

110TH CONGRESS
1ST SESSION

H. R. 2718

To ensure that the goals of the Dietary Supplement Health and Education Act of 1994 are met by authorizing appropriations to fully enforce and implement such Act and the amendments made by such Act, and for other purposes.

IN THE HOUSE OF REPRESENTATIVES

JUNE 14, 2007

Mr. BURTON of Indiana introduced the following bill; which was referred to the Committee on Energy and Commerce

A BILL

To ensure that the goals of the Dietary Supplement Health and Education Act of 1994 are met by authorizing appropriations to fully enforce and implement such Act and the amendments made by such Act, 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 “DSHEA Full Imple-
5 mentation and Enforcement Act of 2007”.

6 **SEC. 2. FINDINGS.**

7 Congress finds the following:

1 (1) Over 158,000,000 Americans regularly con-
2 sume dietary supplements to maintain and improve
3 their health.

4 (2) Consumer expenditures on dietary supple-
5 ments reached a reported \$17,100,000,000 in 2000,
6 double the amount spent in 1994.

7 (3) According to a recent report issued by the
8 Food and Drug Administration (in this Act referred
9 to as the “FDA”) the use of dietary supplements is
10 likely to grow due to factors such as the aging of the
11 baby boom generation, increased interest in self-suf-
12 ficiency, and advances in science that are uncovering
13 new relationships between diet and disease.

14 (4) In 1994, the Dietary Supplement Health
15 and Education Act of 1994 (Public Law 103–417)
16 (in this Act referred to as “DSHEA”) was enacted.
17 This Act balanced continued consumer access to vi-
18 tamins, minerals, and other dietary supplements, in-
19 creased scientific research on the benefits and risks
20 of dietary supplements, public education on dietary
21 supplements, and needed consumer protections.

22 (5) DSHEA requires that claims made on die-
23 tary supplement labels, packaging, and accom-
24 panying material be truthful, non-misleading, and
25 substantiated. Manufacturers are prohibited from

1 making claims that products are intended to diag-
2 nose, treat, mitigate, cure, or prevent a disease.

3 (6) DSHEA provides for good manufacturing
4 practice standards setting requirements for potency,
5 purity, sanitary conditions, and recordkeeping for di-
6 etary supplements.

7 (7) DSHEA requires that manufacturers sub-
8 mit adequate information as to the safety of any
9 new ingredients contained in dietary supplements be-
10 fore those products can be sold.

11 (8) The FDA has updated and expanded its
12 system for the reporting, collection, and analysis of
13 dietary supplement adverse events reports.

14 (9) DSHEA provides the FDA with a number
15 of authorities to remove unsafe dietary supplements
16 from the marketplace.

17 (10) DSHEA created the Office of Dietary
18 Supplements within the National Institutes of
19 Health to expand research and consumer informa-
20 tion about the health effects of dietary supplements.

21 (11) The FDA has not adequately used its au-
22 thority to enforce DSHEA.

23 (12) The FDA needs adequate resources to ap-
24 propriately implement and enforce DSHEA. Con-
25 gress has appropriated additional funds over the last

1 several years beyond those requested in the Presi-
 2 dent's budget to implement and enforce DSHEA,
 3 reaching \$9,700,000 in fiscal year 2003.

4 (13) However, according to the FDA, full im-
 5 plementation of DSHEA would require substantial
 6 additional resources. The FDA asserts that between
 7 \$24,000,000 and \$65,000,000 per year will be need-
 8 ed to fully implement DSHEA.

9 **SEC. 3. AUTHORIZATION AND APPROPRIATION OF RE-**
 10 **SOURCES.**

11 (a) AUTHORIZATION OF APPROPRIATIONS.—There
 12 are authorized to be appropriated to carry out the Dietary
 13 Supplement Health and Education Act of 1994 (Public
 14 Law 103–417), the amendments made by such Act, and
 15 all applicable regulatory requirements for dietary supple-
 16 ments under the Federal Food, Drug, and Cosmetic Act
 17 (21 U.S.C. 301 et seq.)—

- 18 (1) \$20,000,000 for fiscal year 2008;
- 19 (2) \$30,000,000 for fiscal year 2009;
- 20 (3) \$40,000,000 for fiscal year 2010;
- 21 (4) \$50,000,000 for fiscal year 2011; and
- 22 (5) \$65,000,000 for fiscal year 2012.

23 (b) APPROPRIATION OF FUNDS FOR FISCAL YEAR
 24 2008.—There are appropriated, out of any money in the
 25 Treasury not otherwise appropriated, to carry out the Die-

1 tary Supplement Health and Education Act of 1994 (Pub-
2 lic Law 103–417), the amendments made by such Act, and
3 all applicable regulatory requirements for dietary supple-
4 ments under the Federal Food, Drug, and Cosmetic Act
5 (21 U.S.C. 301 et seq.), \$20,000,000 for fiscal year 2008.

6 (c) OFFICE OF DIETARY SUPPLEMENTS.—There are
7 authorized to be appropriated and there are appropriated,
8 out of any money in the Treasury not otherwise appro-
9 priated, for expanded research and development of con-
10 sumer information on dietary supplements by the Office
11 of Dietary Supplements at the National Institutes of
12 Health—

13 (1) \$30,000,000 for fiscal year 2008; and

14 (2) such sums as may be necessary for each of
15 the fiscal years 2009 through 2012.

16 (d) USE OF FUNDS.—The Food and Drug Adminis-
17 tration shall fully and appropriately use the funds appro-
18 priated in subsections (b) and (c) and pursuant to sub-
19 section (a) to regulate dietary supplements.

20 **SEC. 4. ANNUAL ACCOUNTABILITY REPORT ON THE REGU-**
21 **LATION OF DIETARY SUPPLEMENTS.**

22 (a) IN GENERAL.—Not later than January 31, 2008,
23 and annually thereafter, the Secretary of Health and
24 Human Services shall submit a report to Congress on the
25 implementation and enforcement of the Dietary Supple-

1 ment Health and Education Act of 1994 (Public Law
2 103–417).

3 (b) CONTENTS.—The report under subsection (a)
4 shall include the following:

5 (1) The total funding and number of full-time
6 equivalent personnel in the Food and Drug Adminis-
7 tration dedicated to dietary supplement regulation
8 over the prior fiscal year.

9 (2) The total funding and number of full-time
10 equivalent personnel in the Food and Drug Adminis-
11 tration dedicated to administering adverse event re-
12 porting systems as they relate to dietary supplement
13 regulation over the prior fiscal year.

14 (3) The total funding and number of full-time
15 equivalent personnel in the Food and Drug Adminis-
16 tration dedicated to enforcement of dietary supple-
17 ment labeling and claims requirements over the prior
18 fiscal year and an explanation of their activities.

19 (4) The total funding and number of full-time
20 equivalent personnel in the Food and Drug Adminis-
21 tration dedicated to good manufacturing practices
22 inspections of dietary supplement manufacturers
23 over the prior fiscal year and an explanation of their
24 activities.

1 (5) The number of good manufacturing prac-
2 tices inspections of dietary supplement manufactur-
3 ers by the Food and Drug Administration over the
4 prior fiscal year and a summary of the results.

5 (6) The number of new ingredient reviews and
6 safety reviews related to dietary supplements and
7 the results of those reviews.

8 (7) An explanation of all enforcement actions
9 taken by the Food and Drug Administration and the
10 Department of Health and Human Services related
11 to dietary supplements over the prior fiscal year, in-
12 cluding the number and type of actions.

13 (8) The number of dietary supplement claims
14 for which the Food and Drug Administration re-
15 quested substantiation from the manufacturer over
16 the prior fiscal year, and the agency's response.

17 (9) The number of dietary supplement claims
18 determined to be false, misleading, or nonsubstan-
19 tiated by the Food and Drug Administration over
20 the prior fiscal year.

21 (10) The research and consumer education ac-
22 tivities supported by the Office of Dietary Supple-
23 ments of the National Institutes of Health.

1 (11) Any recommendations for administrative
2 or legislative actions regarding the regulation of die-
3 tary supplements.

4 (12) Any other information regarding the regu-
5 lation of dietary supplements determined appropriate
6 by the Secretary of Health and Human Services or
7 the Commissioner of Food and Drugs.

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