

AMENDMENT NO. _____ Calendar No. _____

Purpose: To require studies by the Secretary of Agriculture on the effects of food products from cloned animals entering the food supply.

IN THE SENATE OF THE UNITED STATES—110th Cong., 1st Sess.

H. R. 2419

To provide for the continuation of agricultural programs through fiscal year 2012, and for other purposes.

Referred to the Committee on _____ and ordered to be printed

Ordered to lie on the table and to be printed

AMENDMENT intended to be proposed by Ms. MIKULSKI (for herself and Mr. SPECTER)

Viz:

1 On page 1045, after line 2, insert the following:

2 **SEC. 7505. STUDIES AND REPORTS BY THE DEPARTMENT**
 3 **OF AGRICULTURE AND THE NATIONAL ACAD-**
 4 **EMY OF SCIENCES ON FOOD PRODUCTS**
 5 **FROM CLONED ANIMALS.**

6 (a) STUDY BY THE DEPARTMENT OF AGRICULTURE.—
 7

8 (1) IN GENERAL.—The Secretary of Agri-
 9 culture, in coordination with the Economic Research

1 Service, and after consultation with the Secretary of
2 Health and Human Services, shall conduct a study
3 and report to Congress on the state of domestic and
4 international markets for products from cloned ani-
5 mals, including consumer acceptance. Such report
6 shall be submitted to Congress no later than 180
7 days after the date of enactment of this Act.

8 (2) CONTENT.—The study and report under
9 paragraph (1) shall include a description of how
10 countries regulate the importation of food and agri-
11 cultural products (including dairy products), the
12 basis for such regulations, and potential obstacles to
13 trade.

14 (b) STUDY WITH THE NATIONAL ACADEMY OF
15 SCIENCES.—

16 (1) IN GENERAL.—The Secretary of Agriculture
17 shall contract with the National Academy of
18 Sciences to conduct a study and report to Congress
19 regarding the safety of food products derived from
20 cloned animals and the health effects and costs at-
21 tributable to milk from cloned animals in the food
22 supply. Such report shall be submitted to Congress
23 no later than 1 year after the date of enactment of
24 this Act.

1 (2) CONTENT.—The study and report under
2 paragraph (1) shall include—

3 (A) a review and an assessment of whether
4 the studies (including peer review studies),
5 data, and analysis used in the draft risk assess-
6 ment issued by the Food and Drug Administra-
7 tion entitled *Animal Cloning: A Draft Risk As-*
8 *essment* (issued on December 28, 2006) sup-
9 ported the conclusions drawn by such draft risk
10 assessment and—

11 (i) whether there were a sufficient
12 number of studies to support such conclu-
13 sions; and

14 (ii) whether additional pertinent stud-
15 ies and data exist which were not consid-
16 ered in the draft risk assessment and how
17 this additional information affects the con-
18 clusions drawn in such draft risk assess-
19 ment; and

20 (B) an evaluation and measurement of the
21 potential public health effects and associated
22 health care costs, including any consumer be-
23 havior changes and negative impacts on nutri-
24 tion, health, and chronic diseases that may re-
25 sult from any decrease in dairy consumption,

1 attributable to the commercialization of milk
2 from cloned animals and their progeny.

3 (c) **RULE OF CONSTRUCTION.**—Nothing in this sec-
4 tion shall be construed to impede ongoing scientific re-
5 search in artificial reproductive health technologies.

6 (d) **TIMEFRAME OF FINAL RISK ASSESSMENT.**—Not-
7 withstanding any other provision of law, the Secretary of
8 Health and Human Services (acting through the Commis-
9 sioner of Food and Drugs) shall not issue the final risk
10 assessment on the safety of cloned animals and food prod-
11 ucts derived from cloned animals until the date that the
12 Secretary of Agriculture completes the studies required
13 under this section.

14 (e) **CONTINUANCE OF MORATORIUM.**—Any voluntary
15 moratorium on introducing food from cloned animals or
16 their progeny into the food supply shall remain in effect
17 at least until the date that the Secretary of Health and
18 Human Services (acting through the Commissioner of
19 Food and Drugs) issues the final risk assessment de-
20 scribed in subsection (d).