Employee Productivity Increased And Stress Decreased By Family-Caregiver Digital Platform

Approximately 53 million Americans provide unpaid caregiving, according to Caregiving in the U.S. 2020 by the National Alliance for Caregiving (NAC) and AARP. The percentage increased from 18% in 2015 to 21% in 2020, and the rise is projected to continue.

After navigating the hurdles of being the primary caregiver for her mother before she died, Jessica Kim, a serial entrepreneur, thought there had to be a better way. She and her cofounder, Steven Lee, built an online platform and app, ianacare, that provides a free way to coordinate support within your social circle and easy access to benefits personalized to your employer and insurance companies' plans.

From: Forbes | Published: June 8, 2022

Black and Hispanic Caregivers Hard-Hit by Pandemic, Look to Employers for Support

The pandemic amplified health inequities in several American populations, particularly in those underserved by the medical industry, causing both physical and mental health outcomes to worsen for Black and Hispanic individuals who are older and have disabilities, and at the same time, for their caregivers. Employee benefits professionals are in a position to help provide valuable education and resources to these overlooked groups.

I’m a Working Mom and my 4-year-old Son’s Full-time Medical Caregiver. Here’s What Life is Like

It’s 11:17 p.m., and I just sat down to begin my workday.

Thank God I work from home. Tonight, like so many nights over the past four and a half years, “home” is a hospital room.

My workspace is a fold-out bench-to-bed in the far corner of the room, with a direct line-of-sight to my son asleep in his hospital bed.
More than 1 in 5 Americans are caregivers — 17% of them are Hispanic (non-White, non-African-American) and 13% Black.

From: BenefitsPro | Published: June 13, 2022

This is my workspace. At least, it is one of my workspaces.

From: Today Show | Published: June 10, 2022

NOW AVAILABLE: World Carers Conversation 2022 Session Recordings

World Carers Conversation returned May 19, 2022 as a global, virtual event with the support of EMD Serono and Embracing Carers. Innovators, researchers, policymakers and carer advocates from around the globe learned about advancements in the field, got inspired, and made connections that matter for the future of carers everywhere. These recordings cover the primary conversations and topics discussed.

To view the slide deck used in the event, click here. To view the recordings, click below.

Global Voices of Caregiving: A PhotoVoice Project

As part of the World Carers Conversation 2022, NAC worked with Chief Story Officer Anne Levy and caregivers from around the globe to help them tell their own story through a series of photos. These photos addressed simple questions about being a caregiver, such as what works, what doesn’t work, and what a world that embraces and supports caregivers would look like. Each one of these
Reimagining Clinical Trial Recruitment Through a Family-Centered Lens: Caregiver Recommendations for Enhancing Clinical Trial Participation Diversity

The National Alliance for Caregiving is proud to present Reimagining Clinical Trial Recruitment Through a Family-Centered Lens: Caregiver Recommendations for Enhancing Clinical Trial Participation Diversity, a new report developed with sponsorship from Travere Therapeutics. Building on the priority area of clinical trials recruitment and retention identified in NAC’s 2019 report Paving the Path for Family-Centered Design: A National Report on Family Caregiver Roles in Medical Product Development, this report synthesizes outputs from a series of three roundtables. In collaboration with Health Leads, NAC mobilized its network of researchers, policymakers, innovators, patients, families and caregivers to discuss and develop strategies for improving how researchers engage caregivers from populations most impacted by structural inequities in order to better understand how to increase representation of diverse patients in clinical trials.

Included in this report:
NAC’s report underlines that engaging caregivers more effectively in the clinical trials process to promote better trial enrollment diversity requires investment and collaboration from not only industry partners and sponsors, but also from federal regulatory agencies, policymakers, healthcare leaders, community care providers and patients and caregivers themselves. Recognizing the value – the contributions, the expertise, the health impact – of what caregivers do, as partners in innovation, is a key part of creating a more equitable, person and family-centered health care system.

To read the report and access our findings, click on the red button below.

The Circle of Care Guidebook Series

The National Alliance for Caregiving’s Circle of Care library is a series of guidebooks dedicated to providing resources, support and information to caregivers caring for someone in a specific disease space. Part of an ongoing effort of NAC, these first three guidebooks were developed with the assistance of both patient-advocacy and consumer-facing partners:

- Circle of Care: A Guidebook for Mental Health Caregivers
- Circle of Care: A Guidebook for Caregivers of Children and Adolescents Managing Crohn’s Disease
- The Circle of Care Guidebook for Caregivers of Children with Rare and/or Serious Diseases

NAC is proud to lend their expertise on caregiving through these guidebooks. To access the guidebooks for yourself and to share them with other caregivers in your life, please visit the link below.
The Pride of Caring: The Pride of Community

June 16, 2022 | 11:00 AM - 12:00 PM EST | Online

What can LGBTQ patients teach providers about their health care experiences? What should providers know?

Join World Famous *BOB* and Richard Bondi of the Austin (Texas) LGBT Elder Task Force for this webinar hosted by the WellMed Charitable Foundation to discuss the importance of LGBTQ+ community engagement across all ages to support LGBTQ+ seniors and caregivers.

To register for this event, click the red button below.

REGISTER>

Equity Action Plan Stakeholder Engagement Meeting

June 16, 2022 | 1:00 PM - 2:30 PM EST | Online

Join the Social Security Administration for a national meeting to learn about our Equity Action Plan in support of President Biden’s Executive Order 13985 on Advancing Racial Equity and Support for Underserved Communities Through the Federal Government.

The SSA will discuss its comprehensive approach to advancing equity for people who have been historically underserved, marginalized, and adversely affected by persistent poverty and inequality. Discussion topics will include:

- Identifying inequities in Social Security programs and services;
- Identifying and addressing systemic barriers to program participation;
Ensuring equitable service for unrepresented claimants in the disability application process; 
Increasing gender equity and equality in our programs; and 
Increasing equitable access to SSA research grants and procurement opportunities for Historically Black Colleges and Universities, Minority Serving Institutions, and small and disadvantaged businesses.

Please register by Wednesday, June 15, 2022, to learn more about this historic plan. After registering, you will receive a follow-up email with the meeting agenda and Microsoft Teams link. If you have any questions, please contact us at OEA.Net.Post@ssa.gov.

To register for this event, click the red button below.

**REGISTER >**

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**NCCIH Request for Input on Whole Person Health - Determinants**

**Deadline: June 17, 2022**

The National Center for Complementary and Integrative Health (NCCIH) at the National Institutes of Health (NIH) is seeking your organization's input to delineate a set of common data elements that will guide research on whole person health. We have published a request for information (RFI), "Identification of a Set of Determinants for Whole Person Health," and invite you to share it with your organization's members. Comments are due by June 17, 2022.

The purpose of this Request for Information (RFI) is to solicit public comment on defining a set of key determinants of health that addresses all the elements of the whole person health model, i.e., factors that can influence health either positively or negatively, and that encompass the full continuum of biological, behavioral, social, and environmental domains.

To learn more, click [here](#). To submit comments, click the red button below.

**REGISTER >**

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**Age-Friendly Health Systems Oncology Action Community & Informational Webinar**

**June 22, 2022 | 1:00 PM - 2:00 PM EST | Online**

The Association of Community Cancer Centers (ACCC) is holding a webinar June 22 on how to participate in ACCC’s Age-Friendly Health Systems Oncology Action Community.

Throughout the United States, hospitals, clinics, long-term care and other sites of care are becoming Age-Friendly Health Systems as they implement the 4Ms Framework - what Matters, Medication, Mentation and Mobility - to provide high-quality care that meets the
unique needs of older adults. Join this online event to learn about the Age-Friendly Health Systems movement and how to participate in the ACCC’s free, seven-month learning collaborative that will guide oncology clinical teams in implementing the 4Ms.

To register for this webinar, click the red button below.

REGISTER >

The Pride of Caring: The Pride of Keeping Active

June 23, 2022 | 11:00 AM - 12:00 PM EST | Online

Join Bruce Williams, Active Aging Manager at The Pride Center at Equality Park in Fort Lauderdale, Florida as we discuss the importance of active aging for LGBTQ+ seniors through engagement, enlightenment and entertainment!

This webinar will be hosted by the WellMed Charitable Foundation.

To register for this event, click the red button below.

REGISTER >

Webinar: Your Mental Health as Caregiver in the Times of COVID

June 23, 2022 | 1:00 PM - 2:00 PM EST | Online

An estimated 48 million adults in the US, almost 20% of our country provide informal care to an adult with physical or mental health needs. This unpaid work, which includes everything from trips to the doctor to feeding, bathing, and toileting, has been valued at almost 500 Billion dollars. Even in the best of times, the vast majority of this work is invisible and under supported, leaving millions of caregivers struggling in silence. The Covid-19 pandemic pushed many caregivers into crisis.

And while Covid-19 vaccination has helped some Americans experience a joyful summer, a large number of caregivers are still at home, struggling with the fallout from ongoing
This presentation exposes a national failure to support this vital part of our society — one that many of us will depend on at some point in life and what we need to do to support our own mental health in the face of this new Normal using tried and proven straight forward “Getting Real” tools as our only true solution.

To register for this event, click the red button below.

REGISTER

3rd Annual Virtual Caregiver Support Summit

June 30, 2022 | 9:00 AM - 12:00 PM EST | Online

On behalf of the GV Sonny Montgomery VA Medical Center Caregiver Support Program, we invite you to participate in the 2022 Virtual Caregiver Summit. Our goal is to engage with providers, caregivers, family members and, community partners, “To promote the health and well-being of family caregivers who care for our nation’s Veterans, through education, resources, support, and services.”

VA recognizes the crucial role that Caregivers play in helping Veterans recover from injury and illness and in the daily care of Veterans in the community. VA values the sacrifices Caregivers make to help Veterans remain at home. Caregivers are partners in the care of Veterans and VA is dedicated to providing them with the support and services they need.

To register, email Shalonda.Carlisle@va.gov or Ruth.Drake2@va.gov.

For additional information, contact Caregiver Support Program at 601-364-1559.

REGISTER
The Columbia Aging Center and the Columbia Journalism School will host the 2022 virtual Robert N. Butler-Jack Rosenthal Age Boom Academy: "Caregiving and Our Longer Lives – The $500 Billion Question" over four two-hour online sessions on October 13, 14, 20 and 21, 2022 (exact session times at ageboom.columbia.edu). The deadline for journalists to apply is June 30, 2022. Into its third decade, the Age Boom Academy is Columbia University's signature media training program for reporting on the opportunities and challenges of increased longevity.

If you are an on-staff or freelance journalist, you may apply to become a Fellow in this program co-led by Linda P. Fried, MD, MPH, Dean, Columbia Mailman School of Public Health; and Bruce Shapiro, Executive Director, Dart Center for Journalism and Trauma and Columbia Journalism School faculty. Selected Fellows will convene with researchers for an intensive, multi-session, online workshop focused on how to report effectively on the complex interplay of increased longevity and the U.S. informal caregiving economy. This topic crosscuts journalistic beats, regions and markets.

To learn more and register, click on the link below.

**REGISTER**

**JULY**

Peer Support Mentoring Program – Spirituality Calls

The VA Peer Support Mentoring (PSM) Program and the National Chaplain Office will be hosting a 12-month series addressing spirituality and caregiving. These calls are open to caregivers of Veterans enrolled in both the Program of General Caregiver Support Services (PGCSS) and the Program of Comprehensive Assistance for Family Caregivers (PCAFC).

Calls will be held on the first Friday of each month from 12:00 p.m. to 12:30 p.m. ET (11:00 a.m. to 11:30 a.m. CT; 10:00 a.m. to 10:30 a.m. MT; 9:00 a.m. to 9:30 a.m. Pacific) and will be hosted on WebEx using the dial-in information below. There is NO registration required. Meeting Call in number: USA Toll-free Number 1-833-558-0712. Meeting number (access code): 2762 030 9378

Topics

- July 2022 - To be announced
- August 2022 - Spirituality and Grief
- September 2022 - Spirituality and Self Reflection

To learn more, click the red button below.
Spotlight on Caring for the Caregiver

July 13, 2022 | 6:00 PM - 7:30 PM EST | Online

Please join Partners in Care Foundation and AARP for the last session of a 3-part virtual series, "Spotlight on Caring for the Caregiver," on Zoom.

Moderators:
- June Simmons, President & CEO, Partners in Care Foundation
- Adriana Mendoza, Associate State Director, AARP California

To register for this event, click the red button below.

RELAX: Alternatives to Anger for Parents and Caregivers

July 14, 2022 | 1:00 PM - 2:30 PM EST | Online

Hosted by Michigan State University, this interactive workshop for parents and those who provide care for young children is designed to help adults increase their knowledge about anger and strong emotion in young children and the developmental stages of social emotional self-control. Parents and caregivers will learn what anger is and what triggers anger, explore ways to practice and reinforce positive communication skills, and learn methods of incorporating children’s literature into the process of teaching problem solving skills.

Facilitator: Georgina Guzman, perrygeo@msu.edu.

This workshop is free.

Attendees must register by July 13. Click the red button below to do so.
Powerful Tools for Caregivers – Virtual

July 14, 2022 | 7:00 PM - 8:30 PM EST | Online

In conjunction with Catholic Charities Southwestern Ohio, Powerful Tools for Caregivers presents this second meeting in 6-week self-care class for family caregivers where you will access valuable resources, connect with other caregivers, reduce stress, guilt, anger and depression, learn to communicate effectively with family members, doctors, or other help, and learn how to take care of yourself, relax, set goals and problem solve. Uses companion book "The Caregiver Helpbook", available at www.powerfultoolsforcaregivers.org.

To register for this event, click the red button below. Attendees must register by July 7, 2022.

REGISTER >

Essentials of Cancer Caregiving Lifestyle for Cancer Survivors & Caregivers

July 19, 2022 | 3:00 PM - 4:30 PM EST | Online

Join UT Health San Antonio Caring for the Caregiver for a session designed to help attendees learn more about programs to support cancer survivors and the importance of a healthy lifestyle and social support. Dr. Kyungeh An with UT Health will discuss Cancer survivorship and caregiver roles in a discussion on barriers to a sustain healthy lifestyle.

For more information contact 210-450-8862.

To register for this event, click the red button below.

REGISTER >
Apply to Be a Macy Faculty Scholar

Deadline: August 1, 2022

The Macy Faculty Scholars Program, now in its second decade, aims to identify and nurture promising early career educators in medicine and nursing. The Program will help develop the next generation of national leaders in medical and nursing education by identifying outstanding educators, physicians, nurses, and role models—individuals who represent the breadth of diversity seen in learners, patient populations, and health care settings around the country. By providing the Scholars with resources—especially protected time, mentorship, and a professional network of colleagues—the Program aims to accelerate Scholars’ careers, to turn their teaching practice into scholarship, and to help them become impactful leaders locally, nationally, and beyond.

To learn more about the Macy Faculty Scholars Program, click here. To apply, click the red button below.

Strategies to Support Those Touched by Cancer

August 9, 2022 | 2:00 PM - 3:30 PM EST | Online

Join UT Health San Antonio Caring for the Caregiver for a session designed to help caregivers of individuals with cancer navigate several complexities of the healthcare system. Caregivers will receive ways and means to help identify and advocate for the best pain and symptom management for their loved ones. Finally, caregivers will learn strategies to support self-care.

For more information contact 210-450-8862

Event presented by UT Health Caring for the Caregiver and funding supported by
SEPTEMBER

Survivorship and Cancer Care

September 15, 2022 | 11:00 AM - 12:30 PM EST | Online

Join UT Health School of Nursing and the Caring for the Caregiver program for a special session on Survivorship and Cancer Care. During this session, attendees can expect to learn more about life after Cancer. Participate in a discussion regarding the relationship between what we eat, how we move, and how it impacts the cancer survivors' ability to prepare for treatment, tolerate treatment, and recover from treatment. We will go over strategies to support our loved one's symptoms and recovery from cancer treatment and discuss long-term implications.

For more information call 210-450-8862.

To learn more or register for the conference, click the red button below.

PATIENT AND CAREGIVER STUDIES

Are You Caring for a Loved One With Dementia

Researchers at Weill Cornell Medicine are conducting a research study examining well-being and resource use in dementia caregivers. The aim of this study is to identify and examine the needs of caregivers and the types of resources and support services that would be most beneficial to promote well-being in caregivers.

If you are eligible and agree to participate, you will be asked to complete an online survey lasting 20-30 minutes. Then, you will be given the option to participate in a follow-up phone interview with a member of our research staff. You will be asked to answer some questions about your demographic characteristics, care responsibilities, challenges experienced, and ways you have coped with those challenges. Then, you'll be asked to answer some questions about your use of community resources, barriers to accessing and using resources, and suggestions for resources that you would be interested in using in the future based on your own personal needs.

You will receive a stipend of $25 for your completed surveys. For those who choose to participate in the open-ended interview, you will be compensated with another $40.

If you have any questions about this survey, please feel free to email Francesca Falzarano.
Recruiting Participants for a Study on Accessibility Information in Cities

Our research team at the University of Illinois at Chicago is inviting you to participate in a research study to improve the availability and use of data on sidewalk accessibility for people with disabilities. We would like your input to help understand new ways our tools and the data collected can be used to advocate for more accessible cities and be used by people with disabilities to better understand their community’s accessibility.

Participation includes two 2.5 hours workshops on one of these dates: May 5, 6, 9, 11, 19, 20. The workshop will be either at 10:00am to 12:30pm or 1:00pm to 3:30pm.

To maintain COVID-19 safety precautions the first workshop will be held virtually via Zoom. During these workshops our research team will lead group activities and discussions. We will focus on priorities, needs, and concerns related to accessible sidewalks, and hope to gain your valuable perspectives on the design and usability of a tool we developed. We are also interested in discussing the opportunities and needs around data on accessibility of cities.

Participants will receive $50 per session for their participation.

To Participate in this workshop, participants must be:
· 18 years of age or older
· Speak and understand English
· Identify as a member of one of the following groups:

*People with limitations in mobility OR
*People who are Blind/low vision OR
*A caregiver of a person with a disability
Have experience with or regularly leave home and travel in the community OR travel with someone with a disability

For more information or to schedule a time to participate in the study please contact:

Sierra Berquist
Research Assistant
Email: sberqu2@uic.edu
Study: In Caregiving, These Mindfulness Practices Could Help

Family members who care for a loved one with dementia often experience chronic stress, anxiety and depression. Psychological research into mindfulness remains a fairly nascent field of study, but mindfulness practices have numerous benefits for care partners.

Research conducted by Felipe Jain, Director of Health Aging Studies at Mass General Hospital, has created and established the benefit of a mindfulness and guided imagery approach, Mentalizing Imagery Therapy (MIT), for family dementia caregivers.

The focus of Dr. Jain's research is now to identify strategies to deliver caregiver skills training and relaxation techniques remotely to improve caregiver well-being. The research has been funded by the National Institute on Aging and private foundations.

To learn more about how mindfulness can benefit caregivers, click here. To sign up for an upcoming Mentalizing Imagery Therapy study, click the red button below.

SIGN UP

Mason CARES

George Mason University Department of Social Work invites care partners caring for a loved one with dementia to participate in Mason CARES, a research study assessing the effect of an evidenced-based virtual education program on care partner stress-management. This program serves to reduce care partner stress/burden and increase care partner well-being.

Family care partners provide more than 80% of the daily care of older adults in the U.S. In 2019, more than 16.1 million unpaid family care partners of people with dementia assumed caregiving tasks because of declining cognitive and physical functioning among older adults (Alzheimer's Association, n.d). Non-pharmacological, affordable interventions have evidence of effectiveness for families challenged with dementia in reducing care partner stress and increasing well-being.
Care partners in our research study will enroll in a virtual community of practice, receiving support from the George Mason University Mason CARES team through the 9-week evidence-based Stress-Busting Program (SBP) for Family Caregivers™, with a community of other care partners. We will collect information regarding stress and well-being before, during, and after the study ends. Upon completion of the SBP, the team will follow up with continued support for one month. Time commitment will be no more than 30 – 90 minutes per week depending on weekly activities.

To learn more about the Mason CARES study, click here. To sign up, click the red button below.

IN CASE YOU MISSED IT...

Providing Support When You Need Support: Caregiving While Depressed

As a caregiver, there are resources that can support you, your loved one, and your family as you navigate your path toward wellness. Watch our panel and hear from caregivers with firsthand experience!

RESEARCH & RESOURCES

Making the Invisible Visible: State Strategies for Identifying and Reaching Family Caregivers
Family caregivers — often relatives, friends, and neighbors — provide vital assistance to support the health and functional needs of older adults, but often face physical, emotional, and financial challenges related to their caregiving responsibilities.

Many states are seeking to better support this important workforce, particularly family caregivers in communities of color, but need an effective way to accurately identify the number, diversity, and unique needs of family caregivers in their state.

In identifying family caregivers, states should recognize cultural considerations, including distinct racial and ethnic perspectives associated with caregiving, that may have implications for identification.

This brief by the Center for Health Care Strategies presents key strategies to help states better identify and reach family caregivers.

To view this brief, click on the red button below.

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**PREVIOUSLY APPEARED**

**AARP/National Geographic**

"Second Half of Life Study"

Older Americans are not only the happiest adult Americans as a whole, but also consider themselves healthier and more financially secure than those in their 40s and 50s, reveals a new study published in *AARP Bulletin*. The "Second Half of Life Study," conducted with *National Geographic*, surveyed thousands of adults age 18 to 90 to explore how Americans perceive their current life, their expectations for the future and aging in general as we emerge from the COVID pandemic. About two-thirds of adults over 80 said that they were living their "best possible life" or close to it, compared with just 1 in 5 younger adults.

The study disproves many common misconceptions about aging and suggests that greater focus should be put on adults in their 40s and 50s, who reported higher levels of stress and worry and lower levels of life satisfaction and health than older Americans.

To view the study, click on the red button below.
The engAGED Social Engagement Innovations Hub houses best and emerging social engagement programs. Searchable by a variety of filters—including social engagement intervention type, geographic area served or partners involved—the Innovations Hub gives you access to replicable examples to inspire and inform your organization’s social engagement efforts. Each program summary within the Innovations Hub contains all you need to know about that particular program, such as partners involved, outcomes demonstrated, lessons learned, resources needed and contact information to learn more.

To access the Hub, click the red button below.

ACCESS THE HUB >

Improving Representation in Clinical Trials and Research: Building Research Equity for Women and Underrepresented Groups

The United States has long made substantial investments in clinical research with the goal of improving the health and well-being of our nation. There is no doubt that these investments have contributed significantly to treating and preventing disease and extending human life. Nevertheless, clinical research faces a critical shortcoming. Currently, large swaths of the U.S. population, and those that often face the greatest health challenges, are less able to benefit from these discoveries because they are not adequately represented in clinical research studies. While progress has been made with representation of white women in clinical trials and clinical research, there has been little progress in the last three decades to increase participation of racial and ethnic minority population groups. This underrepresentation is compounding health disparities, with serious consequences for underrepresented groups and for the nation.

At the request of Congress, Improving Representation in Clinical Trials and Research: Building Research Equity for Women and Underrepresented Groups, identifies policies, procedures, programs, or projects aimed at increasing the inclusion of these groups in clinical research and the specific strategies used by those conducting
clinical trials and clinical and translational research to improve diversity and inclusion. This report models the potential economic benefits of full inclusion of men, women, and racial and ethnic groups in clinical research and highlights new programs and interventions in medical centers and other clinical settings designed to increase participation.

To learn more or to access the report, click the red button below.

**LEARN MORE >**

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**How to Get Help from Social Security**

Local Social Security offices are offering more in-person appointments and have resumed in-person service for people without an appointment.

As in-person service expands, the Social Security Administration expects its offices to become increasingly busy. The SSA strongly encourages beneficiaries seeking assistance to continue to go online, call for help, and schedule appointments in advance.

The Social Security Administration has provided a fact sheet for those seeking assistance. To read the fact sheet, click the red button below.

**LEARN MORE >**

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**Home Safety Checklist for Alzheimer's Disease**

Use the following room-by-room checklist provided by the National Institute on Aging to alert you to potential hazards and to record any changes you need to make to help keep a person with Alzheimer's disease safe. You can buy products or gadgets necessary for home safety at stores carrying hardware, electronics, medical supplies, and children's items.

To view the checklist, click the red button below.

**VIEW THE CHECKLIST >**
At a Glance: Reducing Disability in Alzheimer’s Disease

Check out this 1-page infographic for a quick, top-level overview of Reducing Disability in Alzheimer’s Disease (RDAD), a dementia caregiving program that offers 12 in-person (individual or group) exercise training and education sessions for caregivers and persons living with dementia. It’s focused on gentle exercise for the person with dementia and caregiver training to manage behavioral symptoms and identify pleasant events.

Feel free to print it out, save for reference or send to a peer or colleague who may be interested learning more about the program. You can find full information, including program components and characteristics, research evidence, sustainability strategies and more at the program’s full profile page at Best Practice Caregiving.

To view the infographic, click the red button below.

A Caregiver Program that Creates Lasting Improvements

BRI Care Consultation™ is a proven care-coaching solution delivered by telephone and email to adults with health conditions and their family or friend caregivers, empowering both to more effectively manage short- and long-term needs.

BRI Care Consultation uses a combination of ongoing assessment, action planning and follow-up support to create real and lasting changes in the caregiving situation, resulting in a decrease in unmet needs, stress and strain, and symptoms of depression for both caregivers and the person receiving care.

BRI Care Consultation helps to change the cycle of ongoing care, uncovering alternatives to current care methods, identifying helpful new community resources, mobilizing family and friend assistance, and driving lasting changes in the care situation.

Through numerous research studies, BRI Care Consultation has been proven to provide positive outcomes, including:

- Improved satisfaction with care from doctors and other professionals
- Reduced symptoms of depression
- Less relationship strain between the caregiver and person receiving care
- Fewer costly emergency room visits and hospital re-admissions.

If you’d like more information, contact Michelle Palmer at mpalmer@benrose.org, or click the red button below.
New Blood Test Can Help Diagnose Alzheimer’s Disease

Alzheimer’s disease is characterized by the buildup of a protein called beta-amyloid, which forms sticky plaques on the brain and can cause brain cells to die. Testing for the presence of these amyloid plaques on the brain is an important part of Alzheimer’s diagnosis and research.

A study, funded in part by NIA, found that a new blood test can accurately predict the presence of beta-amyloid in the brain. The blood test became even more accurate when the research team took into account the version of APOE (a gene linked to Alzheimer’s risk) that each person had. Scientists note that the blood test performs comparably to existing brain scan- or spinal tap-based tests. However, the blood samples used in the study were from majority white, affluent individuals, and may not be generalizable to other demographic groups.

Using blood samples will make it easier to screen healthy people for potential enrollment in Alzheimer’s clinical trials and could help lower costs and expand the availability of diagnostic studies for Alzheimer’s.

To learn more, click the red button below.

Caregiving While Black Course

Caregiving While Black is a six week culturally appropriate caregiving training and education course. The overall goal of this project is to address the cultural reality of “Caregiving while Black” during a public health crisis. We are aiming to develop and prototype-test a highly accessible program designed to enhance the mastery of Black American caregivers to provide care to family members or friends living with a dementia illness in a time of crisis.

A $40 gift card will be provided for participation in each baseline and post-intervention interview and each semi-structured interview.

If you’d like more information, click here. If you’d like to participate, click the red button below.
Free Course on Diversity, Equity, and Inclusion for Home-Based Care

During National Minority Health Month, the Home Centered Care Institute (HCCI) has launched a new free course entitled, “Diversity, Equity, and Inclusion for Home-Based Care.”

This on-demand course, supported (in part) by the Josiah Macy Jr. Foundation, addresses the complex and multifaceted topic of health equity by discussing power and privilege, implicit and explicit bias, inferred cultural preferences, and more, which are especially relevant for providers working with patients and families in their home.

The course, designated for a maximum of 0.5 AMA PRA Category 1 Credit ™, provides guidance in developing and implementing best practices for culturally sensitive care in the home environment and within the workplace. While the course targets home-based primary care (HBPC) providers and those interested in HBPC, non-clinical staff members can also benefit from understanding implicit bias as they support interactions with patients and caregivers (such as scheduling or billing), while those in leadership will learn how to better engage and retain high-performing, diverse teams.

To take the course, click the red button below.

LEARN MORE

Caregiver TLC

What is Caregiver TLC?

The Caregiver TLC Research Program offers FREE ONLINE support to adults providing care to persons with memory loss, dementia, or chronic illness.

Caregivers will complete six weekly Zoom sessions led by trained facilitators. This workshop will teach coping skills to deal with stress, depression and burden, and strategies to improve quality of life.

Is Caregiver TLC For Me?

- Do you provide at least 4 hours of care a week to a person with a chronic illness or memory loss?
Do you help your family member to:
- Remember appointments?
- Get bathed or dressed?
- Take their medications on time?
- Could you attend six weekly 2-hour Zoom* sessions?
- Would you like to connect with other caregivers near you?
- Do you have a personal email address?

To register for Caregiver TLC, click here. To learn more, click the red button below.

LEARN MORE

Older Adults and Healthy Aging (podcast)

Older-adult health has been increasingly on the radar of public health agencies over the course of the COVID-19 pandemic. For many, it prompted new or enhanced partnerships with their sibling agencies. As the population of older adults continues to grow, it is crucial for public health agencies to join partners in the aging network to create systems that support healthy aging.

In this podcast episode, Jewel Mullen (alumni-CT) offers perspectives on the value of cultivating partnerships between state public health agencies and state units on aging. In addition, Elizabeth Head (Deputy Director, Injury Prevention Section, Georgia Department of Public Health) and Aline Stone (Georgia Alzheimer’s and Related Dementias Coordinator, Division of Aging Services, Georgia Department of Human Services) describe their collaboration to improve the health of older adults in Georgia. They share their experience participating in a pilot project with ASTHO and Trust for America’s Health to cross reference their state health improvement and aging plans.

To listen to the podcast, click the red button below.

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Frequently Asked Questions About Caregiving

Caregiving can be overwhelming for anyone. If you’re new to caregiving, take a deep breath, then tackle one task at a time. Consider these tips when getting started:

- Assess your loved one’s needs. Figure out what types of help the person you are caring for needs.
- Ask family members and friends to share caregiving tasks. Splitting up tasks among different people can help both the person needing care and the primary caregiver.
- Look for resources available in your community. Local resources may help you find in-home help, transportation, and home-modification services.
- Learn about different options for paying for care. The person needing care may qualify for certain financial options.
To learn more, click the red button below.