in section
8 201(z) of the Federal Food, Drug, and Cosmetic Act
9 (21 U.S.C. 321(z)).