International Parental Child Abduction Month'' and expressing the sense of the Senate that Congress should raise awareness of the harm caused by international parental child abduction.

S. RES. 585

At the request of Mr. MENENDEZ, the name of the Senator from Illinois (Mr. DURBIN) was added as a cosponsor of S. Res. 585, a resolution honoring the life, achievements, and legacy of the Honorable Madeleine K. Albright.

STATEMENTS ON INTRODUCED BILLS AND JOINT RESOLUTIONS

By Mr. DURBIN:

S. 4089. A bill to restore entitlement to educational assistance under Veterans Rapid Retraining Program in cases of a closure of an educational institution or a disapproval of a program of education, and for other purposes; considered and passed.

Mr. DURBIN. Mr. President, I ask unanimous consent that the text of the bill be printed in the RECORD.

Without objection, the text of the bill was ordered to be printed in the RECORD, as follows:

S. 4089

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

SECTION 1. SHORT TITLE.

This Act may be cited as the ''Veterans Rapid Retraining Assistance Program Restoration and Recovery Act of 2022''.

SEC. 2. RESTORATION OF ENTITLEMENT UNDER VETERANS RAPID RETRAINING ASSISTANCE PROGRAM.

(a) IN GENERAL.—Section 8006 of the American Rescue Plan Act of 2021 (Public Law 117–2), as amended by the Training in High-demand Roles to Improve Veteran Employment Act (Public Law 117–16), is further amended—

(1) by redesignating subsection (n) as subsection (o); and

(2) by inserting after subsection (m), the following new subsection (n):

''(n) EFFECTS OF CLOSURE OF AN EDUCATIONAL INSTITUTION OR DISAPPROVAL OF A PROGRAM OF EDUCATION.—

''(1) IN GENERAL.—Any payment of retraining assistance under subsection (d)(1) shall not be charged against any entitlement to retraining assistance described in subsection (a) if the Secretary determines that an individual was unable to complete a course or program of education as a result of —

''(A) the closure of an educational institution; or

''(B) the disapproval of a program of education by the State approving agency or the Secretary when acting in the role of the State approving agency.

''(2) PERIOD NOT CHARGED.—The period for which, by reason of this subsection, retraining assistance is not charged shall be equal to the full amount of retraining assistance provided for enrollment in the program of education.

''(3) HALT OF PAYMENTS TO CERTAIN EDUCATIONAL INSTITUTIONS.—In the event of a closure or disapproval, as described in paragraph (1), the educational institution shall not receive any further payments under subsection (d).

''(4) RECOVERY OF FUNDS.—In the event of a closure or disapproval, as described in paragraph (1), any payment already made under subsection (d) to the educational institution shall be considered an overpayment and constitute a liability of such institution to the United States.''.

(b) CONFORMING AMENDMENT.—In subsection (b)(3) of such section, strike the period and insert '', except for an individual described in subsection (n).''.

(c) EFFECTIVE DATE.—The amendments made by this section shall apply as if included in the American Rescue Plan Act of 2021 (Public Law 117–2).

By Mr. DURBIN (for himself and Mr. BRAUN):

S. 4090. A bill to improve transparency and availability of information regarding dietary supplements by amending the Federal Food, Drug, and Cosmetic Act to require manufacturers of dietary supplements to list dietary supplements with the Food and Drug Administration; to the Committee on Health, Education, Labor, and Pensions.

Mr. DURBIN. Mr. President, I ask unanimous consent that the text of the bill be printed in the RECORD.

Without objection, the text of the bill was ordered to be printed in the RECORD, as follows:

S. 4090

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

SECTION 1. SHORT TITLE.

This Act may be cited as the ''Dietary Supplement Listing Act of 2022''.

SEC. 2. REGULATION OF DIETARY SUPPLEMENTS.

(a) IN GENERAL.—Chapter IV of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 341 et seq.) is amended by inserting after section 403C of such Act the following:

''SEC. 403D. DIETARY SUPPLEMENT LISTING REQUIREMENT.

''(a) IN GENERAL.—Each dietary supplement shall be listed with the Secretary in accordance with this section.

''(b) LISTING SUBMISSIONS.—

''(1) IN GENERAL.—Each responsible person, or, if the responsible person is a foreign entity, the United States agent, shall submit to the Secretary in accordance with this section the following information for each dietary supplement that will be marketed:

''(A) Any proprietary name of the dietary supplement and the statement of identity, including brand name and specified flavors, if applicable.

''(B) The full name, address, and telephone number for the responsible person, and the name and e-mail address of the owner, operator, or agent in charge of the responsible person.

''(C) The full name, address, telephone number, and e-mail address for the United States agent, if the responsible person is a foreign entity.

''(D) The full business name and address of all locations at which the responsible person manufactures, packages, labels, or holds the dietary supplement.

''(E) An electronic copy of the label for the dietary supplement, and an electronic copy of the package insert, if any.

''(F) A list of all ingredients in the dietary supplement required to appear on the label under sections 101.4 and 101.36 of title 21, Code of Federal Regulations, including—

''(i) the amount per serving of each listed ingredient, if such information is required to appear on the label; and

''(ii) if required by section 101.36 of title 21, Code of Federal Regulations, the percent of the daily value of each listed ingredient.

''(G) The number of servings per container for each container size.

''(H) The conditions of use.

''(I) Warnings and precautions.

''(J) Statements regarding major food allergens, as defined in section 201(qq) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321(qq)).

''(K) The dosage form, such as pill, capsule, liquid, or powder.

''(L) Any claim that—

''(i) characterizes the relationship of any nutrient which is of the type required by section 403(q)(1) or section (q)(2) to be in the label or labeling of the food to a disease or a health-related condition; or

''(ii) is subject to notification under section 403(r)(6) that appears in the supplement's labeling.

''(M) The unique dietary supplement identifier for the product, provided in accordance with paragraph (3).

''(2) FORMAT.—A listing submitted under this section shall be in such electronic form and manner as the Secretary may prescribe. The Secretary shall promptly confirm, electronically, receipt of a complete listing under this section.

''(3) UNIQUE LISTING IDENTIFICATION NUMBERS.—

''(A) IN GENERAL.—The Secretary shall establish a unique dietary supplement identifier system that shall be used by the responsible person under this section.

''(B) RESERVATION OF NUMBERS.—The system shall allow a responsible person to reserve multiple dietary supplement identifier numbers in advance of listing.

''(C) USE REQUIREMENT.—Any unique dietary supplement identifier shall be used only in connection with the product for which the identifier was used during the listing process.

''(4) SUBMISSION DATES.—A responsible person under this section shall report to the Secretary the listing information described in paragraph (1) pursuant to the following timelines:

''(A) IN GENERAL.—

''(i) EXISTING DIETARY SUPPLEMENTS.—In the case of a dietary supplement that is being offered in interstate commerce on the date that is 18 months after the date of enactment of the Dietary Supplement Listing Act of 2022, a listing for each such dietary supplement formulation introduced or delivered for introduction into interstate commerce by the responsible person for commercial distribution shall be submitted by the responsible person with the Secretary under this section not later than 60 days after the date that is 18 months after the date of enactment of such Act.

''(ii) NEW DIETARY SUPPLEMENTS.—In the case of a dietary supplement that is not being offered in interstate commerce on the date that is 18 months after the date of enactment. of the Dietary Supplement Listing Act of 2022, a listing for each such dietary supplement formulation introduced or delivered for introduction into interstate commerce by the responsible person for commercial distribution which has not been included in any listing previously submitted by the responsible person to the Secretary under this section shall be submitted to the Secretary prior to introducing the dietary supplement into interstate commerce.

''(B) REFORMULATIONS.—A listing of each dietary supplement formulation introduced by the responsible person for commercial distribution that has a label that differs for such dietary supplement from the representative label provided under subsection (a) with respect to the product name, amount of dietary ingredients, or other distinguishing characteristics such as dosage form (such as pill, capsule, liquid, or powder) shall be submitted to the Secretary not later than 15 business days after introducing the dietary