Mr. COTTON, Mr. CRAPO, Mr. CRUZ, Mr. DAINES, Mr. DONNELLY, Mr. DURBIN, Mr. ENZI, Mrs. ERNST, Mrs. FEINSTEIN, Mrs. FISCHER, Mr. FLAKE, Mr. FRANKEN, Mr. GARDNER, Mrs. GILLIBRAND, Mr. GRAHAM, Mr. GRASSLEY, Mr. HATCH, Mr. HEINRICH, Ms. HEITKAMP, Mr. HELLER, Ms. HIRONO, Mr. HOEVEN, Mr. INHOFE, Mr. ISAKSON, Mr. JOHNSON, Mr. KAINE, Mr. KING, Mr. KIRK, Ms. KLOBUCHAR, Mr. LANKFORD, Mr. LEAHY, Mr. LEE, Mr. MANCHIN, Mr. MARKEY, Mr. MCCAIN, Mrs. MCCASKILL, Mr. MENENDEZ, Mr. MERKLEY, Ms. MIKULSKI, Mr. MORAN, Ms. MURKOWSKI, Mr. MURPHY, Mrs. MURRAY, Mr. PAUL, Mr. PERDUE, Mr. PETERS, Mr. PORTMAN, Mr. REED, Mr. RISCH, Mr. ROBERTS, Mr. ROUNDS, Mr. SANDERS, Mr. SASSE, Mr. SCHATZ, Mr. SCHUMER, Mr. SCOTT, Mr. SESSIONS, Mrs. SHAHEEN, Mr. SHELBY, Ms. STABENOW, Mr. SULLIVAN, Mr. TESTER, Mr. THUNE, Mr. TILLIS, Mr. TOOMEY, Mr. UDALL, Mr. VITTER, Mr. WARNER, Ms. WARREN, Mr. WHITEHOUSE, Mr. WICKER, and Mr. WYDEN):

S. Res. 496. A resolution condemning the terrorist attack on the Pulse Orlando nightclub, honoring the memory of the victims of the attack, offering condolences to and expressing support for their families and friends and all those affected, and applauding the dedication and bravery of law enforcement, emergency response, and counterterrorism officials in responding to the attack; considered and agreed to.

By Ms. STABENOW (for herself and Mr. PETERS):

S. Res. 497. A resolution honoring the life and legacy of Gordon ''Gordie'' Howe; considered and agreed to.

By Mr. BLUMENTHAL (for himself, Ms. COLLINS, Ms. AYOTTE, Mrs. MCCASKILL, Mr. GRASSLEY, Mr. CASEY, Mr. COTTON, Mr. TILLIS, Mr. MURPHY, and Mr. HELLER):

S. Res. 498. A resolution designating June 15, 2016, as ''World Elder Abuse Awareness Day'' ; considered and agreed to.

By Mr. TOOMEY (for himself and Mr. CASEY):

S. Res. 499. A resolution congratulating the Pittsburgh Penguins for winning the 2016 Stanley Cup hockey championship; considered and agreed to.

By Mr. CORNYN (for himself, Mrs. BOXER, Ms. BALDWIN, Mr. BENNET, Mr. BOOKER, Mr. BROWN, Mr. BURR, Mr. CASEY, Mr. COCHRAN, Mr. CRAPO, Mr. CRUZ, Mr. DURBIN, Mrs. FEINSTEIN, Mr. FRANKEN, Mrs. GILLIBRAND, Ms. HIRONO, Mr. INHOFE, Mr. KAINE, Ms. KLOBUCHAR, Mr. LANKFORD, Mr. LEAHY, Mr. LEE, Mr. MARKEY, Mr. MERKLEY, Ms. MURKOWSKI, Mr. MURPHY, Mrs. MURRAY, Mr. NELSON, Mr. PAUL, Mr. PETERS, Mr. REID, Mr. RUBIO, Mr. SCHUMER, Mr. SCOTT, Ms. STABENOW, Mr. TILLIS, Mr. WARNER, Mr. WHITEHOUSE, Mr. WICKER, Mr. WYDEN, and Ms. WARREN):

S. Res. 500. A resolution designating June 19, 2016, as ''Juneteenth Independence Day'' in recognition of June 19, 1865, the date on which slavery legally came to an end in the United States; considered and agreed to.

By Mrs. ERNST (for herself and Mrs. BOXER):

S. Con. Res. 41. A concurrent resolution expressing the sense of Congress on the Peshmerga of the Kurdistan Region of Iraq; to the Committee on Foreign Relations.

ADDITIONAL COSPONSORS

S. 804

At the request of Ms. COLLINS, the name of the Senator from Missouri (Mr. BLUNT) was added as a cosponsor of S. 804, a bill to amend title XVIII of the Social Security Act to specify coverage of continuous glucose monitoring devices, and for other purposes.

S. 1555

At the request of Ms. HIRONO, the names of the Senator from New Hampshire (Ms. AYOTTE), the Senator from South Carolina (Mr. GRAHAM), the Senator from Utah (Mr. HATCH), the Senator from Maine (Mr. KING) and the Senator from Idaho (Mr. CRAPO) were added as cosponsors of S. 1555, a bill to award a Congressional Gold Medal, collectively, to the Filipino veterans of World War II, in recognition of the dedicated service of the veterans during World War II.

S. 2213

At the request of Mr. BLUMENTHAL, the name of the Senator from Pennsylvania (Mr. CASEY) was added as a cosponsor of S. 2213, a bill to prohibit firearms dealers from selling a firearm prior to the completion of a background check.

S. 2218

At the request of Mr. THUNE, the name of the Senator from Nevada (Mr. HELLER) was added as a cosponsor of S. 2218, a bill to amend the Internal Revenue Code of 1986 to treat certain amounts paid for physical activity, fitness, and exercise as amounts paid for medical care.

S. 2311

At the request of Mr. HELLER, the name of the Senator from New Mexico (Mr. HEINRICH) was added as a cosponsor of S. 2311, a bill to amend the Public Health Service Act to authorize the Secretary of Health and Human Services, acting through the Administrator of the Health Resources and Services Administration, to make grants to States for screening and treatment for maternal depression.

S. 2373

At the request of Ms. CANTWELL, the name of the Senator from Iowa (Mrs. ERNST) was added as a cosponsor of S. 2373, a bill to amend title XVIII of the Social Security Act to provide for Medicare coverage of certain lymphedema compression treatment items as items of durable medical equipment.

S. 2604

At the request of Mr. WARNER, the name of the Senator from North Dakota (Ms. HEITKAMP) was added as a cosponsor of S. 2604, a bill to establish in the legislative branch the National Commission on Security and Technology Challenges.

S. 2736

At the request of Mr. THUNE, the name of the Senator from Mississippi (Mr. WICKER) was added as a cosponsor of S. 2736, a bill to improve access to durable medical equipment for Medicare beneficiaries under the Medicare program, and for other purposes.

S. 2873

At the request of Mr. HATCH, the name of the Senator from Massachusetts (Ms. WARREN) was added as a cosponsor of S. 2873, a bill to require studies and reports examining the use of, and opportunities to use, technology-enabled collaborative learning and capacity building models to improve programs of the Department of Health and Human Services, and for other purposes.

S. 2912

At the request of Mr. JOHNSON, the name of the Senator from Texas (Mr. CRUZ) was added as a cosponsor of S. 2912, a bill to authorize the use of unapproved medical products by patients diagnosed with a terminal illness in accordance with State law, and for other purposes.

S. 2924

At the request of Mr. REID, the name of the Senator from New York (Mrs. GILLIBRAND) was added as a cosponsor of S. 2924, a bill to award a Congressional Gold Medal to former United States Senator Max Cleland.

S. 3053

At the request of Mr. CASEY, the name of the Senator from New Hampshire (Mrs. SHAHEEN) was added as a cosponsor of S. 3053, a bill to prevent a person who has been convicted of a misdemeanor hate crime, or received an enhanced sentence for a misdemeanor because of hate or bias in its commission, from obtaining a firearm.

S. 3059

At the request of Ms. CANTWELL, the name of the Senator from Alaska (Ms. MURKOWSKI) was added as a cosponsor of S. 3059, a bill to reauthorize and amend the John H. Prescott Marine Mammal Rescue and Response Grant Program and for other purposes.

S. 3060

At the request of Ms. HEITKAMP, the names of the Senator from Montana (Mr. TESTER) and the Senator from Virginia (Mr. WARNER) were added as cosponsors of S. 3060, a bill to provide an exception from certain group health plan requirements for qualified small employer health reimbursement arrangements.

S. RES. 482

At the request of Mrs. SHAHEEN, the name of the Senator from Michigan (Mr. PETERS) was added as a cosponsor of S. Res. 482, a resolution urging the European Union to designate Hizballah in its entirety as a terrorist organization and to increase pressure on the organization and its members to the fullest extent possible.

AMENDMENT NO. 4691

At the request of Mr. MURPHY, the name of the Senator from Connecticut (Mr. BLUMENTHAL) was added as a cosponsor of amendment No. 4691 intended to be proposed to H.R. 2578, a bill making appropriations for the Departments of Commerce and Justice,

Science, and Related Agencies for the fiscal year ending September 30, 2016, and for other purposes.

AMENDMENT NO. 4719

At the request of Ms. BALDWIN, the names of the Senator from Connecticut (Mr. BLUMENTHAL), the Senator from Minnesota (Mr. FRANKEN) and the Senator from Vermont (Mr. LEAHY) were added as cosponsors of amendment No. 4719 intended to be proposed to H.R. 2578, a bill making appropriations for the Departments of Commerce and Justice, Science, and Related Agencies for the fiscal year ending September 30, 2016, and for other purposes.

AMENDMENT NO. 4720

At the request of Mrs. FEINSTEIN, the names of the Senator from Rhode Island (Mr. WHITEHOUSE), the Senator from Virginia (Mr. WARNER), the Senator from Oregon (Mr. WYDEN), the Senator from New York (Mrs. GILLIBRAND), the Senator from Vermont (Mr. SANDERS), the Senator from Wisconsin (Ms. BALDWIN), the Senator from Colorado (Mr. BENNET), the Senator from Pennsylvania (Mr. CASEY), the Senator from Rhode Island (Mr. REED) and the Senator from Maryland (Mr. CARDIN) were added as cosponsors of amendment No. 4720 proposed to H.R. 2578, a bill making appropriations for the Departments of Commerce and Justice, Science, and Related Agencies for the fiscal year ending September 30, 2016, and for other purposes.

STATEMENTS ON BILLS AND JOINT RESOLUTIONS

By Mr. DURBIN:

S. 3075. A bill to establish programs related to prevention of prescription opioid misuse, and for other purposes; to the Committee on Finance.

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

There being no objection, the text of the bill was ordered to be printed in the RECORD, as follows:

S. 3075

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 ''Addiction Prevention and Responsible Opioid Practices Act''.

SEC. 2. OPIOID ACTION PLAN.

(a) ADVISORY COMMITTEE.—

(1) NEW DRUG APPLICATION.—Except as provided in paragraph (4), prior to the approval of a new drug that is an opioid under section 505 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355), the Commissioner of Food and Drugs shall refer such drug to an advisory committee of the Food and Drug Administration to seek recommendations from such Committee.

(2) PEDIATRIC OPIOID LABELING.—The Commissioner of Food and Drugs shall convene the Pediatric Advisory Committee of the Food and Drug Administration to seek recommendations from such Committee regarding a framework for the inclusion of information in the labeling of drugs that are opioids relating to the use of such drugs in pediatric populations before such Commissioner approves any labeling changes for drugs that are opioids intended for use in pediatric populations.

(3) PUBLIC HEALTH EXEMPTION.—If the Commissioner of Food and Drugs finds that referring a new opioid drug or drugs to an advisory committee of the Food and Drug Administration as required under paragraph (1) is not in the interest of protecting and promoting public health, and has submitted a notice containing the rationale for such a finding to the Committee on Health, Education, Labor, and Pensions of the Senate and the Committee on Energy and Commerce of the House of Representatives, or if the matter that would be considered by such advisory committee with respect to any such drug or drugs concerns bioequivalence, sameness of active ingredient, or other criteria applicable to applications submitted under section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)), the Commissioner shall not be required to refer such drug or drugs to an advisory committee as required under paragraph (1).

(4) SUNSET.—Unless Congress reauthorizes paragraphs (1) and (2), the requirements of such paragraphs shall cease to be effective on October 1, 2022.

(b) EDUCATION FOR PRESCRIBERS OF OPIOIDS.—Not later than 1 year after the date of enactment of this Act, the Secretary of Health and Human Services, acting through the Commissioner of Food and Drugs, as part of the Food and Drug Administration's evaluation of the Extended-Release/Long-Acting Opioid Analgesics Risk Evaluation and Mitigation Strategy, and in consultation with the Director of the Centers for Disease Control and Prevention, the Director of the National Institutes of Health, the Administrator of the Agency for Healthcare Research and Quality, the Administrator of the Drug Enforcement Administration, and relevant stakeholders, shall develop recommendations regarding education programs for prescribers of opioids required to be disseminated under section 505–1 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355–1), including recommendations for which prescribers should participate in such programs and how often participation in such programs is necessary.

(c) GUIDANCE.—Not later than 1 year after the date of enactment of this Act, the Commissioner of Food and Drugs shall issue guidance on if and how the approved labeling of a drug that is an opioid and is the subject of an application under section 505(j) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)) may include statements that such drug deters abuse.

SEC. 3. OPIOID INFORMATIONAL DOCUMENTS.

(a) IN GENERAL.—Subchapter A of chapter V of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 351 et seq.) is amended by inserting after section 505–1 the following:

''SEC. 505–2. OPIOID INFORMATIONAL DOCUMENTS.

''(a) DEVELOPMENT OF MATERIALS.—The Commissioner shall develop informational documents describing to consumers of opioid drugs the risk factors for opioid-related harm, and shall submit such documents to the Director of the Centers for Disease Control and Prevention for approval.

''(b) LABELING REQUIREMENT.—The manufacturer of any opioid drug approved under section 505 shall ensure that the appropriate informational documents developed under subsection (a), and approved by the Director of the Centers for Disease Control and Prevention, are included in the labeling of such drug.''.

(b) ENFORCEMENT.—Section 502 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 352) is amended by adding at the end the following:

''(dd) If it is an opioid drug and the labeling does not include the informational documents required under section 505–2.''.

SEC. 4. STRENGTHENING CONSIDERATIONS FOR DEA NARCOTIC QUOTAS.

Section 306 of the Controlled Substances Act (21 U.S.C. 826) is amended by adding at the end the following:

''(i)(1) In fixing manufacturing quotas under this section the Attorney General shall take into consideration the impact of the manufacturing quotas on diversion and efforts to reduce the costs, injuries, and deaths associated with the abuse of prescription opioids and heroin in the United States.

''(2)(A) Not later than 1 year after the date of enactment of this subsection and every year thereafter, the Attorney General shall publish the approved manufacturing quota for each manufacturer of fentanyl, oxycodone, hyrdocodone, oxymorphone, and hyrdomorphone for that year.

''(B) For any year in which the approved manufacturing quota for a manufacturer for any substance described in subparagraph (A) is higher than the approved manufacturing quota for a manufacturer for the substance in the previous year, the Attorney General shall publish a report explaining why the public health benefits of increasing such quota outweigh the consequences of having an increased volume of such substance available for sale, and potential diversion, in the United States.

''(C) For any substance described in subparagraph (A) that is approved under section 505 of the Federal Food, Drug, and Cosmetic Act after the date of enactment of this subsection, the Attorney General shall publish a report explaining what factors were taken into consideration in setting the manufacturing quota for the substance.

''(3) Not later than 90 days after the date of enactment of this subsection, the Attorney General shall submit to Congress a report on—

''(A) how the Attorney General will ensure that the process of fixing manufacturing quotas under this section takes into consideration efforts to reduce the costs, injuries, and deaths associated with the abuse of prescription opioids and heroin;

''(B) formal steps that will be taken to improve data collection from approved drug collection receptacles, mail-back programs, and take-back events on the volume and class of controlled substances that are collected; and

''(C) how the information described in subparagraphs (A) and (B) will influence the quota-setting process of the Attorney General in the following year.''.

SEC. 5. CONTINUING MEDICAL EDUCATION AND PRESCRIPTION DRUG MONITORING PROGRAM REGISTRATION FOR PRESCRIBERS.

Section 303 of the Controlled Substances Act (21 U.S.C. 823) is amended by adding at the end the following:

''(k)(1) The Attorney General shall not register, or renew the registration of, a practitioner under subsection (f) who is licensed under State law to prescribe controlled substances in schedule II, III, or IV, unless the practitioner submits to the Attorney General, for each such registration or renewal request, a written certification that—

''(A)(i) the practitioner has, during the 1-year period preceding the registration or renewal request, completed a training program described in paragraph (2); or

''(ii) the practitioner, during the applicable registration period, will not prescribe such controlled substances in amounts in excess of a 72-hour supply (for which no refill is available); and