

119TH CONGRESS
2D SESSION

S. 4974

AN ACT

To amend the Federal Food, Drug, and Cosmetic Act with
respect to food safety.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

1 **SECTION 1. SHORT TITLE.**

2 This Act may be cited as the “Making America’s
3 Food Safer Act”.

4 **SEC. 2. EXPANSION OF THE ACCREDITED THIRD-PARTY**
5 **CERTIFICATION PROGRAM.**

6 (a) REVISED DEFINITIONS.—Section 808(a) of the
7 Federal Food, Drug, and Cosmetic Act (21 U.S.C.
8 384d(a)) is amended—

9 (1) by striking paragraph (6) and inserting the
10 following:

11 “(6) ELIGIBLE ENTITY.—The term ‘eligible en-
12 tity’ means a foreign or domestic entity, including a
13 foreign or domestic facility subject to registration
14 under section 415, in the food supply chain that
15 chooses to be audited by an accredited third-party
16 auditor or the audit agent of such accredited third-
17 party auditor.”; and

18 (2) in paragraph (7)(B)—

19 (A) in clause (i), by striking “; or” and in-
20 serting a semicolon;

21 (B) in clause (ii), by striking the period
22 and inserting “; or”; and

23 (C) by adding at the end the following:

24 “(iii) whether a facility is eligible to
25 receive a food or facility certification for

1 other purposes described in subsection
 2 (c)(2)(B)(ii).”.

3 (b) REMOVING LIMITATIONS ON THE USE OF CER-
 4 TIFICATIONS.—Section 808(c)(2) of the Federal Food,
 5 Drug, and Cosmetic Act (21 U.S.C. 384d(c)(2)) is amend-
 6 ed—

7 (1) in subparagraph (A), by striking “food cer-
 8 tification, described in section 801(q), or facility cer-
 9 tification under section 806(a), as appropriate, to
 10 accompany each food shipment for import into the
 11 United States from an eligible entity” and inserting
 12 “food certification or facility certification for pur-
 13 poses described in subparagraph (B), as appro-
 14 priate,”; and

15 (2) by striking subparagraph (B) and inserting
 16 the following:

17 “(B) PURPOSE OF CERTIFICATION.—

18 “(i) IN GENERAL.—The Secretary
 19 shall use certification provided by accred-
 20 ited third-party auditors to—

21 “(I) determine, in conjunction
 22 with any other assurances the Sec-
 23 retary may require under section
 24 801(q), whether a food satisfies the
 25 requirements of such section; and

1 “(II) determine whether a facility
 2 is eligible to be a facility from which
 3 food may be offered for import under
 4 the voluntary qualified importer pro-
 5 gram under section 806.

6 “(ii) OTHER CONSIDERATIONS.—The
 7 Secretary may consider the results of regu-
 8 latory audits and food or facility certifi-
 9 cations provided by accredited third-party
 10 auditors under this section in analyzing
 11 risks and prioritizing inspections and other
 12 regulatory activities, as appropriate for the
 13 protection of public health.”.

14 (c) TECHNICAL AND CONFORMING AMENDMENTS.—
 15 Section 808 of the Federal Food, Drug, and Cosmetic Act
 16 (21 U.S.C. 384d) is amended—

17 (1) in subsection (b)(1)(A)—

18 (A) by striking “ACCREDITATION BODIES”
 19 in the subparagraph heading and all that fol-
 20 lows through “Not later than” in clause (i) and
 21 inserting the following: “ACCREDITATION BOD-
 22 IES—Not later than”; and

23 (B) by striking clause (ii);

24 (2) in subsection (c)—

1 (A) in paragraphs (1) and (2), by striking
 2 “(or, in the case of direct accreditation under
 3 subsection (b)(1)(A)(ii), the Secretary)” each
 4 place it appears;

5 (B) in paragraph (2)(C)(i), by striking
 6 “food certification under section 801(q) or a fa-
 7 cility certification described under this subpara-
 8 graph (B)” and inserting “food certification or
 9 a facility certification described in this section”;

10 (C) in paragraph (6)(A)(i), by striking
 11 “food certified under section 801(q) or from a
 12 facility certified under paragraph (2)(B)” and
 13 inserting “food or a facility certified under this
 14 section”;

15 (D) in paragraph (6)(C), by striking “re-
 16 quirements under section 801(q), of certifying
 17 the food, or the requirements under paragraph
 18 (2)(B) of certifying the entity” and inserting
 19 “requirements for certifying the food or facility
 20 under this section”; and

21 (E) in paragraph (7)(B)(i), by striking “,
 22 through direct accreditation under subsection
 23 (b)(1)(A)(ii) or”; and
 24 (3) in subsection (d)—

25 (A) in paragraph (1), by striking “or”; and

1 (B) at the end of paragraph (2), by strik-
 2 ing the period and inserting “; or”; and

3 (C) by adding at the end the following new
 4 paragraph:

5 “(3) otherwise seeks certification for purposes
 6 of subsection (c)(2)(B)(ii).”.

7 **SEC. 3. SHARING FOOD SAFETY INFORMATION WITH STATE,**
 8 **LOCAL, TRIBAL, AND TERRITORIAL AUTHORI-**
 9 **TIES.**

10 (a) IN GENERAL.—Section 708 of the Federal Food,
 11 Drug, and Cosmetic Act (21 U.S.C. 379) is amended by
 12 adding at the end the following:

13 “(d) SHARING FOOD SAFETY INFORMATION WITH
 14 STATE, LOCAL, TRIBAL, AND TERRITORIAL AUTHORI-
 15 TIES.—

16 “(1) AUTHORIZATION.—Notwithstanding sec-
 17 tion 301(j) and any other law, regulation, or policy,
 18 the Secretary may share, with a State, local, Tribal,
 19 or territorial authority with counterpart functions
 20 related to the protection of public health, unredacted
 21 information in the possession of the Food and Drug
 22 Administration relating to any of the following:

23 “(A) Foodborne illness surveillance data.

24 “(B) Laboratory sampling testing informa-
 25 tion.

1 “(C) Inspectional information and results.

2 “(D) Distribution lists for recalls and out-
3 breaks.

4 “(E) Consumer complaints.

5 “(F) Any other information the Secretary
6 determines will assist such authority in pro-
7 tecting the public.

8 “(2) TIMING.—The Secretary may share infor-
9 mation pursuant to paragraph (1) as soon as is rea-
10 sonably practicable.

11 “(3) LIMITATION ON FURTHER DISCLOSURE.—
12 A State, local, Tribal, or Territorial authority in re-
13 ceipt of information provided by the Secretary under
14 this subsection shall not further disclose such infor-
15 mation without permission of the Food and Drug
16 Administration unless such authority determines
17 that disclosure of such information is necessary to
18 contain a foodborne illness outbreak, carry out a re-
19 call, or carry out other State enforcement activities.

20 “(4) EFFECT OF SUBSECTION.—Nothing in this
21 subsection affects the authority of the Secretary to
22 enter into any written agreement authorized by
23 other provisions of law to share confidential informa-
24 tion.

1 “(e) INFORMATION DISCLOSURE DURING FOOD
 2 SAFETY INCIDENTS.—The Secretary is authorized to dis-
 3 close commercial information obtained from a person that
 4 is protected under section 1905 of title 18, United States
 5 Code, when the disclosure of such information advances
 6 public health protection during a food safety incident, in-
 7 cluding a foodborne illness outbreak, an investigation re-
 8 lated to contaminated food, or a food recall.”.

9 (b) CONFORMING AMENDMENT.—The first sentence
 10 of section 301(j) of the Federal Food, Drug, and Cosmetic
 11 Act (21 U.S.C. 331(j)) is amended—

12 (1) by inserting “to a State, local, Tribal, or
 13 territorial authority as specified in section 708(d),”
 14 after “of the Department,”; and

15 (2) by striking the second period at the end.

16 **SEC. 4. DESTRUCTION OF CERTAIN REFUSED ARTICLES.**

17 Section 801 of the Federal Food, Drug, and Cosmetic
 18 Act (21 U.S.C. 381) is amended by adding at the end the
 19 following:

20 “(v) ORDER TO DESTROY CERTAIN REFUSED ARTI-
 21 CLES.—

22 “(1) IN GENERAL.—For any article that has
 23 been refused admission and is in violation of this
 24 Act, the Secretary of Health and Human Services
 25 may issue to the owner or consignee an order that

1 the article shall be destroyed, without the oppor-
2 tunity to export, if the Secretary of Health and
3 Human Services finds that the article presents a sig-
4 nificant public health concern. Before issuing an
5 order to destroy an article under this subsection, the
6 Secretary of Health and Human Services shall pro-
7 vide for notice and an opportunity to appear before
8 the Secretary of Health and Human Services and in-
9 troduce testimony on the order to destroy. The Sec-
10 retary of Health and Human Services may combine
11 the opportunity to appear before the Secretary and
12 the opportunity to introduce testimony into a single
13 proceeding with respect to an article. The regula-
14 tions under paragraph (2) shall provide that prior to
15 the destruction of any such article, appropriate due
16 process is available to the owner or consignee seek-
17 ing to challenge the decision of the Secretary of
18 Health and Human Services to order destruction.
19 Such process may be combined with the notice and
20 opportunity to appear before the Secretary and in-
21 troduce testimony on the refusal as long as appro-
22 priate notice is provided to the owner or consignee
23 about the potential order to destroy. The Secretary
24 of the Treasury shall cause the owner or consignee
25 to complete the destruction of any such article with-

1 in 90 days of the order for destruction and the
2 owner or consignee shall be responsible for the costs
3 of such destruction.

4 “(2) REGULATIONS.—

5 “(A) PROPOSED.—Not later than 18
6 months after the date of enactment of the Mak-
7 ing America’s Food Safer Act, the Secretary of
8 Health and Human Services shall issue pro-
9 posed regulations to implement paragraph (1),
10 including a framework for due process, allowing
11 for notice and comment on such proposed regu-
12 lations.

13 “(B) FINAL.—Not later than 1 year after
14 the issuance of the proposed regulations under
15 subparagraph (A), the Secretary of Health and
16 Human Services shall promulgate final regula-
17 tions to implement paragraph (1).

18 “(3) EXCEPTIONS.—With respect to importa-
19 tion by an individual of a prescription drug that is
20 not a controlled substance pursuant to section 804(j)
21 and in a manner that is consistent with personal or
22 household use, the authority provided under para-
23 graph (1) shall not apply.

24 “(4) CLARIFICATION.—For purposes of this
25 section, a prescription drug described in paragraph

1 (3) that is imported as described in such paragraph
2 shall not be considered a ‘significant public health
3 concern’.”.

Passed the Senate September 28, 2026.

Attest:

Secretary.

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