H. R. 3411

To amend the Public Health Service Act to establish a program of research regarding the risks posed by the presence of dioxin, synthetic fibers, and other additives in feminine hygiene products, and to establish a program for the collection and analysis of data on toxic shock syndrome.

IN THE HOUSE OF REPRESENTATIVES

July 22, 2005

Mrs. Maloney (for herself, Mr. Gutierrez, Mr. Waxman, Mr. McNulty, Mr. Sanders, Mrs. Christensen, Ms. Jackson-Lee of Texas, Mr. Abercrombie, and Ms. Woolsey) introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

To amend the Public Health Service Act to establish a program of research regarding the risks posed by the presence of dioxin, synthetic fibers, and other additives in feminine hygiene products, and to establish a program for the collection and analysis of data on toxic shock syndrome.

- 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 "Robin Danielson Act".

TITLE I—RESEARCH REGARDING

- 2 RISKS POSED BY DIOXIN, SYN-
- 3 THETIC FIBERS, AND OTHER
- 4 ADDITIVES IN FEMININE HY-
- 5 **GIENE PRODUCTS**
- 6 SEC. 101. FINDINGS.
- 7 The Congress finds as follows:
- 8 (1) Tampons are used by approximately
- 9 73,000,000 women in the United States today, and
- the average woman may use as many as 16,800
- tampons in her lifetime. A woman on estrogen re-
- placement therapy may use as many as 24,360 tam-
- pons in her lifetime.
- 14 (2) The Environmental Protection Agency and
- the International Agency for Research on Cancer, an
- arm of the World Health Organization, have con-
- 17 cluded that dioxins are a probable human carcinogen
- 18 (cancer causing agent).
- 19 (3) Dioxin is a byproduct of chlorine-bleaching
- processes used in the manufacture of paper prod-
- 21 ucts, including tampons, sanitary pads, panty liners,
- and diapers.
- 23 (4) While bleaching processes that do not
- produce dioxin in any amount are available, most
- 25 pulp and paper manufacturers, which produce the

- raw materials used in tampons, currently use either elemental-chlorine or chlorine-dioxide bleaching processes. Both of these bleaching processes use chlorine and therefore produce dioxin.
 - (5) The effects of dioxin from various sources are cumulative and can be measured 20 to 30 years after exposure. Women may be exposed to dioxin in tampons and other menstrual products for as long as 60 years over the course of their reproductive lives.
 - (6) Internal documents of the Food and Drug Administration suggest the agency has not adequately investigated the danger of dioxin in tampons, according to a 1992 staff report of a subcommittee of the Committee on Government Operations of the House of Representatives.
 - (7) The Food and Drug Administration has historically relied on data provided by manufacturers of feminine hygiene products in determining product safety.
 - (8) Although the Food and Drug Administration currently requires tampon manufacturers to monitor dioxin levels in their finished products, the information is not readily available to the public.

- (9) Recent studies have produced conflicting information about the link between dioxin exposure and increased risks for endometriosis.
 - (10) The Environmental Protection Agency has concluded that people with high levels of exposure to dioxins may be at risk for other noncancer effects that could suppress the immune system, increase the risk of pelvic inflammatory disease, reduce fertility, and interfere with fetal and childhood development.
 - (11) An independent study in 1991 found that tampons commonly included one or more of the following additives: Chlorine compounds, absorbency enhancers (such as surfactants like polysorbate-20), natural and synthetic fibers (such as cotton, rayon, polyester, and polyacrylate), deodorant, and fragrance.
 - (12) Toxic Shock Syndrome (TSS) has been linked to tampon use and the absorbency of the tampon. TSS is a rare bacterial illness that occurs mostly in menstruating women. During 1979 and 1980, the syndrome was responsible for at least 55 deaths and 1,066 nonfatal cases.
 - (13) In response to a 1988 lawsuit, the Food and Drug Administration has required tampons to be labeled with reference to an absorbency standard

1	(e.g., super tampons must absorb between 9 and 12
2	grams of liquid).
3	(14) Independent research has shown that syn-
4	thetic fiber additives in tampons amplify toxin pro-
5	duction, which is associated with toxic shock syn-
6	drome.
7	SEC. 102. NATIONAL INSTITUTES OF HEALTH; RESEARCH
8	ON DIOXIN PURSUANT TO OFFICE OF RE-
9	SEARCH ON WOMEN'S HEALTH.
10	Part F of title IV of the Public Health Service Act
11	(42 U.S.C. 287d et seq.) is amended by adding at the end
12	the following section:
13	"SEC. 486C. CERTAIN PROJECTS REGARDING WOMEN'S
14	HEALTH.
	HEALTH. "(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.—
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15 16	"(a) Dioxin in Feminine Hygiene Products.—
15 16 17	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in
15 16 17 18	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in collaboration with the Director of the Office, shall
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115 116 117 118 119 220	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in collaboration with the Director of the Office, shall provide for the conduct or support of research to determine the extent to which the presence of dioxin,
14 15 16 17 18 19 20 21	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in collaboration with the Director of the Office, shall provide for the conduct or support of research to determine the extent to which the presence of dioxin, synthetic fibers, and other additives in tampons and
115 116 117 118 119 220 221	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in collaboration with the Director of the Office, shall provide for the conduct or support of research to determine the extent to which the presence of dioxin, synthetic fibers, and other additives in tampons and other feminine hygiene products—
115 116 117 118 119 220 221 222	"(a) DIOXIN IN FEMININE HYGIENE PRODUCTS.— "(1) IN GENERAL.—The Director of NIH, in collaboration with the Director of the Office, shall provide for the conduct or support of research to determine the extent to which the presence of dioxin, synthetic fibers, and other additives in tampons and other feminine hygiene products— "(A) poses any risks to the health of

- system deficiencies, pelvic inflammatory disease,
 and toxic shock syndrome; and
- "(B) poses any risks to the health of children of women who used such products during or before the pregnancies involved, including risks relating to fetal and childhood development.
- 9 "(2) REQUIREMENT REGARDING DATA FROM
 9 MANUFACTURERS.—Research under paragraph (1)
 10 shall include research to confirm the data on tam11 pons and other feminine hygiene products submitted
 12 to the Commissioner of Food and Drugs by manu13 facturers of such products.
- "(3) DEFINITION.—For purposes of paragraph (1), the term 'feminine hygiene products' means tampons, pads, liners, and similar products used by women with respect to menstruation or other genital-tract secretions.
- "(b) Reports.—Reports on the results of research under subsection (a) shall be periodically submitted to the Congress, the Commissioner of Food and Drugs, the Administrator of the Environmental Protection Agency, and the Chairman of the Consumer Product Safety Commission. Such reports shall be made available to the public through the data system and clearinghouse program es-

1	tablished under section 486A, or through other appro-
2	priate means.".
3	TITLE II—COLLECTION AND
4	ANALYSIS OF DATA ON TOXIC
5	SHOCK SYNDROME
6	SEC. 201. FINDINGS.
7	The Congress finds as follows:
8	(1) Of the cases of toxic shock syndrome in the
9	United States, approximately 50 percent are related
10	to tampon use and approximately 50 percent occur
11	in nonmenstruating women and in men and children.
12	(2) The Centers for Disease Control and Pre-
13	vention (CDC) believes that women are at increased
14	risk for developing toxic shock syndrome due to a
15	false sense of security that there is no longer any
16	risk for developing the disease.
17	(3) The CDC has estimated that each year such
18	syndrome strikes more than 1,300 individuals.
19	Among women in the age group 12 through 44 who
20	use tampons or barrier contraceptives, between one
21	and two of every 100,000 will develop the syndrome.
22	(4) Epidemiological data on cases of toxic shock

syndrome are not systematically collected in the

United States, and information on cases seldom

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- travels beyond the victim's circle of family and friends.
- 5 The CDC and the States should cooperate
 to collect and analyze such data. Increasing the
 amount of information on toxic shock syndrome will
 lead to increased awareness about the disease in the
 medical community, and may also lead to an increased understanding of the causes of the syn-
- 10 SEC. 202. CENTERS FOR DISEASE CONTROL AND PREVEN-
- 11 TION; ESTABLISHMENT OF PROGRAM FOR
- 12 COLLECTION AND ANALYSIS OF DATA ON
- 13 TOXIC SHOCK SYNDROME.
- Part B of title III of the Public Health Service Act
- 15 (42 U.S.C. 243 et seq.) is amended by inserting after sec-
- 16 tion 317S the following section:

9

drome.

- 17 "SEC. 317T. COLLECTION AND ANALYSIS OF DATA ON TOXIC
- 18 SHOCK SYNDROME.
- 19 "(a) IN GENERAL.—The Secretary, acting through
- 20 the Director of the Centers for Disease Control and Pre-
- 21 vention, shall carry out a program to collect, analyze, and
- 22 make available data on toxic shock syndrome, including
- 23 data on the causes of such syndrome.
- 24 "(b) National Incidence and Prevalence.—In
- 25 carrying out the program under subsection (a), the Sec-

- 1 retary shall to the extent practicable determine the na-
- 2 tional incidence and prevalence of toxic shock syndrome.
- 3 "(c) Cooperation With States.—The Secretary
- 4 may carry out the program under subsection (a) directly
- 5 and through grants to States and local health depart-
- 6 ments.
- 7 "(d) AUTHORIZATION OF APPROPRIATIONS.—For the
- 8 purpose of carrying out this section, there are authorized
- 9 to be appropriated such sums as may be necessary for
- 10 each of the fiscal years 2006 through 2010.".

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