

America's Cup Race Management during the 1-year period beginning on the last date of such defense.

(2) **AMERICA'S CUP RACE MANAGEMENT.**—The term "America's Cup Race Management" means the entity established to provide for independent, professional, and neutral race management of the America's Cup sailing competitions.

(3) **ELIGIBILITY CERTIFICATION.**—The term "Eligibility Certification" means a certification issued under section 4.

(4) **ELIGIBLE VESSEL.**—The term "eligible vessel" means a competing vessel or supporting vessel of any registry that—

(A) is recognized by America's Cup Race Management as an official competing vessel, or supporting vessel of, the 34th America's Cup, as evidenced in writing to the Administrator of the Maritime Administration of the Department of Transportation;

(B) transports not more than 25 individuals, in addition to the crew;

(C) is not a ferry (as defined under section 2101(10b) of title 46, United States Code);

(D) does not transport individuals in point-to-point service for hire; and

(E) does not transport merchandise between ports in the United States.

(5) **SUPPORTING VESSEL.**—The term "supporting vessel" means a vessel that is operating in support of the 34th America's Cup by—

(A) positioning a competing vessel on the race course;

(B) transporting equipment and supplies utilized for the staging, operations, or broadcast of the competition; or

(C) transporting individuals who—

(i) have not purchased tickets or directly paid for their passage; and

(ii) who are engaged in the staging, operations, or broadcast of the competition, race team personnel, members of the media, or event sponsors.

SEC. 3. AUTHORIZATION OF ELIGIBLE VESSELS.

Notwithstanding sections 55102, 55103, and 55111 of title 46, United States Code, an eligible vessel, operating only in preparation for, or in connection with, the 34th America's Cup competition, may position competing vessels and may transport individuals and equipment and supplies utilized for the staging, operations, or broadcast of the competition from and around the ports in the United States.

SEC. 4. CERTIFICATION.

(a) **REQUIREMENT.**—A vessel may not operate under section 3 unless the vessel has received an Eligibility Certification.

(b) **ISSUANCE.**—The Administrator of the Maritime Administration of the Department of Transportation is authorized to issue an Eligibility Certification with respect to any vessel that the Administrator determines, in his or her sole discretion, meets the requirements set forth in section 2(4).

SEC. 5. ENFORCEMENT.

Notwithstanding sections 55102, 55103, and 55111 of title 46, United States Code, an Eligibility Certification shall be conclusive evidence to the Secretary of the Department of Homeland Security of the qualification of the vessel for which it has been issued to participate in the 34th America's Cup as a competing vessel or a supporting vessel.

SEC. 6. PENALTY.

Any vessel participating in the 34th America's Cup as a competing vessel or supporting vessel that has not received an Eligibility Certification or is not in compliance with section 12112 of title 46, United States Code, shall be subject to the applicable penalties provided in chapters 121 and 551 of title 46, United States Code.

SEC. 7. WAIVERS.

(a) **IN GENERAL.**—Notwithstanding sections 12112 and 12132 and chapter 551 of title 46,

United States Code, the Secretary of the department in which the Coast Guard is operating may issue a certificate of documentation with a coastwise endorsement for each of the following vessels:

(1) M/V GEYSIR (United States official number 622178).

(2) OCEAN VERITAS (IMO number 7366805).

(3) LUNA (United States official number 280133).

(b) **DOCUMENTATION OF LNG TANKERS.**—

(1) **IN GENERAL.**—Notwithstanding sections 12112 and 12132 and chapter 551 of title 46, United States Code, the Secretary of the department in which the Coast Guard is operating may issue a certificate of documentation with a coastwise endorsement for each of the following vessels:

(A) LNG GEMINI (United States official number 595752).

(B) LNG LEO (United States official number 595753).

(C) LNG VIRGO (United States official number 595755).

(2) **LIMITATION ON OPERATION.**—Coastwise trade authorized under paragraph (1) shall be limited to carriage of natural gas, as that term is defined in section 3(13) of the Deepwater Port Act of 1974 (33 U.S.C. 1502(13)).

(3) **TERMINATION OF EFFECTIVENESS OF ENDORSEMENTS.**—The coastwise endorsement issued under paragraph (1) for a vessel shall expire on the date of the sale of the vessel by the owner of the vessel on the date of enactment of this Act to a person who is not related by ownership or control to such owner.

(c) **OPERATION OF A DRY DOCK.**—A vessel transported in Dry Dock #2 (State of Alaska registration AIDEA FDD-2) is not merchandise for purposes of section 55102 of title 46, United States Code, if, during such transportation, Dry Dock #2 remains connected by a utility or other connecting line to pierside moorage.

The amendment was ordered to be engrossed and the bill to be read a third time.

The bill (H.R. 3321), as amended, was read the third time and passed.

AMERICAN MEDICAL ISOTOPES PRODUCTION ACT OF 2011

Mr. LEVIN. Mr. President, I now ask unanimous consent that the Senate proceed to the consideration of Calendar No. 53, S. 99.

The PRESIDING OFFICER. The clerk will report the bill by title.

The bill clerk read as follows:

A bill (S. 99) to promote the production of molybdenum-99 in the United States for medical isotope production, and to condition and phase out the export of highly enriched uranium for the production of medical isotopes.

There being no objection, the Senate proceeded to consider the bill, which had been reported from the Committee on Energy and Natural Resources, with an amendment to strike all after the enacting clause and insert in lieu thereof the following:

SECTION 1. SHORT TITLE.

This Act may be cited as the "American Medical Isotopes Production Act of 2011".

SEC. 2. DEFINITIONS.

In this Act:

(1) **DEPARTMENT.**—The term "Department" means the Department of Energy.

(2) **HIGHLY ENRICHED URANIUM.**—The term "highly enriched uranium" means uranium enriched to 20 percent or greater in the isotope U-235.

(3) **LOW ENRICHED URANIUM.**—The term "low enriched uranium" means uranium enriched to less than 20 percent in the isotope U-235.

(4) **SECRETARY.**—The term "Secretary" means the Secretary of Energy.

SEC. 3. IMPROVING THE RELIABILITY OF DOMESTIC MEDICAL ISOTOPE SUPPLY.

(a) **MEDICAL ISOTOPE DEVELOPMENT PROJECTS.**—

(1) **IN GENERAL.**—The Secretary shall establish a technology-neutral program—

(A) to evaluate and support projects for the production in the United States, without the use of highly enriched uranium, of significant quantities of molybdenum-99 for medical uses;

(B) to be carried out in cooperation with non-Federal entities; and

(C) the costs of which shall be shared in accordance with section 988 of the Energy Policy Act of 2005 (42 U.S.C. 16352).

(2) **CRITERIA.**—Projects shall be judged against the following primary criteria:

(A) The length of time necessary for the proposed project to begin production of molybdenum-99 for medical uses within the United States.

(B) The capability of the proposed project to produce a significant percentage of United States demand for molybdenum-99 for medical uses.

(C) The cost of the proposed project.

(3) **EXEMPTION.**—An existing reactor in the United States fueled with highly enriched uranium shall not be disqualified from the program if the Secretary determines that—

(A) there is no alternative nuclear reactor fuel, enriched in the isotope U-235 to less than 20 percent, that can be used in that reactor;

(B) the reactor operator has provided assurances that, whenever an alternative nuclear reactor fuel, enriched in the isotope U-235 to less than 20 percent, can be used in that reactor, it will use that alternative in lieu of highly enriched uranium; and

(C) the reactor operator has provided a current report on the status of its efforts to convert the reactor to an alternative nuclear reactor fuel enriched in the isotope U-235 to less than 20 percent, and an anticipated schedule for completion of conversion.

(4) **PUBLIC PARTICIPATION AND REVIEW.**—The Secretary shall—

(A) develop a program plan and annually update the program plan through public workshops; and

(B) use the Nuclear Science Advisory Committee to conduct annual reviews of the progress made in achieving the program goals.

(5) **AUTHORIZATION OF APPROPRIATIONS.**—There are authorized to be appropriated to the Secretary for carrying out the program under paragraph (1) \$143,000,000 for the period encompassing fiscal years 2011 through 2014.

(b) **DEVELOPMENT ASSISTANCE.**—The Secretary shall establish a program to provide assistance for—

(1) the development of fuels, targets, and processes for domestic molybdenum-99 production that do not use highly enriched uranium; and

(2) commercial operations using the fuels, targets, and processes described in paragraph (1).

(c) **URANIUM LEASE AND TAKE-BACK.**—

(1) **IN GENERAL.**—The Secretary shall establish a program to make low-enriched uranium available, through lease contracts, for irradiation for the production of molybdenum-99 for medical uses.

(2) **TITLE.**—The lease contracts shall provide for the producers of the molybdenum-99 to take title to and be responsible for the molybdenum-99 created by the irradiation, processing, or purification of uranium leased under this section.

(3) **DUTIES.**—

(A) **SECRETARY.**—The lease contracts shall require the Secretary—

(i) to retain responsibility for the final disposition of spent nuclear fuel created by the irradiation, processing, or purification of uranium

leased under this section for the production of medical isotopes; and

(i) to take title to and be responsible for the final disposition of radioactive waste created by the irradiation, processing, or purification of uranium leased under this section for which the Secretary determines the producer does not have access to a disposal path.

(B) PRODUCER.—The producer of the spent nuclear fuel and radioactive waste shall accurately characterize, appropriately package, and transport the spent nuclear fuel and radioactive waste prior to acceptance by the Department.

(4) COMPENSATION.—

(A) IN GENERAL.—Subject to subparagraph (B), the lease contracts shall provide for compensation in cash amounts equivalent to prevailing market rates for the sale of comparable uranium products and for compensation in cash amounts equivalent to the net present value of the cost to the Federal Government for—

(i) the final disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3); and

(ii) other costs associated with carrying out the uranium lease and take-back program authorized by this subsection.

(B) DISCOUNT RATE.—The discount rate used to determine the net present value of costs described in subparagraph (A)(ii) shall be not greater than the average interest rate on marketable Treasury securities.

(5) AUTHORIZED USE OF FUNDS.—The Secretary may obligate and expend funds received under leases entered into under this subsection, which shall remain available until expended, for the purpose of carrying out the activities authorized by this Act, including activities related to the final disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3).

(6) EXCHANGE OF URANIUM FOR SERVICES.—The Secretary shall not barter or otherwise sell or transfer uranium in any form in exchange for—

(A) services related to the final disposition of the spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3); or

(B) any other services associated with carrying out the uranium lease and take-back program authorized by this subsection.

(d) COORDINATION OF ENVIRONMENTAL REVIEWS.—The Department and the Nuclear Regulatory Commission shall ensure to the maximum extent practicable that environmental reviews for the production of the medical isotopes shall complement and not duplicate each review.

(e) OPERATIONAL DATE.—The Secretary shall establish a program as described in subsection (c)(3) not later than 3 years after the date of enactment of this Act.

(f) RADIOACTIVE WASTE.—Notwithstanding section 2 of the Nuclear Waste Policy Act of 1982 (42 U.S.C. 10101), radioactive material resulting from the production of medical isotopes that has been permanently removed from a reactor or subcritical assembly and for which there is no further use shall be considered low-level radioactive waste if the material is acceptable under Federal requirements for disposal as low-level radioactive waste.

(g) AUTHORIZATION OF APPROPRIATIONS.—There are authorized to be appropriated to the Secretary \$5,000,000 for the establishment of a program for the final disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under subsection (c).

SEC. 4. EXPORTS.

Section 134 of the Atomic Energy Act of 1954 (42 U.S.C. 2160d) is amended by striking subsection c. and inserting the following:

“c. Effective 7 years after the date of enactment of the American Medical Isotopes Production Act of 2011, the Commission may not issue a license for the export of highly enriched uranium from the United States for the purposes of medical isotope production.

“d. The period referred to in subsection b. may be extended for no more than 6 years if, no earlier than 6 years after the date of enactment of the American Medical Isotopes Production Act of 2011, the Secretary of Energy certifies to the Committee on Energy and Commerce of the House of Representatives and the Committee on Energy and Natural Resources of the Senate that—

“(1) there is insufficient global supply of molybdenum-99 produced without the use of highly enriched uranium available to satisfy the domestic United States market; and

“(2) the export of United States-origin highly enriched uranium for the purposes of medical isotope production is the most effective temporary means to increase the supply of molybdenum-99 to the domestic United States market.

“e. To ensure public review and comment, the development of the certification described in subsection c. shall be carried out through announcement in the Federal Register.

“f. At any time after the restriction of export licenses provided for in subsection b. becomes effective, if there is a critical shortage in the supply of molybdenum-99 available to satisfy the domestic United States medical isotope needs, the restriction of export licenses may be suspended for a period of no more than 12 months, if—

“(1) the Secretary of Energy certifies to the Congress that the export of United States-origin highly enriched uranium for the purposes of medical isotope production is the only effective temporary means to increase the supply of molybdenum-99 necessary to meet United States medical isotope needs during that period; and

“(2) the Congress enacts a Joint Resolution approving the temporary suspension of the restriction of export licenses.

“g. As used in this section—

“(1) the term ‘alternative nuclear reactor fuel or target’ means a nuclear reactor fuel or target which is enriched to less than 20 percent in the isotope U-235;

“(2) the term ‘highly enriched uranium’ means uranium enriched to 20 percent or more in the isotope U-235;

“(3) a fuel or target ‘can be used’ in a nuclear research or test reactor if—

“(A) the fuel or target has been qualified by the Reduced Enrichment Research and Test Reactor Program of the Department of Energy; and

“(B) use of the fuel or target will permit the large majority of ongoing and planned experiments and medical isotope production to be conducted in the reactor without a large percentage increase in the total cost of operating the reactor; and

“(4) the term ‘medical isotope’ includes molybdenum-99, iodine-131, xenon-133, and other radioactive materials used to produce a radiopharmaceutical for diagnostic or therapeutic procedures or for research and development.”

SEC. 5. REPORT ON DISPOSITION OF EXPORTS.

Not later than 1 year after the date of the enactment of this Act, the Chairman of the Nuclear Regulatory Commission, after consulting with other relevant agencies, shall submit to the Congress a report detailing the current disposition of previous United States exports of highly enriched uranium used as fuel or targets in a nuclear research or test reactor, including—

- (1) their location;
- (2) whether they are irradiated;
- (3) whether they have been used for the purpose stated in their export license;
- (4) whether they have been used for an alternative purpose and, if so, whether such alternative purpose has been explicitly approved by the Commission;
- (5) the year of export, and reimportation, if applicable;
- (6) their current physical and chemical forms; and
- (7) whether they are being stored in a manner which adequately protects against theft and unauthorized access.

SEC. 6. DOMESTIC MEDICAL ISOTOPE PRODUCTION.

(a) IN GENERAL.—Chapter 10 of the Atomic Energy Act of 1954 (42 U.S.C. 2131 et seq.) is amended by adding at the end the following:

“SEC. 112. DOMESTIC MEDICAL ISOTOPE PRODUCTION.—a. The Commission may issue a license, or grant an amendment to an existing license, for the use in the United States of highly enriched uranium as a target for medical isotope production in a nuclear reactor, only if, in addition to any other requirement of this Act—

“(1) the Commission determines that—

“(A) there is no alternative medical isotope production target, enriched in the isotope U-235 to less than 20 percent, that can be used in that reactor; and

“(B) the proposed recipient of the medical isotope production target has provided assurances that, whenever an alternative medical isotope production target can be used in that reactor, it will use that alternative in lieu of highly enriched uranium; and

“(2) the Secretary of Energy has certified that the United States Government is actively supporting the development of an alternative medical isotope production target that can be used in that reactor.

“b. As used in this section—

“(1) the term ‘alternative medical isotope production target’ means a nuclear reactor target which is enriched to less than 20 percent of the isotope U-235;

“(2) a target ‘can be used’ in a nuclear research or test reactor if—

“(A) the target has been qualified by the Reduced Enrichment Research and Test Reactor Program of the Department of Energy; and

“(B) use of the target will permit the large majority of ongoing and planned experiments and medical isotope production to be conducted in the reactor without a large percentage increase in the total cost of operating the reactor;

“(3) the term ‘highly enriched uranium’ means uranium enriched to 20 percent or more in the isotope U-235; and

“(4) the term ‘medical isotope’ includes molybdenum-99, iodine-131, xenon-133, and other radioactive materials used to produce a radiopharmaceutical for diagnostic or therapeutic procedures or for research and development.”

(b) TABLE OF CONTENTS.—The table of contents for the Atomic Energy Act of 1954 is amended by inserting the following new item at the end of the items relating to chapter 10 of title I:

“Sec. 112. Domestic medical isotope production.”

SEC. 7. ANNUAL DEPARTMENT REPORTS.

(a) IN GENERAL.—Not later than 1 year after the date of enactment of this Act, and annually thereafter for 5 years, the Secretary shall report to Congress on Department actions to support the production in the United States, without the use of highly enriched uranium, of molybdenum-99 for medical uses.

(b) CONTENTS.—The reports shall include the following:

(1) For medical isotope development projects—

(A) the names of any recipients of Department support under section 3;

(B) the amount of Department funding committed to each project;

(C) the milestones expected to be reached for each project during the year for which support is provided;

(D) how each project is expected to support the increased production of molybdenum-99 for medical uses;

(E) the findings of the evaluation of projects under section 3(a)(2); and

(F) the ultimate use of any Department funds used to support projects under section 3.

(2) A description of actions taken in the previous year by the Secretary to ensure the safe disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under section 3(c).

SEC. 8. NATIONAL ACADEMY OF SCIENCES REPORT.

(a) *IN GENERAL.*—The Secretary shall enter into an arrangement with the National Academy of Sciences to conduct a study of the state of molybdenum-99 production and utilization, to be provided to Congress not later than 5 years after the date of enactment of this Act.

(b) *CONTENTS.*—The report shall include the following:

(1) *For molybdenum-99 production—*
(A) a list of all facilities in the world producing molybdenum-99 for medical uses, including an indication of whether these facilities use highly enriched uranium in any way;

(B) a review of international production of molybdenum-99 over the previous 5 years, including—

(i) whether any new production was brought online;

(ii) whether any facilities halted production unexpectedly; and

(iii) whether any facilities used for production were decommissioned or otherwise permanently removed from service; and

(C) an assessment of progress made in the previous 5 years toward establishing domestic production of molybdenum-99 for medical uses, including the extent to which other medical isotopes that have been produced with molybdenum-99, such as iodine-131 and xenon-133, are being used for medical purposes.

(2) An assessment of the progress made by the Department and others to eliminate all world-

wide use of highly enriched uranium in reactor fuel, reactor targets, and medical isotope production facilities.

SEC. 9. BUDGETARY EFFECTS.

The budgetary effects of this Act, for the purpose of complying with the Statutory Pay-As-You-Go-Act of 2010, shall be determined by reference to the latest statement titled “Budgetary Effects of PAYGO Legislation” for this Act, submitted for printing in the Congressional Record by the Chairman of the Senate Budget Committee, provided that such statement has been submitted prior to the vote on passage.

Mr. LEVIN. Mr. President, I ask unanimous consent that the committee-reported substitute amendment be considered, the Bingaman amendment, which is at the desk, be agreed to, the committee-reported amendment, as amended, be agreed to, the bill, as amended, be read a third time, and the budgetary pay-go statement at the desk be read.

The PRESIDING OFFICER. Without objection, it is so ordered.

The amendment (No. 1223) was agreed to, as follows:

On page 15, line 14, strike “establish” and insert “carry out”.

On page 17, strike lines 15 through 19.

On page 17, line 21, strike “establish” and insert “carry out”.

On page 21, strike lines 12 through 16.

On page 29, after line 23, add the following:

SEC. 9. REPEAL.

The Nuclear Safety Research, Development, and Demonstration Act of 1980 (42 U.S.C. 9701 et seq.) is repealed.

On page 30, line 1, strike “9” and insert “10”.

The committee amendment in the nature of a substitute, as amended, was agreed to.

The bill, as amended, was ordered to be engrossed for a third reading and was read the third time.

The PRESIDING OFFICER. The clerk will read the pay-go statement.

The bill clerk read as follows:

Mr. Conrad: This is the Statement of Budgetary Effects of PAYGO Legislation for S. 99 as amended.

Total Budgetary Effects of S. 99 for the 5-year Statutory PAYGO Scorecard: \$0.

Total Budgetary Effects of S. 99 for the 10-year Statutory PAYGO Scorecard: \$0.

Also submitted for the Record as part of this statement is a table prepared by the Congressional Budget Office, which provides additional information on the budgetary effects of this Act, as follows:

CBO ESTIMATE OF THE STATUTORY PAY-AS-YOU-GO EFFECTS FOR S. 99, THE AMERICAN MEDICAL ISOTOPES PROTECTION ACT OF 2011, AS REPORTED BY THE SENATE COMMITTEE ON ENERGY AND NATURAL RESOURCES ON MAY 18, 2011, AND WITH A SUBSEQUENT AMENDMENT PROVIDED TO CBO ON NOVEMBER 17, 2011

By fiscal year, in millions of dollars—

	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021	2012-2016	2012-2021
Statutory Pay-As-You-Go Impact	0	0	0	0	0	0	0	0	0	0	0	0

S. 99 would direct the Secretary of Energy to lease low-enriched uranium to producers of molybdenum-99. CBO estimates that enacting S. 99 would affect receipts generated from such resources, but that any net changes to such receipts would be negligible in any given year.
Source: Congressional Budget Office.

Mr. LEVIN. Mr. President, I ask unanimous consent that the bill, as amended, be passed, the motions to reconsider be laid upon the table with no intervening action or debate, and that any related statements be printed in the RECORD.

The PRESIDING OFFICER. Without objection, it is so ordered.

The bill (S. 99), as amended, was passed, as follows:

S. 99

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 “American Medical Isotopes Production Act of 2011”.

SEC. 2. DEFINITIONS.

In this Act:

(1) **DEPARTMENT.**—The term “Department” means the Department of Energy.

(2) **HIGHLY ENRICHED URANIUM.**—The term “highly enriched uranium” means uranium enriched to 20 percent or greater in the isotope U-235.

(3) **LOW ENRICHED URANIUM.**—The term “low enriched uranium” means uranium enriched to less than 20 percent in the isotope U-235.

(4) **SECRETARY.**—The term “Secretary” means the Secretary of Energy.

SEC. 3. IMPROVING THE RELIABILITY OF DOMESTIC MEDICAL ISOTOPE SUPPLY.

(a) **MEDICAL ISOTOPE DEVELOPMENT PROJECTS.**—

(1) *IN GENERAL.*—The Secretary shall carry out a technology-neutral program—

(A) to evaluate and support projects for the production in the United States, without the

use of highly enriched uranium, of significant quantities of molybdenum-99 for medical uses;

(B) to be carried out in cooperation with non-Federal entities; and

(C) the costs of which shall be shared in accordance with section 988 of the Energy Policy Act of 2005 (42 U.S.C. 16352).

(2) **CRITERIA.**—Projects shall be judged against the following primary criteria:

(A) The length of time necessary for the proposed project to begin production of molybdenum-99 for medical uses within the United States.

(B) The capability of the proposed project to produce a significant percentage of United States demand for molybdenum-99 for medical uses.

(C) The cost of the proposed project.

(3) **EXEMPTION.**—An existing reactor in the United States fueled with highly enriched uranium shall not be disqualified from the program if the Secretary determines that—

(A) there is no alternative nuclear reactor fuel, enriched in the isotope U-235 to less than 20 percent, that can be used in that reactor;

(B) the reactor operator has provided assurances that, whenever an alternative nuclear reactor fuel, enriched in the isotope U-235 to less than 20 percent, can be used in that reactor, it will use that alternative in lieu of highly enriched uranium; and

(C) the reactor operator has provided a current report on the status of its efforts to convert the reactor to an alternative nuclear reactor fuel enriched in the isotope U-235 to less than 20 percent, and an anticipated schedule for completion of conversion.

(4) **PUBLIC PARTICIPATION AND REVIEW.**—The Secretary shall—

(A) develop a program plan and annually update the program plan through public workshops; and

(B) use the Nuclear Science Advisory Committee to conduct annual reviews of the progress made in achieving the program goals.

(b) **DEVELOPMENT ASSISTANCE.**—The Secretary shall carry out a program to provide assistance for—

(1) the development of fuels, targets, and processes for domestic molybdenum-99 production that do not use highly enriched uranium; and

(2) commercial operations using the fuels, targets, and processes described in paragraph (1).

(c) **URANIUM LEASE AND TAKE-BACK.**—

(1) *IN GENERAL.*—The Secretary shall establish a program to make low-enriched uranium available, through lease contracts, for irradiation for the production of molybdenum-99 for medical uses.

(2) **TITLE.**—The lease contracts shall provide for the producers of the molybdenum-99 to take title to and be responsible for the molybdenum-99 created by the irradiation, processing, or purification of uranium leased under this section.

(3) **DUTIES.**—

(A) **SECRETARY.**—The lease contracts shall require the Secretary—

(i) to retain responsibility for the final disposition of spent nuclear fuel created by the irradiation, processing, or purification of uranium leased under this section for the production of medical isotopes; and

(ii) to take title to and be responsible for the final disposition of radioactive waste

created by the irradiation, processing, or purification of uranium leased under this section for which the Secretary determines the producer does not have access to a disposal path.

(B) PRODUCER.—The producer of the spent nuclear fuel and radioactive waste shall accurately characterize, appropriately package, and transport the spent nuclear fuel and radioactive waste prior to acceptance by the Department.

(4) COMPENSATION.—

(A) IN GENERAL.—Subject to subparagraph (B), the lease contracts shall provide for compensation in cash amounts equivalent to prevailing market rates for the sale of comparable uranium products and for compensation in cash amounts equivalent to the net present value of the cost to the Federal Government for—

(i) the final disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3); and

(ii) other costs associated with carrying out the uranium lease and take-back program authorized by this subsection.

(B) DISCOUNT RATE.—The discount rate used to determine the net present value of costs described in subparagraph (A)(ii) shall be not greater than the average interest rate on marketable Treasury securities.

(5) AUTHORIZED USE OF FUNDS.—The Secretary may obligate and expend funds received under leases entered into under this subsection, which shall remain available until expended, for the purpose of carrying out the activities authorized by this Act, including activities related to the final disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3).

(6) EXCHANGE OF URANIUM FOR SERVICES.—The Secretary shall not barter or otherwise sell or transfer uranium in any form in exchange for—

(A) services related to the final disposition of the spent nuclear fuel and radioactive waste for which the Department is responsible under paragraph (3); or

(B) any other services associated with carrying out the uranium lease and take-back program authorized by this subsection.

(d) COORDINATION OF ENVIRONMENTAL REVIEWS.—The Department and the Nuclear Regulatory Commission shall ensure to the maximum extent practicable that environmental reviews for the production of the medical isotopes shall complement and not duplicate each review.

(e) OPERATIONAL DATE.—The Secretary shall establish a program as described in subsection (c)(3) not later than 3 years after the date of enactment of this Act.

(f) RADIOACTIVE WASTE.—Notwithstanding section 2 of the Nuclear Waste Policy Act of 1982 (42 U.S.C. 10101), radioactive material resulting from the production of medical isotopes that has been permanently removed from a reactor or subcritical assembly and for which there is no further use shall be considered low-level radioactive waste if the material is acceptable under Federal requirements for disposal as low-level radioactive waste.

SEC. 4. EXPORTS.

Section 134 of the Atomic Energy Act of 1954 (42 U.S.C. 2160d) is amended by striking subsection c. and inserting the following:

“c. Effective 7 years after the date of enactment of the American Medical Isotopes Production Act of 2011, the Commission may not issue a license for the export of highly enriched uranium from the United States for the purposes of medical isotope production.

“d. The period referred to in subsection b. may be extended for no more than 6 years if,

no earlier than 6 years after the date of enactment of the American Medical Isotopes Production Act of 2011, the Secretary of Energy certifies to the Committee on Energy and Commerce of the House of Representatives and the Committee on Energy and Natural Resources of the Senate that—

“(1) there is insufficient global supply of molybdenum-99 produced without the use of highly enriched uranium available to satisfy the domestic United States market; and

“(2) the export of United States-origin highly enriched uranium for the purposes of medical isotope production is the most effective temporary means to increase the supply of molybdenum-99 to the domestic United States market.

“e. To ensure public review and comment, the development of the certification described in subsection c. shall be carried out through announcement in the Federal Register.

“f. At any time after the restriction of export licenses provided for in subsection b. becomes effective, if there is a critical shortage in the supply of molybdenum-99 available to satisfy the domestic United States medical isotope needs, the restriction of export licenses may be suspended for a period of no more than 12 months, if—

“(1) the Secretary of Energy certifies to the Congress that the export of United States-origin highly enriched uranium for the purposes of medical isotope production is the only effective temporary means to increase the supply of molybdenum-99 necessary to meet United States medical isotope needs during that period; and

“(2) the Congress enacts a Joint Resolution approving the temporary suspension of the restriction of export licenses.

“g. As used in this section—

“(1) the term ‘alternative nuclear reactor fuel or target’ means a nuclear reactor fuel or target which is enriched to less than 20 percent in the isotope U-235;

“(2) the term ‘highly enriched uranium’ means uranium enriched to 20 percent or more in the isotope U-235;

“(3) a fuel or target ‘can be used’ in a nuclear research or test reactor if—

“(A) the fuel or target has been qualified by the Reduced Enrichment Research and Test Reactor Program of the Department of Energy; and

“(B) use of the fuel or target will permit the large majority of ongoing and planned experiments and medical isotope production to be conducted in the reactor without a large percentage increase in the total cost of operating the reactor; and

“(4) the term ‘medical isotope’ includes molybdenum-99, iodine-131, xenon-133, and other radioactive materials used to produce a radiopharmaceutical for diagnostic or therapeutic procedures or for research and development.”

SEC. 5. REPORT ON DISPOSITION OF EXPORTS.

Not later than 1 year after the date of the enactment of this Act, the Chairman of the Nuclear Regulatory Commission, after consulting with other relevant agencies, shall submit to the Congress a report detailing the current disposition of previous United States exports of highly enriched uranium used as fuel or targets in a nuclear research or test reactor, including—

- (1) their location;
- (2) whether they are irradiated;
- (3) whether they have been used for the purpose stated in their export license;
- (4) whether they have been used for an alternative purpose and, if so, whether such alternative purpose has been explicitly approved by the Commission;
- (5) the year of export, and reimportation, if applicable;

(6) their current physical and chemical forms; and

(7) whether they are being stored in a manner which adequately protects against theft and unauthorized access.

SEC. 6. DOMESTIC MEDICAL ISOTOPE PRODUCTION.

(a) IN GENERAL.—Chapter 10 of the Atomic Energy Act of 1954 (42 U.S.C. 2131 et seq.) is amended by adding at the end the following: “SEC. 112. DOMESTIC MEDICAL ISOTOPE PRODUCTION.—

“a. The Commission may issue a license, or grant an amendment to an existing license, for the use in the United States of highly enriched uranium as a target for medical isotope production in a nuclear reactor, only if, in addition to any other requirement of this Act—

“(1) the Commission determines that—

“(A) there is no alternative medical isotope production target, enriched in the isotope U-235 to less than 20 percent, that can be used in that reactor; and

“(B) the proposed recipient of the medical isotope production target has provided assurances that, whenever an alternative medical isotope production target can be used in that reactor, it will use that alternative in lieu of highly enriched uranium; and

“(2) the Secretary of Energy has certified that the United States Government is actively supporting the development of an alternative medical isotope production target that can be used in that reactor.

“b. As used in this section—

“(1) the term ‘alternative medical isotope production target’ means a nuclear reactor target which is enriched to less than 20 percent of the isotope U-235;

“(2) a target ‘can be used’ in a nuclear research or test reactor if—

“(A) the target has been qualified by the Reduced Enrichment Research and Test Reactor Program of the Department of Energy; and

“(B) use of the target will permit the large majority of ongoing and planned experiments and medical isotope production to be conducted in the reactor without a large percentage increase in the total cost of operating the reactor;

“(3) the term ‘highly enriched uranium’ means uranium enriched to 20 percent or more in the isotope U-235; and

“(4) the term ‘medical isotope’ includes molybdenum-99, iodine-131, xenon-133, and other radioactive materials used to produce a radiopharmaceutical for diagnostic or therapeutic procedures or for research and development.”

(b) TABLE OF CONTENTS.—The table of contents for the Atomic Energy Act of 1954 is amended by inserting the following new item at the end of the items relating to chapter 10 of title I:

“Sec. 112. Domestic medical isotope production.”

SEC. 7. ANNUAL DEPARTMENT REPORTS.

(a) IN GENERAL.—Not later than 1 year after the date of enactment of this Act, and annually thereafter for 5 years, the Secretary shall report to Congress on Department actions to support the production in the United States, without the use of highly enriched uranium, of molybdenum-99 for medical uses.

(b) CONTENTS.—The reports shall include the following:

(1) For medical isotope development projects—

(A) the names of any recipients of Department support under section 3;

(B) the amount of Department funding committed to each project;

(C) the milestones expected to be reached for each project during the year for which support is provided;

(D) how each project is expected to support the increased production of molybdenum-99 for medical uses;

(E) the findings of the evaluation of projects under section 3(a)(2); and

(F) the ultimate use of any Department funds used to support projects under section 3.

(2) A description of actions taken in the previous year by the Secretary to ensure the safe disposition of spent nuclear fuel and radioactive waste for which the Department is responsible under section 3(c).

SEC. 8. NATIONAL ACADEMY OF SCIENCES REPORT.

(a) IN GENERAL.—The Secretary shall enter into an arrangement with the National Academy of Sciences to conduct a study of the state of molybdenum-99 production and utilization, to be provided to Congress not later than 5 years after the date of enactment of this Act.

(b) CONTENTS.—The report shall include the following:

(1) For molybdenum-99 production—

(A) a list of all facilities in the world producing molybdenum-99 for medical uses, including an indication of whether these facilities use highly enriched uranium in any way;

(B) a review of international production of molybdenum-99 over the previous 5 years, including—

(i) whether any new production was brought online;

(ii) whether any facilities halted production unexpectedly; and

(iii) whether any facilities used for production were decommissioned or otherwise permanently removed from service; and

(C) an assessment of progress made in the previous 5 years toward establishing domestic production of molybdenum-99 for medical uses, including the extent to which other medical isotopes that have been produced with molybdenum-99, such as iodine-131 and xenon-133, are being used for medical purposes.

(2) An assessment of the progress made by the Department and others to eliminate all worldwide use of highly enriched uranium in reactor fuel, reactor targets, and medical isotope production facilities.

SEC. 9. REPEAL.

The Nuclear Safety Research, Development, and Demonstration Act of 1980 (42 U.S.C. 9701 et seq.) is repealed.

SEC. 10. BUDGETARY EFFECTS.

The budgetary effects of this Act, for the purpose of complying with the Statutory Pay-As-You-Go-Act of 2010, shall be determined by reference to the latest statement titled "Budgetary Effects of PAYGO Legislation" for this Act, submitted for printing in the Congressional Record by the Chairman of the Senate Budget Committee, provided that such statement has been submitted prior to the vote on passage.

AMERICAN EDUCATION WEEK

Mr. LEVIN. Mr. President, I ask unanimous consent that the Senate proceed to the immediate consideration of S. Res. 332 which was submitted earlier today by Senator HAGAN.

The PRESIDING OFFICER. The clerk will report the resolution by title.

The bill clerk read as follows:

A resolution (S. Res. 332) supporting the goals and ideals of American Education Week.

There being no objection, the Senate proceeded to consider the resolution.

Mr. LEVIN. Mr. President, I ask unanimous consent that the resolution be agreed to, the preamble be agreed to, the motions to reconsider be laid upon the table, with no intervening action or debate, and that any statements relating to the measure be printed in the RECORD.

The PRESIDING OFFICER. Without objection, it is so ordered.

The resolution (S. Res. 332) was agreed to.

The preamble was agreed to.

The resolution, with its preamble, reads as follows:

S. RES. 332

Whereas the National Education Association has designated November 13 through November 19, 2011, as the 90th annual observance of American Education Week;

Whereas public schools are the backbone of the Nation's democracy, providing young people with the tools they need to maintain the Nation's precious values of freedom, civility, and equality;

Whereas by equipping young people in the United States with both practical skills and broader intellectual abilities, public schools give them hope for, and access to, a productive future;

Whereas people working in the field of public education, be they teachers, principals, higher education faculty and staff, custodians, substitute educators, bus drivers, clerical workers, food service professionals, workers in skilled trades, health and student service workers, security guards, technical employees, or librarians, work tirelessly to serve children and communities throughout the Nation with care and professionalism; and

Whereas public schools are community linchpins, bringing together adults, children, educators, volunteers, business leaders, and elected officials in a common enterprise: Now, therefore, be it

Resolved, That the Senate—

(1) supports the goals and ideals of American Education Week; and

(2) encourages the people of the United States to observe National Education Week by reflecting on the positive impact of all those who work together to educate children.

WELCOMING AND COMMENDING THE GOVERNMENT OF JAPAN

Mr. LEVIN. I ask unanimous consent that the Senate proceed to the immediate consideration of S. Res. 333 which was submitted earlier today by Senator FEINSTEIN.

The PRESIDING OFFICER. The clerk will report the resolution by title.

The bill clerk read as follows:

A resolution (S. Res. 333) welcoming and commending the Government of Japan for extending an official apology to all United States former prisoners of war from the Pacific War and establishing in 2010 a visitation program to Japan for surviving veterans, family members, and descendants.

There being no objection, the Senate proceeded to consider the resolution.

Mrs. FEINSTEIN. President, I rise today in support of this resolution honoring former World War II U.S. POWs from the Pacific theater and acknowledging the steps the Japanese Government has made to heal the wounds of the past.

My friend and colleague from California, Representative MIKE HONDA, introduced this resolution in the House and I am proud to follow suit here in the Senate. I applaud his leadership on this important matter.

Our resolution welcomes and commends the Government of Japan for extending an official apology to all U.S. former prisoners of war from the Pacific War and establishing in 2010 a visitation program to Japan for surviving veterans, their families, and descendants.

The resolution appreciates the recent efforts by the Government of Japan toward historic apologies for the war crimes of Imperial Japan.

The resolution requests that the Government of Japan continue its new Japanese/American POW Friendship Program of reconciliation and remembrance.

It requests that the Government of Japan respect the wishes and sensibilities of the United States former prisoners of war by supporting and encouraging programs for lasting remembrance and reconciliation that recognize their sacrifices, history, and forced labor.

It acknowledges the work of the Department of State in advocating for the United States Prisoners of War from the Pacific war, and it applauds the persistence, dedication, and patriotism of the members and descendants of the American Defenders of Bataan and Corregidor.

According to the Congressional Research Service, approximately 27,000 U.S. prisoners of war were held by Imperial Japanese forces during World War II.

They were often subject to brutal and inhumane treatment.

They were starved and denied adequate medical care and were forced to perform slave labor for private Japanese companies.

American POWs toiled in mines, factories, shipyards, and steel mills for hours every day under extremely dangerous conditions. Many suffered health problems long after their time as POWs had ended.

Some 40 percent of POWs perished and never returned home to their loved ones.

We owe these brave heroes a debt that can never be fully repaid. It is critical that we never forget their sacrifice.

A lot has changed since the end of the war.

Japan has emerged from the ashes of war to develop into one of our closest friends and allies and a responsible member of the international community.

Our relationship is sustained by shared values of democracy, human rights, and the rule of law.

The American POWs—those that survived—returned home and tried to move on with their lives.

They completed their education, got married, started families, began new