

118TH CONGRESS
1ST SESSION

S. 3512

To amend the Federal Food, Drug, and Cosmetic Act to strengthen requirements related to nutrient information on food labels.

IN THE SENATE OF THE UNITED STATES

DECEMBER 13, 2023

Mr. BLUMENTHAL (for himself, Mr. BOOKER, Mr. WHITEHOUSE, and Mr. MARKEY) introduced the following bill; which was read twice and referred to the Committee on Health, Education, Labor, and Pensions

A BILL

To amend the Federal Food, Drug, and Cosmetic Act to strengthen requirements related to nutrient information on food labels.

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 “Transparency, Read-
5 ability, Understandability, Truth, and Helpfulness in La-
6 beling Act” or the “TRUTH in Labeling Act”.

7 **SEC. 2. FINDINGS.**

8 Congress finds the following:

1 (1) The average American consumes substan-
2 tially more added sugars, sodium, and saturated fat
3 than is recommended by the Dietary Guidelines for
4 Americans published under section 301 of the Na-
5 tional Nutrition Monitoring and Related Research
6 Act of 1990 (7 U.S.C. 5341), potentially increasing
7 their risk for hypertension, type-2 diabetes, and
8 heart disease.

9 (2) A large body of experimental and real-world
10 evidence has demonstrated that front-of-package la-
11 bels that highlight high levels of added sugars, so-
12 dium, and saturated fat can significantly improve
13 the nutritional quality of foods that consumers pur-
14 chase or select.

15 (3) Simplified, contextual information on a food
16 package that compliments the Nutrition Facts label
17 can help consumers make healthy food choices. Ro-
18 bust research shows that front-of-package nutrition
19 labels can be particularly beneficial for busy shop-
20 pers and for those with lower nutrition knowledge.

21 (4) Front-of-package nutrition labeling gives
22 consumers quick and easy access to key information
23 about the healthfulness of foods and can support
24 healthier choices.

1 (5) Studies also show that front-of-package la-
2 beling can improve consumers’ understanding of the
3 relative healthfulness of different foods.

4 **SEC. 3. ADDITIONAL REQUIREMENTS FOR FRONT-OF-PACK-**
5 **AGE LABELING FOR FOODS.**

6 (a) INTERPRETIVE NUTRITION INFORMATION.—Sec-
7 tion 403 of the Federal Food, Drug, and Cosmetic Act
8 (21 U.S.C. 343) is amended by adding at the end the fol-
9 lowing:

10 “(z)(1) Except as provided in subparagraphs (3), (4),
11 and (5) of paragraph (q), if it is food intended for human
12 consumption and is offered for sale and otherwise required
13 to bear nutrition labeling, unless its principal display panel
14 bears interpretive nutrition information.

15 “(2) Final regulations regarding the interpretive nu-
16 trition information required under subparagraph (1) shall
17 meet the following criteria:

18 “(A) There shall be a standardized symbol sys-
19 tem that displays calorie information related to the
20 serving size determined under paragraph (q)(1)(A)
21 and interpretative nutrition information related to
22 the content of any nutrients that the Secretary de-
23 termines the highlighting of which will assist con-
24 sumers in maintaining healthy dietary practices
25 (such as added sugars, sodium, or saturated fat), in-

1 including by highlighting products containing high lev-
2 els of such nutrients.

3 “(B) The information shall—

4 “(i) appear in a consistent location on the
5 principal display panels across products;

6 “(ii) have a prominent design that visually
7 contrasts with existing packaging design; and

8 “(iii) be sufficiently large to be easily leg-
9 ible.

10 “(3) In promulgating regulations regarding the inter-
11 pretive nutrition information required under subpara-
12 graph (1) and the standardized symbol system required
13 under subparagraph (2)(A), the Secretary shall take into
14 account published reports by the Health and Medicine Di-
15 vision of the National Academy of Sciences, Engineering,
16 and Medicine regarding such information, and base regu-
17 lations on the following principles:

18 “(A) Consumers should be able to quickly and
19 easily comprehend the meaning of the system as an
20 indicator of a product’s contribution to a healthy
21 diet without requiring specific or sophisticated nutri-
22 tional knowledge.

23 “(B) The information should be provided to fa-
24 cilitate consumer selection of healthy product op-

1 tions, including among nutritionally at-risk sub-
2 populations.

3 “(C) The Secretary should periodically evaluate
4 the standardized symbol system to assess its effec-
5 tiveness in providing information to facilitate con-
6 sumer selection of healthy product options and the
7 extent to which manufacturers are offering healthier
8 products as a result of the disclosure.

9 “(D) The implementation of the information
10 disclosure should be accompanied by appropriate
11 consumer education and promotion campaigns deter-
12 mined by the Secretary.”.

13 (b) REPORT.—

14 (1) IN GENERAL.—Not later than 5 years after
15 the effective date specified in final regulations issued
16 by the Secretary pursuant to section 4(b), the Sec-
17 retary of Health and Human Services (referred to in
18 this Act as the “Secretary”) shall submit to Con-
19 gress a report that—

20 (A) evaluates whether implementation of
21 the amendment made by subsection (a) has
22 been associated with an increase in the preva-
23 lence of products containing low- or no-calorie
24 sweeteners in the United States food supply;
25 and

1 (B) describes actions that will be taken by
2 the Secretary to further monitor the use of low-
3 and no-calorie sweeteners in such products, if
4 there has been an increase described in sub-
5 paragraph (A).

6 (2) UPDATE.—Not later than 3 years after
7 completion of the report described in paragraph (1),
8 the Secretary shall submit to Congress an update to
9 such report based on more recent data.

10 **SEC. 4. REGULATIONS.**

11 (a) PROPOSED REGULATIONS.—Not later than 2
12 years after the date of enactment of this Act, the Sec-
13 retary shall issue proposed regulations to carry out the
14 amendment made by section 3(a).

15 (b) FINAL REGULATIONS.—Not later than 3 years
16 after the date of enactment of this Act, the Secretary shall
17 finalize the regulations proposed pursuant to subsection
18 (a), which regulations shall specify the date on which the
19 amendment made by section 3(a) shall take effect.