

critical strategy to improve the educational attainment of those students: Now, therefore, be it

Resolved, That the Senate—

(1) congratulates Vision To Learn on helping 200,000 students;

(2) recognizes Vision To Learn as a national leader in providing school-based vision care and commends the strides the organization has made in that effort; and

(3) supports the mission of Vision To Learn to ensure that no child goes without the glasses needed for that child to succeed in school and in life.

SENATE RESOLUTION 223—EXPRESSING SUPPORT FOR THE DESIGNATION OF MAY 17, 2019, AS “DIPG PEDIATRIC BRAIN CANCER AWARENESS DAY” TO RAISE AWARENESS OF AND ENCOURAGE RESEARCH ON DIFFUSE INTRINSIC PONTINE GLIOMA TUMORS AND PEDIATRIC CANCERS IN GENERAL

Mr. RUBIO (for himself, Mr. REED, Mrs. HYDE-SMITH, and Mr. CASEY) submitted the following resolution; which was considered and agreed to:

S. RES. 223

Whereas diffuse intrinsic pontine glioma (referred to in this preamble as “DIPG”) tumors regularly affect 200 to 400 children in the United States each year;

Whereas brain tumors are the leading cause of cancer-related death among children;

Whereas, during childhood, DIPG tumors are—

(1) the second most common type of malignant brain tumor; and

(2) the leading cause of pediatric brain cancer deaths;

Whereas, with respect to a child who is diagnosed with a DIPG tumor and receives treatment for a DIPG tumor, the median amount of time that the child survives after diagnosis is only 9 months;

Whereas, with respect to an individual who is diagnosed with a DIPG tumor, the rate of survival 5 years after diagnosis is approximately 2 percent;

Whereas the average age at which a child is diagnosed with a DIPG tumor is between 5 and 9 years, resulting in a life expectancy approximately 70 years shorter than the average life expectancy in the United States; and

Whereas the prognosis for children diagnosed with DIPG tumors has not improved over the past 40 years: Now, therefore, be it

Resolved, That the Senate—

(1) supports—

(A) designating May 17, 2019, as “DIPG Pediatric Brain Cancer Awareness Day”; and

(B) efforts—

(i) to better understand diffuse intrinsic pontine glioma tumors;

(ii) to develop effective treatments for diffuse intrinsic pontine glioma tumors; and

(iii) to provide comprehensive care for children with diffuse intrinsic pontine glioma tumors and their families; and

(2) encourages all individuals in the United States to become more informed about—

(A) diffuse intrinsic pontine glioma tumors;

(B) pediatric brain cancer in general; and

(C) challenges relating to research on pediatric cancers and ways to advance such research.

SENATE RESOLUTION 224—SUPPORTING THE GOALS AND IDEALS OF NATIONAL NURSES WEEK, TO BE OBSERVED FROM MAY 6 THROUGH MAY 12, 2019

Mr. WICKER (for himself and Mr. MERKLEY) submitted the following resolution; which was considered and agreed to:

S. RES. 224

Whereas, beginning in 1991, National Nurses Week is celebrated annually from May 6, also known as “National Recognition Day for Nurses”, through May 12, the birthday of Florence Nightingale, the founder of modern nursing;

Whereas National Nurses Week is a time of year to reflect on the important contributions that nurses make to provide safe, high-quality health care;

Whereas nurses are known to be patient advocates, acting to protect the lives of individuals under their care;

Whereas nurses represent the largest single component of the health care professions, with an estimated population of 4,000,000 registered nurses in the United States;

Whereas nurses are leading in the delivery of quality care in a transformed health care system that improves patient outcomes and safety;

Whereas the Future of Nursing report of the Institute of Medicine has called for the nursing profession to meet the call for leadership in a team-based delivery model;

Whereas, when nurse staffing levels increase, the risk of patient complications and lengthy hospital stays decreases, resulting in cost savings;

Whereas nurses are experienced researchers, and the work of nurses encompasses a wide scope of scientific inquiry, including clinical research, health systems and outcomes research, and nursing education research;

Whereas nurses provide care that is sensitive to the regional and community customs of individuals needing care;

Whereas nurses are well-positioned to provide leadership to eliminate health care disparities that exist in the United States;

Whereas nurses are the cornerstone of the public health infrastructure, promoting healthy lifestyles and educating communities on disease prevention and health promotion;

Whereas nurses help inform, educate, and work closely with legislators to improve—

(1) the education, retention, recruitment, and practice of all nurses; and

(2) the health and safety of the patients for whom the nurses care;

Whereas there is a need—

(1) to strengthen nursing workforce development programs at all levels, including the number of doctorally prepared faculty members; and

(2) to provide education to the nurse research scientists who can develop new nursing care models to improve the health status of the diverse population of the United States;

Whereas nurses touch the lives of the people of the United States through every stage of life; and

Whereas nursing has been voted the most honest and ethical profession in the United States: Now, therefore, be it

Resolved, That the Senate—

(1) supports the goals and ideals of National Nurses Week, as founded by the American Nurses Association;

(2) recognizes the significant contributions of nurses to the health care system in the United States; and

(3) encourages the people of the United States to observe National Nurses Week with

appropriate recognition, ceremonies, activities, and programs to demonstrate the importance of nurses to the everyday lives of patients.

SENATE RESOLUTION 225—SUPPORTING THE GOALS OF INTERNATIONAL MYALGIC ENCEPHALOMYELITIS/CHRONIC FATIGUE SYNDROME AWARENESS DAY

Mr. MARKEY (for himself, Ms. COLLINS, Mr. CRAMER, Mr. BLUMENTHAL, Mr. COONS, Mr. HOEVEN, Mr. BOOKER, Mr. VAN HOLLEN, Mrs. FEINSTEIN, Mr. KING, Ms. STABENOW, Ms. SINEMA, Mr. CASEY, Ms. HARRIS, and Ms. WARREN) submitted the following resolution; which was considered and agreed to:

S. RES. 225

Whereas the National Academy of Medicine (referred to in this preamble as “NAM”), formerly known as the Institute of Medicine, has found Myalgic Encephalomyelitis/Chronic Fatigue Syndrome (referred to in this preamble as “ME/CFS”) to be “a serious, chronic, complex, and systemic disease that frequently and dramatically limits the activities of affected patients”;

Whereas between 836,000 and 2,500,000 individuals of all ages, races, and sexes in the United States are believed to be afflicted with ME/CFS, with millions more afflicted by ME/CFS worldwide, and the vast majority of individuals with ME/CFS are undiagnosed or misdiagnosed;

Whereas ME/CFS is approximately 4 times more prevalent in women than in men;

Whereas ME/CFS is a chronic disease with no known cure and leaves ¼ of individuals with ME/CFS housebound or bedbound for extended periods of time;

Whereas 50 to 75 percent of individuals with ME/CFS cannot work or attend school;

Whereas medical expenses and lost productivity related to ME/CFS cost the economy of the United States an estimated \$17,000,000,000 to \$24,000,000,000 annually;

Whereas the cause of ME/CFS is unknown, there is no diagnostic test for ME/CFS, and there is no treatment for ME/CFS that is approved by the Food and Drug Administration;

Whereas NAM has noted a “paucity of research” on ME/CFS and that “more research is essential”;

Whereas the Centers for Disease Control and Prevention has called ME/CFS “America’s Hidden Health Crisis”;

Whereas individuals with ME/CFS struggle to find doctors to care for them, and ME/CFS is included in less than ⅓ of medical school curricula;

Whereas, in recognition of the dearth of research on ME/CFS and the profound impact that the disease has on individuals with ME/CFS and their loved ones and caretakers, the National Institutes of Health (referred to in this preamble as the “NIH”) is “committed to unraveling the underlying biologic cause(s) of ME/CFS as swiftly as possible, and promoting research that will inform the development of effective strategies for treatment and prevention of this devastating condition”;

Whereas, in 2017, 11 Institutes at the NIH and the Office of the Director of the NIH contributed more than \$7,000,000 in grants to assist in establishing Collaborative Research Centers and a Data Management Coordinating Center to improve the coordination of ME/CFS research and help accelerate understanding of ME/CFS; and