Evolving into New Ways of Living our Mission
Multiple sclerosis (MS) is one of the most common neurological disorders diagnosed in young adults. Many experts estimate that 2.5 million people worldwide have MS. Nearly one million individuals are living with MS in the United States.

MS is an autoimmune disease of the central nervous system, affecting the protective covering (myelin) that surrounds the nerves of the brain and spinal cord. The nerves (axons) may also become damaged. MS is not contagious and researchers continue to seek a cause and a cure.

Individuals with MS may experience one or more of a variety of symptoms, such as visual changes, fatigue, numbness, bladder and bowel problems, depression, cognitive issues, and changes in mobility. Initially, most individuals with MS experience periods of symptom flare-ups (also referred to as relapses or exacerbations), followed by periods of remission. A smaller population of individuals begin with a progressive form of MS, characterized by a slow but steady accumulation of symptoms. More than 20 disease-modifying therapies (DMTs) have been approved by the United States Food and Drug Administration (FDA) to slow disease activity.

The Multiple Sclerosis Association of America (MSAA) was founded in 1970 and is a leading resource for the entire MS community, Improving Lives Today through vital services and support. MSAA provides free, ongoing support and direct services to individuals with MS, their families, and their care partners.

Overseeing MSAA's activities is a national Board of Directors comprised of accomplished professionals from across the country, volunteering their time for MSAA. Providing medical consultation is MSAA's Chief Medical Officer who reviews all of MSAA's medical information. MSAA's Healthcare Advisory Council (HAC) is comprised of leading members of the MS healthcare professional community and works in collaboration with MSAA's Board of Directors and staff to advance and promote the organization's mission and goals. MSAA's African American Advisory Board and Latinx Advisory Board are comprised of leading MS healthcare professionals, people with MS, and care partners. The Advisory Boards identify unmet needs and focus on developing comprehensive and innovative programmatic and educational initiatives to address those needs.

MSAA is a long-standing member of America’s Charities and has achieved Platinum Level status with GuideStar. Additionally, MSAA is a founding member of the Multiple Sclerosis Coalition (MSC), an affiliation of independent MS organizations dedicated to the enhancement of the quality of life for all those affected by MS.
March is always a very active time for MSAA while recognizing this month as MS Awareness Month. During the 2020/2021 fiscal year, our awareness campaign theme, as mentioned earlier, focused on Improving Mental Health and Wellness. Specific programs addressed Purpose in Life, depression and anxiety in MS, care partnering, and wellness strategies to improve symptom management and overall quality of life.

Moving on to our strategic plan covering 2021, 2022, and into 2023, we incorporated a wide range of voices, including those within the MS and healthcare communities. These opinions and ideas were critical in creating our strategic plan and continue to be vital as we seek to plan for the future.

As we look ahead, we will continue to focus on providing innovative programs, timely information, and educational resources. At the same time, we remain committed to adapting to best serve the needs of those with MS and their care partners.

We extend our sincerest gratitude to all who continue to support MSAA in our mission to Improve Lives Today for the MS community.

“Thank you. During this unprecedented time in our lives, it is nice to know we have an organization that is still there for us no matter what.”

— Jonathan from New York
MSAA’s Programs and Services

Assistive Equipment Distribution
More than 700 Items Provided
MSAA provides safety and mobility equipment at no charge and ships directly to the client. Items distributed include a variety of products such as shower chairs and grab bars, as well as mobility devices including canes, walkers, and wheelchairs.

Cooling Equipment Distribution
Nearly 2,000 Cooling Items Provided
MSAA offers cooling vests and accessories at no charge and ships them directly to both adults and children diagnosed with MS. Individuals with MS using this type of technology may experience less fatigue and a reduction in other debilitating symptoms, allowing them to perform better in warm environments and take part in outdoor activities.

My MS Manager™
More than 46,600 Cumulative Downloads by End of Fiscal Year
MSAA’s mobile phone application, My MS Manager, is provided free of charge to individuals with MS or their care partner to use on their iPhone, iPad, iPod touch, or Android mobile phone device. It allows people to track disease activity, store information, generate reports, and securely share information with their healthcare team.

MRI Access Program
Nearly 1,000 Individuals Received Financial Support for MRI Scans
MSAA’s magnetic resonance imaging (MRI) Access Program assists individuals who are uninsured, under-insured, or financially unable to pay for this important exam. MRI scans are vital to the diagnosis of MS, as well as the follow-up evaluation to track the progression of the disease and the effectiveness of long-term treatments.

Helpline
Nearly 16,000 Inquiries Made to MSAA’s Helpline
Our Helpline allows individuals with MS, family members, care partners, and friends to connect directly with one of MSAA’s experienced Client Services Specialists – via phone, email, or our online Chat feature – to receive valuable information, guidance, and support.

MS Conversations Blog
38,759 Visits to Our Blog
MSAA’s blog features timely, interactive discussions on topics important to the entire MS community, from symptom management to the impact of MS on everyday life.

My MSAA Community
More than 6,600 Cumulative Members by End of Fiscal Year
This peer-to-peer online forum welcomes individuals with MS, their families, and their care partners to share information and their experiences with multiple sclerosis.

“I am so very grateful for the financial assistance that I received for my MRI. It made a tremendous difference in my outlook. Knowing that others cared enough about me to offer financial assistance was a gift that lifted my spirits while lifting the financial burden that I was facing. Saying thank you seems inadequate to express my sincere gratitude, but I simply must say it. Thank you!”

— Lora from Georgia
In 2010, I was diagnosed with multiple sclerosis, my second autoimmune illness while 6 months pregnant. As an artist and Visual Arts teacher, I immediately noticed that I could not see the color red or the saturation of color from my right eye. I foolishly thought that At that time, not only was my right eye affected but I also had little control of my right hand which had small tremor like movements. With treatment my vision was restored but illustrating again was a tougher journey. By re-directing my focus, I learned to live in the now. With time, I pushed myself to create art on a higher technical level, yet with deeper meaning for me. I have had unexpected flare-ups in disease activity. Art will always be my guiding light through the darkness that often fills my vision when I relapse. It enables me to cleanse myself spiritually from the dark thoughts and emotions that are often overlooked by people with MS.

Omaya Rivera-Filardi

MSAA’s Website at mymsaa.org

Nearly 800,000 Sessions During this Fiscal Year

MSAA’s comprehensive website provides a wealth of targeted information addressing all aspects of the disease, including experimental and approved long-term treatments, symptom-management strategies, and resources to help individuals adjust to the illness. Visitors to mymsaa.org may access a number of specialized sections as well as educational videos and webinars, publications, and research articles, plus details on MSAA’s vital programs and ways to support MSAA.

MSi Videos

More than 235,000 Cumulative Views* of 96 On-Demand Video Programs

Updating viewers on the latest advances in MS research, disease and symptom management, wellness strategies, and more, this growing library of on-demand video programming and archived webinars serves as a vital resource for the entire MS community.

* Based on videos and webinars currently published on MSAA’s website.

Publications

MSAA publishes: a national magazine, The Motivator, twice per year; My MSAA Today e-newsletter on a bimonthly basis; and other publications focused on areas such as relapse management. Additionally, MSAA’s “What’s New in MS Research” online article series is published bimonthly and features the latest findings in MS treatments, disease development and progression, plus other vital topics.

MSAA Podcast

2,357 Total Downloads of MSAA’s Podcast Episodes
MSAA has long been committed to addressing health inequities for MS communities of color. Listed below are updates on several important initiatives:

MSAA’s African American Advisory Board is comprised of leading healthcare professionals, individuals with MS, and care partners from across the country. The Advisory Board was formed to help MSAA develop programming and initiatives for the African American MS community as well as education for neurologists specializing in MS, addressing the specific issues that face the African American MS community. Since its establishment, the African American Advisory Board has produced a comprehensive publication and patient educational programs as well as a series of educational programs for healthcare professionals titled, All Roads Lead to Change.

MSAA has continued to develop bilingual educational programs focusing on the Hispanic MS experience and in fiscal year 2021, MSAA initiated plans to launch our Hispanic/Latinx Advisory Board. Preparations began for upcoming meetings featuring experts in MS care. Outcomes learned at these meetings will help to guide future strategies and programs.

MSAA conducted a series of national patient educational programs specifically developed for the African American and Latinx MS community, featuring African American and Latinx MS neurologist presenters.

MSAA conducted various webinars and provided on-demand videos throughout the fiscal year for underserved populations, including:

- The Hispanic American MS Experience
  Experiencia Hispanoamericana sobre Esclerosis Múltiple
- How to Navigate the COVID-19 Pandemic as a Person of Color Living with MS
- All Roads Lead to Change: Addressing Unmet Needs in the African American MS Community (provider training webcasts):
  • Providers as Champions of Research
  • Transition of the Adolescent
  • Part I: The Importance of Cultural Sensitivity & Early Diagnosis
  • Part II: The Importance of Cultural Sensitivity & Communication
- Brain Power: Maintaining a Healthy Mind While Living with MS (presented entirely in Spanish) El Poder del Cerebro: Mantener una Mente Sana Mientras se Vive con EM
- New Directions: Making Sense of “Me” and My MS Diagnosis (highlights how MS affects different racial and ethnic communities)

COVID-19 Initiatives

MSAA has responded to the COVID-19 pandemic in several ways to best serve the MS community. Through an informational “hub” on our website, we link the MS community to vital webinars, useful articles, important resources, and our unique COVID-19 and MS Pathfinder online tool.

COVID-19 Webinars

Beginning at the start of the COVID-19 pandemic, MSAA launched a series of webinars titled, “What You Need to Know about COVID-19 and MS,” featuring our Chief Medical Officer Dr. Barry Hendin and the Chair of MSAA’s Healthcare Advisory Council Dr. Carrie Hersh. The webinars continued throughout FY20 to provide the MS community with important updates on COVID-19.

MSAA hosted a series of patient education programs on the topics of the experience and impact of COVID-19 on the African American community.

“This thank you for these wonderful opportunities to educate ourselves about diseases affecting MS. Excellent presentation.”
— Jacqueline G.

COVID-19 and MS Pathfinder

MSAA launched the COVID-19 and MS Pathfinder, a digital and innovative tool created to assist the MS community with accessing vital information. Developed in partnership with Wondros, the COVID-19 and MS Pathfinder tool is designed to enable easy access to important questions. This digital tool features up-to-date information about COVID-19 specifically tailored to the MS community and updated on a regular basis.

“This is actually a really awesome tool. I was “exploring it” last night and it leaves everything out so simply. Definitely check out this tool.”
— Via Twitter re: MSAA’s COVID-19 and MS Pathfinder Tool

Additional COVID-19 Program Initiatives

MSAA continued to offer expanded Helpline hours to meet the increased needs of callers from across the country, plus the addition of a COVID-19 economic hardship waiver for three of our programs: Equipment Distribution, Cooling Distribution, and the MRI Access Program. This waiver enables individuals who would otherwise not meet annual income guidelines, but who are experiencing financial crisis due to the pandemic, to qualify for these programs.
### Awards and Recognition

- **Silver Award** from the National Health Information Awards for MSAA’s publication, *Multiple Sclerosis and the African American Experience*
- **APEX Grand Award** for COVID-19 and MS Pathfinder digital tool
- **Bronze Award** from the Digital Health Awards for the Winter/Spring 2020 edition of *The Motivator* (edition features the cover story, “Coping with the Emotional, Physical, and Mental Effects of a Pandemic”)

“Geeked... zippy... elated... jaunty – you pick! My copy of MSAA’s *The Motivator* came in the mail today! Now that’s a good sign that things are looking up. It’s the little things.”

— My MSAA Community member

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### Financial Performance

#### MULTIPLE SCLEROSIS ASSOCIATION OF AMERICA, INC.

**CONSOLIDATED STATEMENT OF ACTIVITIES FOR THE YEAR ENDED June 30, 2021**

#### ASSETS

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current Assets</td>
<td>$2,204,792</td>
</tr>
<tr>
<td>Property Plant and Equipment (net)</td>
<td>$1,210,063</td>
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<tr>
<td>Other</td>
<td>$4,508,053</td>
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<tr>
<td><strong>Total Assets</strong></td>
<td><strong>$7,922,908</strong></td>
</tr>
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</table>

#### LIABILITIES AND NET ASSETS

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Current Liabilities</td>
<td>$470,612</td>
</tr>
<tr>
<td>Long Term Liabilities</td>
<td>$879,923</td>
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<tr>
<td><strong>Total Liabilities</strong></td>
<td><strong>$1,350,535</strong></td>
</tr>
<tr>
<td>Unrestricted</td>
<td>$5,454,259</td>
</tr>
<tr>
<td>Temporarily Restricted</td>
<td>$1,118,114</td>
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<tr>
<td><strong>Total Net Assets</strong></td>
<td><strong>$6,572,373</strong></td>
</tr>
<tr>
<td><strong>Total Liabilities &amp; Net Assets</strong></td>
<td><strong>$7,922,908</strong></td>
</tr>
</tbody>
</table>

#### REVENUE AND SUPPORT

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Contributions</td>
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<tr>
<td>Rental Income</td>
<td>$120,374</td>
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<tr>
<td>Other</td>
<td>$1,660,980</td>
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<tr>
<td><strong>Total Revenue and Support</strong></td>
<td><strong>$9,802,576</strong></td>
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</table>

#### PROGRAM EXPENSES

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
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</thead>
<tbody>
<tr>
<td>Program Services: Patient services</td>
<td>$4,501,534</td>
</tr>
<tr>
<td>Program Services: Housing</td>
<td>$133,010</td>
</tr>
<tr>
<td><strong>Total Program Expenses</strong></td>
<td><strong>$4,634,544</strong></td>
</tr>
</tbody>
</table>

#### SUPPORTING SERVICES EXPENSES

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>General and administrative</td>
<td>$1,122,119</td>
</tr>
<tr>
<td>Fundraising</td>
<td>$2,452,030</td>
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<tr>
<td><strong>Total Supporting Services Expenses</strong></td>
<td><strong>$3,574,149</strong></td>
</tr>
<tr>
<td><strong>Total Expenses</strong></td>
<td><strong>$8,208,693</strong></td>
</tr>
<tr>
<td>Change in Net Assets</td>
<td>$987,728</td>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>Description</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>NET ASSETS, BEGINNING OF YEAR</td>
<td>$5,584,645</td>
</tr>
<tr>
<td>NET ASSETS, END OF YEAR</td>
<td>$6,572,373</td>
</tr>
</tbody>
</table>
**Fundraising Events**

**MSAA Pivots in a New Virtual Age of Events**

Between continued pandemic-related concerns, and the need to engage a large nationwide audience, MSAA has found a silver lining through a dedicated focus on virtual event planning leading to new growth in event income overall.

MSAA was excited to kick-off 2021 with the launch of a new event series, Improving Lives Through Art, which incorporated elements of the long-standing MSAA Art Showcase. The initial series included two paint-along events featuring an artist with MS, and provided participants with the necessary materials to learn step-by-step how to recreate a piece of art in a safe environment while connecting with the artist, event host, and participants around the country. In addition, the series featured an art lecture focused on artists throughout history who incorporated art as a coping mechanism for an array of challenges. After a successful first season connecting more than 280 participants and raising close to $60,000, MSAA launched a fall season series as well, with two additional series planned for 2022! Learn more about upcoming Improving Lives Through Art events by visiting mymsaa.org.

Hannah Garrison, MSAA Art Showcase participant led members of the MS community through two virtual paint-along events along with host and art enthusiast, Joe Caliva.

Hannah Garrison, MSAA Art Showcase participant led members of the MS community through two virtual paint-along events along with host and art enthusiast, Joe Caliva.

We couldn’t let another year pass without an Improving Lives Benefit, so we worked to launch the very first “Together at Home” virtual Improving Lives Benefit on May 13th. Hosted by Tyler Campbell, the event featured unique content highlighting client stories, the impact of programs such as the MSAA MRI Access Program, as well as our Cooling and Equipment programs. The event highlighted our Corporate Honoree, Medscape, and our Mission Honoree, The MS Foundation of Puerto Rico. Involving more than 300 participants across the country, the event raised more than $188,000 – a 40% increase over the largest Improving Lives Benefit to date and will become an annual tradition even when it is safe to connect in person again. Check out last year’s Benefit by visiting: https://youtu.be/mjpvHE9OGTQ

**MSAA New Donor-Friendly DIY Platform – A Continued Success**

Last year, MSAA launched our new DIY platform to make fundraising even easier for the individuals and groups around the country who wanted to raise awareness and funds in support of MSAA’s mission. In 2021, the Do-It-Yourself platform resulted in almost $100,000 in donations in support of MSAA.

Here are some of the key fundraisers hosted for MSAA in 2021:

• **Swim for MS** is MSAA’s signature fundraiser – You can swim any pool anytime – or in our friend Richard “Dix’s” case, any lake anytime! Dix swam three lakes in California over a weekend in July, beginning his fundraising efforts in June and raised more than $12,000. If you’d like to swim for MS, you can sign-up for free by visiting: engage.mymsaa.org/DIYSwim

Tyler Campbell, motivational speaker and member of the MS community, hosted MSAA's first virtual Improving Lives Benefit.
• Dine & Donate – MSAA’s supporter Sheryl connected with a local restaurant in her area and hosted a multi-day fundraiser. The restaurant donated a portion of sales to MSAA. In addition, they had a tricky tray auction and 50/50 raffle, raising more than $4,000 for the MS community.

• Golf for MS – The Cobler Group Realty in St. Louis hosted their first Golf MS Invitational. They sold sponsorships and foursomes and invited their clients, friends, and community members to join them. You can set-up a golf tournament in your area – it is a great way to network with other local businesses and raise much-needed funds for the MS community.

• Run for MS – One of MSAA’s best fundraising partners is Runs for a Purpose. Every March they host a virtual 1k, 5k, 10k, or half marathon during MS Awareness Month. In 2021 they raised more than $32,000.

If you’re interested in participating in one of the existing fundraisers, or want to start your own, please check out mymsaa.org/DIY

We thank all those associated with the following fundraising events, providing greatly needed support to MSAA and the MS community!

<table>
<thead>
<tr>
<th>Event</th>
<th>Amount</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gary Wallace Golf 2021</td>
<td>$30,817.03</td>
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<tr>
<td>Sunnybrook Golf 2021</td>
<td>$29,175</td>
</tr>
<tr>
<td>Ridgewood Golf 2021</td>
<td>$168,697.63</td>
</tr>
<tr>
<td>Anniversary Challenge 2021</td>
<td>$23,326</td>
</tr>
</tbody>
</table>

235k+ cumulative views of MSAA’s 96 on-demand video programs
Nearby 16,000 inquiries to MSAA’s Helpline, providing needed reassurance and support
Nearly 3,000 pieces of safety, mobility, and symptom-management equipment were distributed
More than 6,600 cumulative members on My MSAA Community, MSAA’s online peer-to-peer forum
Nearly 1,000 people received a vital MRI scan to diagnose or monitor their MS
46,000+ total downloads of MSAA’s mobile app, My MS Manager™
2,357 total downloads of episodes of the MSAA Podcast

Award-winning publications, including a national magazine

Figures shown are for the 2020-2021 fiscal year.

(800) 532–7667 • msaa@mymsaa.org • mymsaa.org

Improving Lives Today
in Many Ways!

WHO GETS MS?

Nearly 1 million Americans and 2.5 million people worldwide have MS

People are most frequently diagnosed between the ages of 15 and 50

Women are 3X more likely than men to develop MS
## Our Generous SUPPORTERS

### CHAMPIONS ($100,000 and above)
- Alan Lionel Fried Trust
- Biogen
- Bristol-Myers Squibb Foundation
- EMD Serono, Inc.
- Genentech
- Johnson & Johnson
- Novartis Pharmaceuticals Corp.
- Sanofi US Services Inc.

### VISIONARIES ($50,000 to $99,999)
- America’s Charities Distribution Account
- Cameron and Jane Baird Foundation
- Impact Education

### MOTIVATORS ($10,000-$49,999)
- Annette Cadwell
- At Point of Care
- Bernard Berkowitz
- Butler/Till Media Services, Inc.
- Celgene
- Commonwealth Insurance Advantage, LLC
- Dianne Boerger
- Discovery Communications, Llc
- Estate of William Crimbring
- Facebook, Inc.
- Greenwich Biosciences
- Healthline
- HelenCarlos
- IBM Employee Services Center
- James Warren
- Mark and Muriel Wexler Foundation
- Mylan Pharmaceuticals
- New Imagitas, Inc.
- Patient Access Network Foundation
- Publicis Health
- Rita Kernen
- Robert Keeley
- Runs for a Purpose
- Santo Petroleum
- The Benevity Community Impact Fund
- The Foster Family Private Foundation, Inc.
- The Piraino/McWilliams Family Foundation
- Virginia T. Dashiel Charitable Foundation
- WarrenKaplan
- WebMD, LLC

### ADDITIONAL DONOR LISTING

#### $5,000 - $9,999
- A + E Networks
- ALC, Inc.
- AmazonSmile Foundation
- American Endowment Foundation
- Barry Black
- Kathi Blair
- David Bock
- Alan Brattesani
- Fred Brunke
- Suzanne Cabaniss
- Cadent Network
- Giuseppe Cecchi
- Richard Chadakoff
- Mike Chevalier
- Cmrp, Inc
- Data Centrum Communications, Inc
- Disney Worldwide Services, Inc.
- Estate of Duane F. Jelinek
- Estate of Florence Rasmussen
- Estate of Jane Breininger
- Estate of Richard Zerbe
- Fiducialy Trust
- William Fleming
- Frontstream, Inc
- Michael Geary
- Genuine Parts Company
- David Herzog
- Kerrin Holsteen
- Deborah Horwitz
- Hulu
- Hyatt
- Intl. Union Of Operating Engineers Local 825
- George Kahler
- Theodoros Kouris
- David Lindner
- Geoffrey Mareaca
- Margaret A. Darrin Charitable Trust
- Melinda McChesney
- Martin McGraw
- Kevin McGurn
- Mildred Burg Private Foundation
- Sharon Myer
- National Philanthropic Trust
- Laura Neufeldt
- New Tradition Media LLC
- NFL Ventures LP
- James Page
- Pearl Media
- PhRMA
- Polar Products Inc.
- ReelzChannel, LLC
- Lynn Riggs
- Michael Schoenhaut
- Ian Scott
- Marcia Stehling
- Sheldon Taft
- TG Therapeutics
- Alice Tollefson
- James Warren
- Justin Warshowsky
- Brett Wein
- Kerry Yates

#### $1,000 - $4,999
- T Active Media Services, Inc.
- Timothy Adams
- Cliff Adams
- Myrtle Adams
- Rachel Adier
- Avaradhesh Agarwal
- Nancy Albacker
- Margaret Aldredge
- Margaret Aldredge
- Ameriprise Financial
- Ann Amicone
- Elma Andrada
- Susan Anthony
- Antidote
- Jane Arnold
- Matt Arnt
- Robert Asson
- Jim Barden
- Jim Barden
- Michael Barish
- Bruce Bartley
- Michele Bass
- Ann Beach
- Mary Bernis
- Bergen County United
- Way Charitable Flex Fund
- Larry Berglund
- Laura Berry
- Herbert Black
- Matthew Blasi
- Patrice Bogart
- Ronald Bonelli
- Frank Brenner
- Bright Funds Foundation
- John Brown
- Bruchis
- Joan Bunker
- Cheryl Burbano
- Carl Busner
- Susan Bycraft
- Hafer Family Foundation c/o
  - Ronnie Rubin, Trustee
- Annette Cadwell
- CAF America
- Debbie Camp
- Catherine Caneau
- CapTech Ventures, Inc.
- Jay Carella
- Mary K Carlson
- CAS Foundation
- Tim Castelli
- Carol Cetrino
- Thomas Chisholm
- Christiana Care
- Barbara Clark
- Paul Clavellle
- June Clayton
- CME Associates
- Richard Conlin
- Jane Coppola
- Teresa Corl
- Linda Costner
- Tiffany Craft
- Ida Craft
- Lina Craft Pujat
- Susan Creighton
- Crum Manufacturing, Inc.
- Tanya Datel
- Mark Davis
- Troy Desai
- Lydia Diaz
- Digitas Health
- Direct Mail Processors
- Patrick Donaghy
- Fred Dorsett
- Double Eagle Trading Solutions LLC
- William Dow
- Martin Draper
- Jeanne Driscoll
- Paul Ducham
- Margaret Duclos
- Duke Energy Corporation
- Roger Dumas
- Roger Dumas
- Allen Edelson
- Estate of Alice Smith Franks
- Estate of Everett and
  - Petal Turner
- Exelon Foundation Corporation
- Steven Farella
- James Farnsworth

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|(800) 532-7667 / mymsaa.org | 17|
“Thank you for all you do! Giving us the tools, education, and support to endure this disease is life-changing.”

— Ellen via Twitter
“I’m grateful for MSAA and your kind staff and their helpfulness! You have been a blessing to me in my time of need!”

— DM from California

“I thought MSAA had a lot of wonderful programs to highlight MS Awareness Month, such as the importance of cultural sensitivity and early diagnosis for healthcare professionals managing black MS patients. Much needed.”

— Cathy C. via Twitter
MSAA’s Mission:
The Multiple Sclerosis Association of America (MSAA) is a leading resource for the entire MS community, improving lives today through vital services and support.

To help support MSAA’s vital programs and services, please visit mymsaa.org/donate or call (800) 532-7667.