Caregivers are not well integrated into U.S. healthcare systems. Across care settings, caregivers lack formal and consistent roles on the care receiver’s care team and can struggle to manage their care recipient’s needs with their own. Only 29 percent of caregivers report being asked by a healthcare professional about their caregiving needs; this rate drops to 13 percent when the question referred to what the caregiver needed to be able to care for themselves (2020 Report of Caregiving in the U.S.; AARP and National Alliance for Caregiving). As discussed in a new white paper published by NAC, no single solution that can address this complex area of need. Variations in payment models, insurance coverages, healthcare settings, and types of provider challenge innovators to generate flexible solutions. Emerging models of caregiver support show promise that integrating caregivers more fully not only benefits caregivers, but may improve patient outcomes.

READ MORE

A Psychiatrist Shares Tips For Balancing Your Career With Caregiving

When I was 31 years old, my father was

Millions of Americans are abandoning their jobs. The caregiving crisis has pushed me to join them
diagnosed with advanced pancreatic cancer, and my whole world changed. My mother, sister, and I, who all had very busy careers, had to figure out how to balance work with caregiving. Even though we were lucky to be able to share responsibilities, it was so exhausting, I felt like I had jet-lag until long after his 15-month cancer journey had ended.

In my professional life, while it did force me to set boundaries with my time and energy, I was so terrified of doing a sub-par job with my patients and projects, the anxiety kept me up at night. I was showing up every day with a smiling face, excited to help people live healthier lives, but behind the scenes, I felt like I was falling apart.

November is National Family Caregivers Month, a time when we acknowledge issues caregivers face and how to provide support. I interviewed Dr. Anisha Patel-Dunn, who serves as Chief Medical Officer at LifeStance Health, provider of virtual and in-person outpatient mental health care. With nearly 20 years of experience as a practicing psychiatrist and managing a large group practice, she is responsible for overseeing all clinical services delivered through the mental healthcare platform.

I never imagined that I would have to choose between my career and caring for my aging mom. But at 50 years of age, I’m back on the job market, looking for flexible work that will allow me to stay home with her. Without any support, millions of women across America encounter a similar caregiving crisis.

They’re calling it the Great Resignation: The Labor Dept. recently reported that 4.3 million people quit their jobs in August. I was just a few steps ahead of them. I quit mine in July.

While the top reasons people are leaving their jobs are for better pay and job security, many of us are quitting because we have to care for our aging parents at home. I love what I do, but I need to help care for my aging mother. It turns out that given the available options, it was the only workable solution to provide my mom the care she needed.

Women of my generation often face the daunting challenge of finding long-term care for our aging parents. As the years passed, I’ve watched my 81-year-old mother’s physical and cognitive abilities decline and my caregiving responsibilities increase.

Webinar: Elevating Dementia Caregiving as a Public Health Priority

November 19, 2021 | 12:00-1:30 PM

The number of people in the U.S. living with dementia and the family members and friends that care for them continue to rise, yet dementia caregiving remains an emerging public health priority. This webinar presentation will highlight recent, John A. Hartford-funded work on the part of the National Alliance for Caregiving that addresses how and why dementia caregiving is indeed a public health concern. We will conclude with a discussion panel that reflects on how we can ensure that dementia caregiving becomes and remains a public health priority.

Click on the red button below to register for the presentation.
Online Caregiver Party: National Family Caregiver Month

**November 22, 2021 | 11:00 AM-12:25 PM**

Join Family Caregiving Alliance and fellow caregivers for a special party in your honor. November is National Family Caregiver Month and this year FCA will be celebrating with a virtual (online) event. Activities will include dance, games, animals, and some surprises!

**Cost:** No charge  
**Where:** Online (Zoom)

To register, click on the red button below.

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**AHHQI has an open RFP: 2022 Home Health Grant Request for Proposals**

**The Deadline is December 23, 2021**

The Alliance for Home Health Quality and Innovation (AHHQI) seeks to sponsor a research project to help advance home health care and may seek to improve care delivery at home through quality and innovation. Funding will be awarded up to a total of $75,000 maximum (including indirect costs). Multiple projects may be funded if they do not exceed the total funding allocated for all projects of $75,000. Proposals are due by close of business on Thursday, December 23, 2021.

**APPLY HERE**

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**PATIENT AND CAREGIVER STUDIES**

**Black Male Dementia Caregiver Burden Study**

GW School of Medicine and Health Sciences is actively recruiting Black men aged 30-85 who are either caregivers or non-caregivers of loved ones diagnosed with dementia. Participants will engage in a series of questionnaires, surveys, and a focus group, and can receive up to $125 in compensation. Click the link below for additional information.

**LEARN MORE >**

**COVID-19 Study**
The University of Tennessee College of Nursing is seeking adults who are currently providing care (at least 8 hours a week) to an older adult with a chronic condition. Are you a caregiver for an older adult with a chronic condition? Please consider taking part in this important research study to understand caregiving during COVID-19. The University of Tennessee College of Nursing is seeking adults who are currently providing care (at least 8 hours a week) to an older adult with a chronic condition for a research study. The purpose of the study is to understand the experiences of caregivers during COVID-19. The study involves responding to an online survey questionnaire with questions related to caregiving and one interview to talk about their experiences using online caregiving resources. The one-time interview takes approximately 30–45 minutes to complete. Participants will receive a $25 Amazon gift card for completing the interview and a separate $10 Amazon gift card for completing the survey.

LEARN MORE

Intuition Study

Biogen has officially opened enrollment for the virtual Intuition Study in the United States. Using everyday devices, this first-of-its-kind study aims to measure changes in thinking and memory in adults, as well as identify longer-term changes in brain health. For more information about who is eligible and how to enroll, click the link below.

LEARN MORE

IN CASE YOU MISSED IT...

NAC Elevates The Voice of The Nation's Caregivers In a New Video Campaign

NAC has produced a series of vignettes designed to capture the voice of our nation’s caregivers. Included in the RAISE Council’s final recommendations to Congress, the vignettes offer a timely, added layer of social support and recognition for family and friends providing unpaid care.

To help support caregivers, NAC supports the implementation of policies that will reinforce those caring for a diverse range of people across different lifespans and experiencing different health conditions.

Click below to view the video.
Livanta #MyRole Social Media Toolkit

To help improve public awareness of family caregivers and increase family caregivers’ self-identification, Livanta has created #MyRoleCounts, a social media toolkit for partners and stakeholders. The social media toolkit consists of a variety of recommended graphic files, social media posts, and hashtags. Content is written using simple, plain language. Graphics are vibrant, use eye-catching colors, and cultural diversity to engage followers. Items from the toolkit can easily be downloaded below from Livanta’s website or cross-posted on social media.

Click the link below to download and share the kit today!

DOWNLOAD HERE
Lessons from the Workplace: Caregiving During COVID-19

In the first article of its new Spotlight series, Social Innovations in Caregiving, the National Alliance for Caregiving details the hardships the COVID-19 crisis magnified for millions of working caregivers and explores the mismatch between what family caregivers need to meet the demands of working while caregiving and what benefits and supports employers offer. “The pandemic has exposed the reality that working Americans must too often choose between caring for loved ones and holding onto their jobs,” says Grace Whiting, President, and CEO of NAC.

Offering potential solutions on how employers can address this mismatch, NAC advances recommendations for workplace policies and practices that employers can provide to caregivers. These recommendations include flextime, compressed workweeks, and public policy strategies to help reinforce workplace solutions such as redefined paid leave criteria.

READ HERE
NAC's Circle of Care Guidebook for Caregivers of Children with Rare and/or Serious Illnesses

The National Alliance for Caregiving recently produced The Circle of Care Guidebook for Caregivers of Children with Rare and/or Serious Illnesses, resource designed in partnership with Global Genes and with support by Mallinckrodt Pharmaceuticals to provide caregivers with the support, services and specialized information they need to care for a child with a rare and/or serious illness. This guidebook offers an extensive list of resources to help a caregiver in any situation throughout the entirety of their journey as a rare disease caregiver, compiled by those in the rare disease space and caregivers themselves. This list is supplemented by the specialized information needed in order to care for a child with a rare and/or serious illness. If you or someone you know cares for a child living with a rare and/or serious illness, make sure to check out the guidebook at the link below.

You’ll learn about:

- The process of getting an accurate diagnosis for a rare and/or serious illness;
- Genetic testing, clinical trials and support groups that can help;
- Information on treatment and care coordination with specialized teams;
- Understanding the cost of care and treatment;
- Advocating for your child, their care and in their disease space;
- Empowering your child to manage their rare and/or serious illness through all aspects of their life, including when they become an adult; and
- Caring for yourself and your family.

There is also an appendix with a comprehensive list of online resources, supports and services for caregivers, the child living with the rare and/or serious illness, and his or her family that are referenced throughout the guidebook.

Click on the button below to access the Guidebook.

Dental Help for Adults With Disabilities

Special care dentistry (SCD) is a branch of dentistry that facilitates care for impaired patients with physical, intellectual, sensory, mental, emotional, medical, or social disabilities.

Special care dentistry is taught as a postgraduate course that equips dentists to offer dental services to patients with special care needs.

International organizations like the Special Care Dentistry Association (SCDA) are made up of oral dental professionals dedicated to promoting the oral health of persons with special needs.

Those qualified for special-care dentistry are people who need free comprehensive dental treatment. This includes:
- The elderly
- Disabled people
- Mentally and medically compromised persons
Best Practice Caregiving: Infographic Series on Dementia Caregiving Program

The Family Caregiver Alliance (FCA) is publishing a series of blogs and one-page infographics as part of a series about specific dementia caregiving programs that are found in Best Practice Caregiving.

Best Practice Caregiving (bpc.caregiver.org) is a free online database that helps health care and social service organizations identify, compare and adopt best-fit programs for their clientele and community.

Plain Language Resource for Crime Victims With Disabilities

A new fact sheet, created by The Arc's National Center on Criminal Justice and Disability, is designed for people with disabilities to understand their rights if they are victimized. The document outlines different types of abuse, what people with disabilities -- or the caregivers who help them -- can do if they become a crime victim, and where to go for help. Download it now and share it in your community!

What’s Public Health Got To Do With… Family Caregiving?

Trust for America’s Health (TFAH) has published a blog, "What’s Public Health Got To Do With… Family Caregiving?", to launch its new monthly blog series on Age-Friendly Public Health Systems. This series is designed to stir up conversation, generate interest, and challenge healthy aging stakeholders to engage more deeply in age-friendly public health issues.

Author of this month’s blog on "What’s Public Health Got To Do With… Family Caregiving?,” Megan Wolfe, TFAH’s Senior Policy Development Manager, outlines five potential roles for public health departments to support caregivers, organized according to TFAH’s AFPHS 5Cs Framework. The roles include collecting and disseminating data, coordinating existing supports and services, connecting and convening multiple sectors, communicating, and complementing existing supports.

Diverse Family Caregivers Toolkit

Download the Diverse Elders Coalition's Resources for Providers: Meeting the Needs of Diverse Family Caregivers Toolkit. This toolkit offers topline information on what providers need to know, and key pieces from our comprehensive training curriculum, Caring For Those Who Care: Meeting the Needs of Diverse Family Caregivers. Whether you've already attended one or more of our trainings, or this is your first time looking into what's available to help you support diverse family caregivers, we think you'll find these resources to be invaluable in building a more welcoming, supportive practice.
Dementia Care Providers Database

Best Practice Caregiving is a free online database of proven dementia programs for family caregivers. It offers a searchable, interactive, national database of vetted, effective programs that offer much-needed information and support. The database is an invaluable tool for healthcare and community-based organizations, as well as funders and policy makers to discover and share high quality programs for caregivers.

In the Best Practice database you will find detailed information about:

- focus of each program
  - (e.g., reducing stress, understanding dementia, planning care, skill-building, health & wellness, etc.)
- program implementation
- research findings
- direct utilization experiences of delivery sites
- program developer information.

Click [here](#) to learn more about the database or click the button below to access it.

Home Is Where the Care Is

What if you were one of the 2 million adults who are homebound in the United States? Wouldn’t you want care to come to you? With the nation's older adult population increasing, home-based primary care is quickly becoming the future of health care for patients who are medically complex. New resources share needed information about delivering primary care in the home:

- The Better Care Playbook's Home-Based Primary Care Collection provides research, resources and tools for delivering care at home. The collection includes blogs from Rush@Home detailing its care model elements and implementation insights, and the Department of Veterans Affairs highlighting its successful program.

- A Home Centered Care Institute (HCCI) House Calls 101 course offers a simulated educational experience to help you get started with home-based primary care.

Click [here](#) to the online course. Click the button below to browse the collection.

Report: The United States Should Recognize and Support Caregiving Youth

In the United States, more than 5.4 million children and adolescents under age 18 provide care for family members who are aging or have chronic illness, disability, or other health conditions that require assistance.
In this policy study published in the Society for Research in Child Development's Social Policy Report, the authors describe youth’s care for the family and highlight the increasing prevalence, global challenges, and uneven successes of measurement and categorization. They briefly summarize research on how caregiving affects youth’s academic, social, and emotional well-being. Next, they present novel, emerging evidence from the public school-based 2019 Youth Risk Behavior Survey for the State of Florida, which suggests that as many as 24 percent of middle school students and 16 percent of high school students provide at least some care to the family on a regular basis. Drawing on this evidence, the authors’ discuss targeted social programs which have been shown to promote the well-being of caregiving youth outside of the United States, as well as a 13-year-old school-based intervention in The School District of Palm Beach County, Florida.

The report concludes with specific recommendations for a path toward recognizing and supporting caregiving youth via policy and practice in the United States. Its aim is to increase the awareness and feasibility of identifying and supporting caregiving youth and their families via government-organized data collection and targeted social policies.

**2021 Global State of Caring Report**

The International Alliance of Carer Organization’s 2021 Global State of Care report provides an in-depth profile of the issues, approaches, policies, and innovations that are needed to support carers. It profiles the carer policies and practices in 18 countries. This interacting tool features carer initiatives addressing six universal carer priorities:

- Recognition: Legislation and awareness campaigns
- Financial support: Income support, pensions, and benefits
- Work and education: Carer-friendly workplaces and educational initiatives
- Health and well-being: Health and social supports
- Information and knowledge: Resources and education
- Evidence-informed practices: Innovative policies and practices

**The John A. Hartford Foundation's Dissemination Center**

Funded and disseminated by The John A. Hartford Foundation, The American Association of Retired Persons (AARP) has released a new series of how-to videos and accompanying resource guides that walk family caregivers through what to expect before, during, and after a planned or emergency hospital stay.

The videos and resource guides, many of which are available in both English and Spanish, are free of charge and were developed by Home Alone Alliance members—the

The “How-To” videos and resource guides for family caregivers are on specific medical/nursing tasks – including preparing special diets, managing incontinence, wound care, mobility, and managing medications.
Employers play a vital societal role by providing employment, work experience, employee benefits, and the ability for workers to save and invest for a secure retirement. This report examines the pandemic's impact on employers across the company, their response, and timely opportunities. A strong employee benefits package, including retirement benefits, health insurance, workplace wellness programs, and caregiver support, can create a win-win situation for employers and their employees. Especially now, as our nation is emerging from the pandemic, employers need support from policymakers to continue paving the way for their recovery and to make it as easy as possible to enhance their business practices and expand their benefits offerings.