May 18, 2017

The Honorable Susan Collins
U.S. Senate
413 Dirksen Senate Office Building
Washington, D.C. 20510

The Honorable Tammy Baldwin
U.S. Senate
709 Hart Senate Office Building
Washington, D.C. 20510

The Honorable Jim Langevin
U.S. House of Representatives
2077 Rayburn House Office Building
Washington, D.C. 20515

The Honorable Gregg Harper
U.S. House of Representatives
2227 Rayburn House Office Building
Washington, D.C. 20515

Dear Senator Collins, Senator Baldwin, Representative Langevin and Representative Harper:

We, the undersigned national organizations representing all ages and disabilities, are writing to offer our fervent support for and endorsement of the Lifespan Respite Care Reauthorization Act being to reauthorize the Lifespan Respite Care Program at $15 million a year for five years. We also want to thank you for your leadership in supporting the nation’s family caregivers.

Every day, millions of American families are faced with unexpected illness, disease, or disability. A soldier is injured in war, a spouse develops multiple sclerosis or Alzheimer’s disease, or a child is diagnosed with a developmental or physical disability or chronic illness. These are but a few examples of events that can forever change an individual’s and family’s trajectory.

While each situation is unique, the one thing that they often have in common is the incredible value of family caregivers. Forty-three million family caregivers provide a vast majority of our nation’s long-term care, permitting individuals of all ages to remain in their communities and avoid or delay nursing home or foster care placements. AARP has estimated that family caregivers provide $470 billion in uncompensated care annually to adults, a staggering statistic that exceeds 2013 federal and state spending on Medicaid health services and long-term services and supports.

While the benefits of family caregiving are plentiful, caregiving can take its toll—with older spousal family caregivers experiencing higher mortality rates, rates of acute and chronic conditions, and depression than noncaregivers. Respite—short-term care that offers individuals or family members temporary relief from the daily routine and stress of providing care—is a critical component to bolstering family stability and maintaining family caregiver health and well-being. Respite is a frequently requested support service among family caregivers, but 85% of family caregivers receive no respite. Not surprisingly, high burden family caregivers (defined as those who assist their loved one
with personal care such as getting dressed or bathing) cite lack of respite as one of their top three concerns.

To help provide family caregivers the support they need, the Lifespan Respite Care Program was enacted in 2006 with strong bipartisan support. The program provides competitive grants to states to establish or enhance statewide Lifespan Respite systems that maximize existing resources and help ensure that quality respite is available and accessible to all family caregivers. With more than half of care recipients under age 75 and more than one-third under age 50, Lifespan Respite rightly recognizes caregiving as a lifespan issue and serves families regardless of age or disability.

Though the program has been drastically underfunded since its inception, thirty-five states and the District of Columbia have received grants and are engaged in impressive work such as identifying and coordinating respite services available through various state agencies, including veterans caregiver services; helping unserved families pay for respite through participant-directed voucher programs or mini-grants to community-based agencies; building respite capacity by recruiting and training respite workers and volunteers; and raising awareness about respite through public education campaigns. Originally authorized through Fiscal Year 2011, enactment of the Lifespan Respite Care Reauthorization Act is necessary to continue this excellent momentum, better coordinate and supply respite care to our nation’s 43 million family caregivers through statewide Lifespan Respite programs, and ensure that states are able to sustain the great work they have begun and still allow new states to receive a grant.

We thank you for your commitment to individuals living with disabilities, older individuals in need of assistance and support, and the loved ones who care for them and we look forward to continuing to work with you as the bill moves forward. If you would like more information, please contact Jill Kagan at jkagan@archrespite.org.

Sincerely,

AARP
Alzheimer's Association
Alzheimer's Foundation of America (AFA)
Alzheimer's Impact Movement
American Association of Caregiving Youth
American Association on Health and Disability
The Arc of the United States
Association of University Centers on Disabilities
Autism Society
Brain Injury Association of America
Caring Across Generations
Caregiver Action Network
Child Welfare League of America
Christopher and Dana Reeve Foundation
Easterseals
Elizabeth Dole Foundation
Family Voices
Generations United
The Jewish Federations of North America
LeadingAge
Lupus Foundation
The Michael J. Fox Foundation
National Alliance for Caregiving
National Alliance of Children’s Trust & Prevention Funds
National Association for Home Care & Hospice
National Association of Area Agencies on Aging (n4a)
National Association of Head Injury Administrators
National Association of Social Workers (NASW)
National Association of State Directors of Developmental Disabilities Services
National Association of State Directors of Special Education (NASDSE)
National Center on Caregiving, Family Caregiver Alliance
National Council on Aging
National Respite Coalition
National Multiple Sclerosis Society
Paralyzed Veterans of America
Rosalynn Carter Institute for Caregiving
United Cerebral Palsy
United Spinal Association