ALS United
Greater New York

Together, We End ALS®

THE IMPACT OF YOUR GIVING

A REVIEW OF OUR ADVANCEMENTS IN 2025

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A REVIEW OF OUR ADVANCEMENTS IN 2025

For more than 30 years, ALS United Greater New York's work has been shaped by local families, caregivers, clinicians, advocates, volunteers, and supporters who believe no one should face ALS alone. ALS United is not just a support organization but rather we are your family of support built on experience, trust, and care. Our local care professionals provide comprehensive, compassionate support that evolves as needs change, a commitment strengthened in 2025 with the launch of **The Tom and Jill Pulley Mental Health and Wellness Program.**

Because of your generosity, 2025 was a year of meaningful progress across our mission. Care Services continued to grow alongside the increasingly complex needs of individuals and families, while education and connection remained central. Families came together through the **5th Annual Education and Resource Summit, Hudson Valley and Long Island Family Connection events,** and throughout the year, ongoing support that ensured trusted information and a caring connection were always within reach.

Your engagement also fueled advocacy, research, and awareness efforts that extended our impact. Together, we helped pass ALS registry legislation in New York and New Jersey, ensuring more accurate data collection of ALS to inform research, resources, and care, and elevated visibility through Lou Gehrig Day at New York ballparks, ringing the opening bell at the New York Stock Exchange, and ALS Awareness Month messaging in Times Square. Fundraising milestones–including $1.3 million raised through WALK ALS and more than $640,000 from our endurance events–supported care, research, and advocacy, culminating in the Lou Gehrig Legacy Gala, which raised more than $1 million as we honored Tom Pulley, Kasper Roet, PhD, Sarah Langs, and Gerrit Cole, and celebrated the collective commitment that makes our work possible.

ALS
United
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CARE SERVICES

The Tom and Jill Pulley Mental Health and Wellness Program

In 2025, ALS United Greater New York launched **THE TOM AND JILL PULLEY MENTAL HEALTH AND WELLNESS PROGRAM.** This innovative program has been established with support from the Pulley family to provide enhanced mental health services for people living with ALS and their loved ones in the Greater New York area.

The Program will create access to specialized and essential psychotherapy inclusively across the community. Through a new full-time Licensed Clinical Social Worker, as well as grants for patients and families with specific mental health needs, this comprehensive initiative will be the first of its kind for those impacted by ALS. These vital resources will meet a substantially growing need in the community and will offer more opportunities for increased quality of life.

The Pulley family worked with ALS United Greater New York to thoughtfully create a much-needed program which provides focused mental health support that is unique to people navigating ALS. The Tom and Jill Pulley Mental Health and Wellness Program blends seamlessly in coordination with our other Care Service programs to ensure personalized care for all.

> **"With all the healthcare expenses I incur every month in fighting this disease, I was blown away by the availability of a grant for my mental health costs. It has helped me immensely and represents real 'caregiving for the caregiver'."**
>
> **– H.A., Lower Hudson Valley**

In addition, the Pulley family has also made possible the ability for notable expansions in the areas of Assistive Technology and Home Accessibility through their overarching newly established named fund, The Tom and Jill Pulley Expansion Fund. ALS United Greater New York is deeply thankful to the Pulley family for their generosity and philanthropic contributions that are significantly improving life with ALS for this community.

CARE SERVICES

WRAPPED IN CARE: MAUREEN'S STORY

When Maureen reached out to ALS United Greater New York last spring, everyday tasks were becoming harder and the path forward felt uncertain. Through one of our nine multidisciplinary ALS clinic partners, she connected with expert clinicians providing comprehensive, state-of-the-art ALS care, alongside local providers she knew and trusted. Being introduced to her regional Care Coordinator brought immediate relief, helping her understand that she did not have to navigate ALS alone.

As her needs evolved, so did the support around her. Adaptive equipment from our free Equipment Loan Program made her home safer and easier to move through, while home assessments with our Assistive Technology Specialist identified practical solutions to support daily living. When entering and exiting her home became a barrier, our Home Accessibility Grant made it possible to install a ramp, restoring her ability to come and go freely and reclaim a sense of independence. Just as important were the moments of connection, as Maureen found encouragement and understanding through our support groups and the reassurance of being part of a community that shows up.

Maureen's story offers a window into how ALS United Greater New York's wraparound, local and personalized care works in real time. In 2025, this support extended to more than 1,200 individuals and families through comprehensive care addressing physical, emotional, and practical needs, including access to free communication devices and durable medical equipment, assistive technology assessments to improve home safety, and opportunities for families to connect and learn through education.

2025 BY THE NUMBERS

1,062 pieces of equipment were provided

1,171 People with ALS received multidisciplinary care and clinical management

1,183 home visits were conducted

1,279 people provided with care

1,785 people attended support groups

223 rides were provided for patients to attend ALS clinic appointments

Financial assistance was provided to nearly **100** families

These numbers represent only a small part of what this care truly means. Behind every piece of equipment delivered, every home made accessible, every clinic visit coordinated, and every support group held is our shared commitment to dignity, connection, and support, so no one in our ALS community has to navigate this journey alone.

RESEARCH

ALS United Greater New York is committed to supporting the most promising research projects and initiatives in the field, and to date we have invested $17 million in ALS research.

ALS UNITED IS PROUD TO PARTNER WITH AND SUPPORT THE FOLLOWING GROUNDBREAKING RESEARCH ORGANIZATIONS AND INITIATIVES:

Milton Safenowitz Postdoctoral Fellowship Program
The Accelerating Medicines Partnership in ALS (AMP-ALS)
Expanded Access | Target ALS
ALS Network Research Program | Robert Packard Center for ALS Research
The Network for Excellence in ALS Consortium (NEALS)

As a signature program of investment, we proudly support the development of bright, young scientists through the Milton Safenowitz Postdoctoral Fellowship Program. The Safenowitz family, through our organization, founded the program in memory of Mr. Safenowitz, who died of ALS in 1998. Fellows collaborate with a senior mentor and receive extensive exposure to the ALS research community. Upon completion of their fellowship, more than 75% of the awardees remain in the field of ALS research.

Additionally, this year, ALS United Greater New York, along with our fourteen other ALS United member organizations announced a new joint research funding partnership with the Robert Packard Center for ALS Research. This collaboration reflects both organizations' commitment to accelerating breakthrough discoveries that will advance understanding of ALS and bring hope to those affected by the disease.

ALS United continues to play an active role in advancing ALS research through meaningful participation in national and international scientific forums. This commitment to research and partnership extended globally, as Chief Care Services Officer Debbie Schlossberg and Dr. Martin McElhiney of Columbia University contributed to the International Alliance of ALS/MND Associations' 2025 Alliance Meeting and Allied Professionals Forum in Toronto, helping to strengthen cross-sector and international collaboration that accelerates clinical trials, improves care, and advances progress for people living with ALS.

Progress in ALS Treatment: Qalsody® Brings Measurable Improvements

Researchers report that long-term use of Qalsody (tofersen), a new drug approved for the treatment of ALS in adults with a confirmed mutation in the SOD1 gene, delays symptom progression and death and in about one-quarter of participants leads to stabilization or improvement. The new findings, published Dec. 22, 2025 in JAMA Neurology, provide long-term follow-up results from a phase 3 trial of Qalsody and its open label extension, that served as the basis for the FDA's approval in 2023 of the drug for this rare form of ALS which accounts for about 2% of all ALS cases.

Some individuals treated with Qalsody are experiencing not only a halting of disease progression but also actual functional improvement, with researchers noting "meaningful preservation of function and suggestions of sustained improvement." Qalsody offers new hope for eligible patients. SOD1 carriers interested in learning more are encouraged to speak with their ALS neurologist.

ADVOCACY | Federal

At ALS United Greater New York, advocacy is a powerful driver of change, one that directly improves the lives of people living with ALS and their families. For more than 30 years, dedicated advocates have helped shape policies, secure critical funding, and elevate awareness, strengthening care and support across our community. Together, we continue to champion policies that reflect the urgency and realities of ALS.

This year's advocacy efforts, powered by people living with ALS, families, clinicians, and community advocates, helped drive meaningful federal investment in ALS research. Through coordinated trips to Washington, D.C., legislative meetings, and participation in advocacy summits, ALS United Greater New York advanced unified funding requests that resulted in significant outcomes.

Congress increased funding for ACT for ALS to $90 million, a $15 million increase over last year, directly reflecting sustained advocacy for faster access to treatments.

While funding for the Department of Defense ALS Research Program (CDMRP) held steady at $40 million, and the CDC ALS Registry remained at $10 million, these investments were preserved amid broader budget constraints.

NIH ALS research funding was maintained at $143 million, reinforcing the importance of continued pressure to grow this critical pipeline.

Notably, advocacy also helped secure $30 million in new funding through ARPA-H dedicated to ALS research, bringing new momentum and innovation to the field.

*** Numbers at the time of printing reflect funding for the Fiscal Year 2025 which is October 1, 2025 - September 30, 2026.**

In May, ALS United Greater New York proudly sponsored and participated in the I AM ALS Summit in Washington, D.C., marking our second consecutive year of engagement. Our team joined advocates from across the country to learn, collaborate, and participate in panel discussions, leading legislative meetings on Capitol Hill, and encouraging constituents to share their stories directly with lawmakers.

In addition, ALS United Greater New York, alongside our partner members of ALS United, continue to help shape a coordinated, inclusive, and effective national advocacy strategy, positioning ALS United Greater New York as a trusted leader in the ALS advocacy space.

ADVOCACY | State

NEW YORK

New York Enacts ALS Registry

A major milestone this past year was the passage of the ALS/MND Registry legislation (S.6413/A.7845), designating ALS as a reportable disease in New York State and establishing a statewide registry to track incidence and prevalence. Signed into law in October 2025, this critical tool will strengthen research, improve care, and guide resource allocation. It was made possible through the leadership of Senator Scarcella-Spanton, Assemblymember Steve Stern, and the dedicated advocates who traveled to Albany and wrote to their elected officials in support.

Additional Legislation and Funding

ALS United Greater New York also continued to champion legislation to protect individuals from genetic discrimination by insurers (S.6124/A.3229) and supported the creation of a Rare Disease Advisory Council (S.1287/A.1296A). While these bills did not fully advance this session, we remain committed to advocating for their passage in 2026 to improve protection and care for people living with ALS and other rare diseases. Our advocacy efforts also led to continued funding in the New York state budget.

NEW JERSEY

New Jersey Enacts ALS Registry

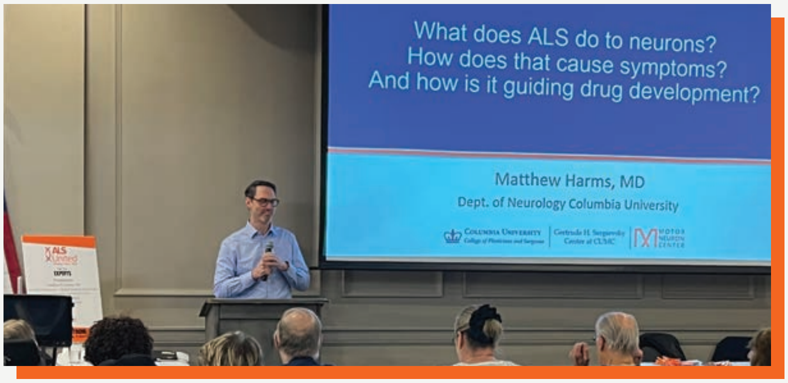
New Jersey achieved a significant advocacy victory with the passage and signing into law legislation designating ALS as a reportable disease statewide. Introduced by Assemblymember Tennille McCoy, with leadership and support from Assemblymember Margie Donlon, M.D., and Senator Vin Gopal, the new law establishes a statewide ALS registry that will improve data collection, inform research, and strengthen care planning. This milestone reflects the power of sustained advocacy and collaboration and marks a crucial step forward for people living with ALS and their families across New Jersey.

New Jersey State Budget Funding

After months of determined advocacy, the New Jersey Legislature reinstated $325,000 in state funding for ALS care in the FY2026 budget, a critical win for the ALS community following a challenging year in which funding for ALS was cut drastically. We are deeply grateful to legislative champions Senator Troy Singleton and Assemblymember Tennille McCoy, along with Assemblymembers Margie Donlon, M.D., Ellen Park, and Senator Andrew Zwicker, whose leadership made this restoration possible. More than a budget line, this funding represents a renewed commitment to people living with ALS and the clinics that care for them, and a reminder of the power of advocacy when a community comes together.

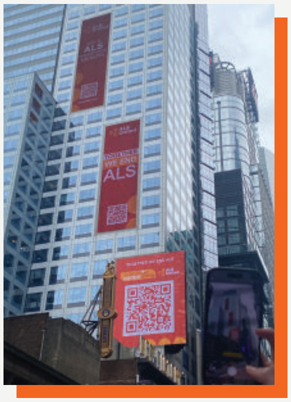
AWARENESS & EDUCATION

Awareness and education are vital in the fight against ALS, empowering families, guiding care, and driving progress in research and advocacy. This past year, ALS United Greater New York brought the community together to learn, connect, and support those living with ALS.



We began the year with a **Talk. Learn. Connect.** research event on Long Island, made possible through the generous partnership and support of the **Joseph Cotrone Foundation** and **Patrizia's Restaurant**. Dr. Matthew Harms of Columbia University shared the latest ALS research updates, while attorney Richard Weinblatt offered practical guidance on Medicaid changes, providing both hope and actionable support for families.

Over the summer, we partnered with the New York Yankees and New York Mets to honor Lou Gehrig's legacy. From **Lou Gehrig Day at Yankee Stadium** to a night of **ALS Awareness at Citi Field**, these community events highlighted ALS awareness and the ongoing need for research and improved care, with on-field presentations alongside I AM ALS, Her ALS Story, MAC Angels, and Target ALS.



During ALS Awareness Month, the 15 members of ALS United across the country collaborated on a high-visibility awareness effort that included digital ads in Times Square, bringing national attention to ALS and the need for continued research and support. This momentum continued with ALS United members participating in the ceremonial ringing of the opening bell of the New York Stock Exchange, further amplifying awareness, and demonstrating the collective strength and unity of the ALS community on a prominent national platform.

AWARENESS & EDUCATION

Together, these events reflect the heart of our mission: bringing people together, sharing knowledge, and building a strong, supportive ALS community. Each connection, conversation, and moment of celebration helps strengthen our resolve, raise awareness, and inspire hope.

Long Island Family Day and **Hudson Valley Family Day** welcomed people living with ALS, their families, and WALK ALS participants. Guests connected with our Care Services team, explored available resources, and built relationships that continue to strengthen our ALS community.

Education and innovation were front and center during a Talk. Learn. Connect. event at the **Yogi Berra Museum & Learning Center**. The event featured a research update by Dr. Jonathon Tiu, Director of Neuro Rehabilitation at Hackensack University Medical Center, while Charles Poeppelman, Market Development at **Synchron**, presented on emerging brain-computer interface technology.

Community spirit was also celebrated through the generosity of **Gary Ragusa and the Team Goose Foundation**, and the **New York Islanders** and **Islanders Children's Foundation**. They provided ALS community members with tickets to John Mulaney, Jon Stewart, and Pete Davidson shows at UBS Arena, as well as a chance to attend a New York Islanders game, complete with custom jerseys and a post-game meet-and-greet with players Casey Cizikas, Anders Lee, Matthew Schaefer, Ryan Pulock, Kyle Palmieri, and Mathew Barzal.

COMMUNITY IN ACTION

Community Advisory Board

ALS United Greater New York is excited to announce the formation of our new Community Advisory Board, made up of people living with ALS and family caregivers. By incorporating the experiences of individuals living with ALS directly into our decision-making, the advisory board helps us stay informed, responsive, and deeply connected to the needs of those we serve. With input from the Community Advisory Board, our team is working together to deliver the best care possible and ensure that no one walks this journey alone. Advisory Board Members include Stephen Fray, Lauren Garcia, Ila Gupta, Les Hanna, Dorothy Pao, Lawrence Modafferi, and Kathleen Ventarola.

Community Fundraisers

Through the creativity and commitment of our local community, independently hosted events raised awareness of ALS and generated more than $450,000 to support care services, advocacy, and research. These gatherings brought people together in meaningful ways—neighbors, friends, and families turning their passion into action. From the annual **Iron Horse Party** to golf outings, sports competitions, and community Walks, each event reflected the power of grassroots leadership, and the difference one idea can make.

Musicians United for ALS: A Night for Wayne Warnecke was a moving tribute that welcomed 300 guests for an unforgettable evening honoring the beloved music producer and sound engineer. Performances by The Bacon Brothers, Patty Smyth, and Rob Thomas filled the room with energy and heart. Proceeds from ticket sales directly supported ALS United, helping to fund care services, advocacy efforts, and research. It was more than a concert—it was a powerful show of solidarity that turned music into meaningful impact.

Bruderhof Community & King Family Summer Farmers' Market for ALS brought neighbors together in a simple but powerful way. Through fresh produce, homemade goods, and shared stories, it became not only a seasonal gathering, but also a way to directly support people living with ALS. Funds raised are helping local families access care services and resources when they need them most. The King Family Market highlights that whether you host a large concert or a neighborhood gathering, your efforts can make a real difference.

Plandome Country Club Ladies' Golf Association hosted their Annual Holiday Fundraiser Luncheon bringing together over 200 attendees to support our Long Island Care Services - providing patient transportation, clinic visits, medical equipment, home accessibility, and mental health services to people and families living with ALS.

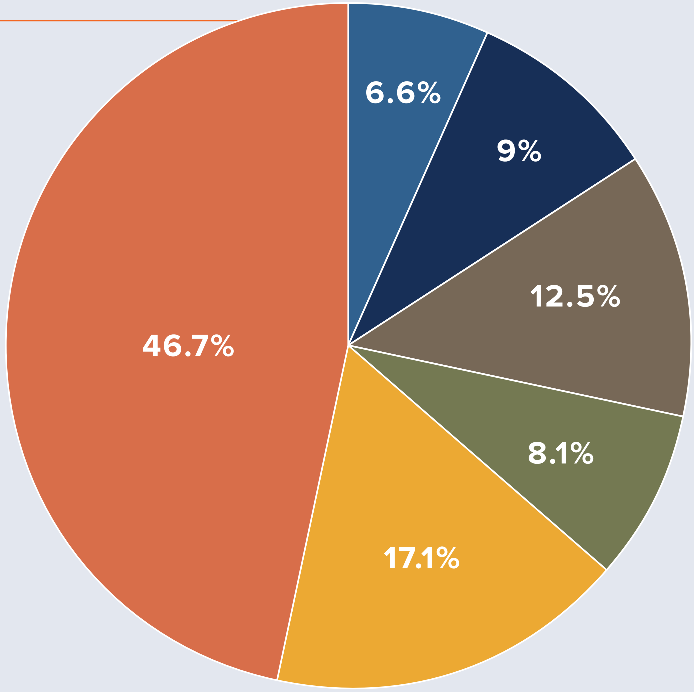
If you are inspired to turn your hobby, interest, or passion into a fundraiser for ALS awareness, we would love to partner with you. Contact Charlotte Seto, Senior Manager, Fundraising & Special Events, at 212-720-3054 or cseto@als-ny.org.

YOUR SUPPORT

The 2025 fiscal year ending January 31, 2026, brought in $7.7 million across all fundraising, including events, major gifts, foundation grants, and bequests. The dedicated support from donors at every level enhances our vital programs to best serve people living with ALS and their families across the Greater New York Community.

WHERE ARE YOUR DONATIONS GOING

- **CARE SERVICES: 46.7%**
- **CLINICAL CENTERS: 17.1%**
- **RESEARCH 8.1%**
- **ADVOCACY, PUBLIC AWARENESS & EDUCATION: 12.5%**
- **FUNDRAISING: 9%**
- **ADMINISTRATION: 6.6%**



*** Based on financial statement for Fiscal Year Ending 1/31/2025**

ALS United Greater New York is proud to be one of 15 member organizations with a shared mission: local care, strong advocacy, and progress toward a cure.

Together, We End ALS®.

ALS United Greater New York is proud to be a part of a very select group of high-performing charities that have earned a Four-Star rating from Charity Navigator, America's largest independent charity evaluator. In 2025, we earned our twelfth consecutive Four-Star rating from Charity Navigator. That means we have a proven record of being accountable, transparent, and fiscally responsible, further validating that our supporters can trust our commitment to good governance and financial health.

Charity Navigator
✦ ✦ ✦ ✦
✦ FOUR-STAR ✦

EVENTS

WALK ALS

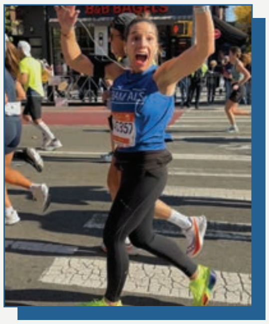
In 2025, we held **Walk ALS** events from May through October in New York City, Westchester, North Jersey, the Jersey Shore, Long Island, and in the Hudson Valley. With more than 4,000 walkers in attendance, $1.3 million was raised to support programs, services, and research. We are grateful to the many dedicated and visionary team captains, event participants, volunteers, sponsors, and ALS advocates who participated. Register to join us for one of our six walks in 2026 at **alswalks.org**.

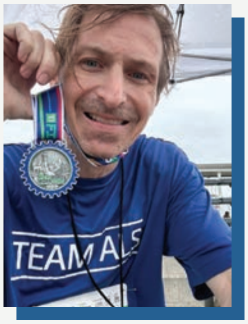
THANK YOU TO THE TOP TEN TEAMS OF 2025

FOR THE LOVE OF JEMMA
Pablo's Pals
Team Yorktel
4K
Team Jim L'Ecuyer & Friends
Dani's Defenders
Our Love for Mona
Katz Walkers
Team Karle
Team Mike Leiderman

ALS IRON HORSE GOLF CLASSIC

The **ALS Iron Horse Golf Classic** marked its eighth successful year on Monday, August 11, 2025, at Sands Point Golf Club. More than 100 golfers came together to enjoy a beautiful day on the course and helped to raise more than $150,000 in support of our services and programs. Save the date for the ninth Annual ALS Iron Horse Golf Classic on Monday, August 10, 2026!





TEAM ALS

Over the course of 2025 **Team ALS** raised more than $475,000, thanks to the efforts of 100 athletes from around the world, and a significant representation from our local community in Greater New York. These athletes participated in the TCS New York City Marathon, United Airlines NYC Half, NYCRuns Brooklyn Experience Half Marathon, TD Five Boro Bike Tour, and individual endurance events. We are actively recruiting participants to join Team ALS for one of our 2026 endurance events. If you are interested, please contact Charlotte Seto at 212-720-3054 or cseto@als-ny.org.

EVENTS

LOU GEHRIG LEGACY GALA

The annual **Lou Gehrig Legacy Gala** took place at Pier Sixty on Tuesday, November 11, 2025. Hosted by ESPN's Jeremy Schaap, with awards presented by Bob Costas, the event celebrated the year's achievements on behalf of the ALS community. The evening paid special attention to veterans living with ALS and highlighted the presentation of several prestigious awards honoring individuals whose exceptional contributions have made a lasting impact on the ALS community.

The Jacob K. Javits Lifetime Achievement Award was presented to **Tom Pulley**, who was diagnosed with ALS in 2024. Tom's award celebrated the launch of The Tom and Jill Pulley Mental Health and Wellness Program, an innovative initiative providing enhanced mental health services for people living with ALS and their loved ones across the Greater New York area.

The Champion Award was presented to **Kasper Roet, PhD,** Founder and CEO of QurAlis Corporation. As a representative of the future of ALS research, Kasper, a neuroscientist specializing in gene and stem cell–based precision medicine for ALS and other neurodegenerative diseases, has a long-standing connection to ALS United Greater New York. In 2015, he was awarded the Milton Safenowitz Postdoctoral Fellowship, funded in part by our organization.

The Lou Gehrig Sports Award was presented to **Gerrit Cole**, New York Yankee, six-time MLB All-Star and 2023 American League Cy Young Award winner. **Sarah Langs**, reporter and researcher for MLB.com, received the **Iron Horse Award** for her outstanding contributions to baseball and her inspiring presence in the sports community, all while living with ALS since 2021.

The Gala's impact extended beyond heartfelt and inspirational speeches and awards. With generous contributions from supporters, the **event raised $1 Million**. These funds directly support ALS United Greater New York's mission, providing free services that include the equipment and assistive technology loan program, care coordination and support, transportation to ALS clinics, and much more.

HONORING THEIR LEGACY

As we remember the lives of three extraordinary individuals who have helped to shape the Greater New York ALS community, their legacies of compassion and service will continue to inspire our mission.

ALAN GRIFFITH, retired Vice Chairman of the Bank of New York and a former Chairman of our Board of Directors, was a guiding force in the fight against ALS for nearly five decades. His commitment began in the 1970s as a board member of the New York-based ALS Foundation, and his steady presence continued through the pivotal 1985 merger that formed The ALS Association, where Alan served on the national Board of Trustees. His leadership and generosity were instrumental in forming the foundation for what would become ALS United Greater New York. To this day, our work reflects Alan's dedication to serving people and families living with ALS in our community.

ALICE CLAAR, who joined our Board in 1995, brought a deeply personal commitment to our cause, honoring the memory of her brother Sheldon, who died of ALS in 1987 at just 53 years old. Over the past 30 years, Alice served on our Board's Patient Services Committee and fought tenaciously for supporting families, improving care, and creating a future without ALS.

Alan and Alice were two of our founding board members whose dedication, leadership, and vision helped to build our organization from its earliest days. We are deeply grateful to them for their invaluable efforts.

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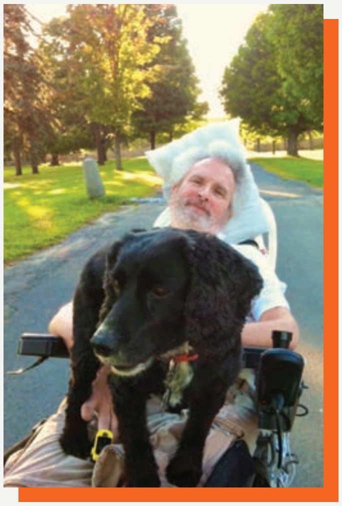

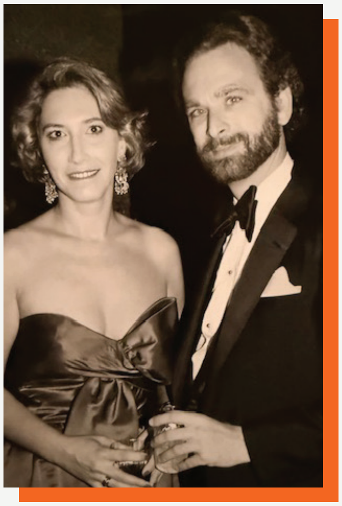

JAY MORTIMER lived with ALS for 18 years until the age of 74. He and his wife Prue have been instrumental in growing the Equipment Loan Program over the past 14 years. Together, along with family support from The Harriman Foundations, they have contributed nearly $1M to ALS United Greater New York. Jay and Prue wanted to contribute in a way that could tangibly help others dealing with ALS. Having experienced firsthand the impact of assistive technology and durable medical equipment, they found a perfect way to support the mission. In reflecting on their giving, Prue shared that, "it is always such a highlight for us and we have been thrilled to do it." Over the years, ALS patients and caregivers have come to Prue at support groups and hospitals thanking them for their incredible support. Greatly missed, Jay is remembered for his warmth and kindness, as well as his bravery in the face of adversity. ALS United Greater New York is honored to serve Jay's legacy of helping others through our deep commitment to continue enhancing the Equipment Loan Program.

PLEASE JOIN US IN THANKING OUR TOP DONORS OF 2025

Andrew Abramsky
John Albuerne Family Foundation
Joseph Alexander Foundation
Allied World
ALS Ride For Life, Inc.
Estate of Lois B. Altman
Anonymous
Linda and Paul Barnett
Fred and Janet Baron
Dan Bass
BBDO
Jeff Beer and Mia Graham-Beer
Eric and Jen Berniker
Vincenzo Bilotta
The Birk's Works Foundation, Inc.
Bloomberg L.P.
Bloomberg Philanthropies
Thomas and Ashley Bradley
Devon and Pete Briger
Jack Broderick
Lee and Rebecca Brodsky
Ann Marie Carr
Craig and Tammi Chalfie
Norm and Noriko Chen
Daniel and Rhonda Chestler
Steven Chestler
Cinque Foundation, Inc.
William and Amy Clark
The Cobb Foundation, Inc.
The Gerrit and Amy Cole Foundation
The Joseph Cotrone Foundation
Louise M. Cox
Crane Charitable Funds
Christine Currence
The Dahlquist Foundation
Kirk and Peggy Davenport
Bayley and Michele Davis
Elizabeth DeJean
Lisa Detwiler
Dial A Bug
Molly and Chris Dillon
Eli Edwards
Ike, Molly & Steven Elias Foundation
Emanuel Lutheran Church
Mark Engel
James and Nina Essey
Paul Falgout
Barbara and Patrick Fallon
Fortress Investment Group
Joel Frank
Concetta Gallo
Genatt V Insurance Solutions
Kevin and Terry Glassman
The Gluck Family Foundation
Sandra Gluck
Eric Golden
Kieran and Catherine Goodwin
Estate of Barbara G. Gottlieb
Eldridge and Kirsten Gray
Greater New York Mutual Insurance Company
Alan and Penny Griffith
Jane W. Griffith
Anita Gupta
Connie Gyftakis and Bryan Yaubrough
Alan and Susan Hammer
David Hammerman and Kyle Gerstenschlager
Christine Harper
Mary and Kathleen Harriman Foundation
Mary W. Harriman Foundation
Livia Helmer
Jane and Chris Hentemann
Caren Hoffman and Family
Helen Hoffritz Charitable Trust
Holzer Family Foundation
Hospital for Special Surgery
Hudson Valley Walk ALS Committee
IBM Corporation
Intex Solutions, Inc.
Leslie Jorgensen
Sonny and Michelle Kalsi
John and Elaine Kanas Family Foundation
Amanda and Daniel Kaplan
Katz Realty Group
The Deana and Sheldon Katz Fund
Sheldon C. Katz Memorial Fund
Laurie Kayden Foundation
Jeffrey Kelly
Cristina and David Kepner
Keskin Family Fund
The Danielle Kirrane Foundation
Roger Klauber Family Foundation, Inc.
Gloria J. Kosek
Beth Kougasian
Marc and Carol Lane
Robert Larson
Stanley and Gene Lasdon Foundation
Alan and Susan Levine
The Fay J. Lindner Foundation
The Jeffrey H. Loria Family Charitable Foundation
Erica Miles
Cathy and Jonathan Miller
William and Francine Monti
Prue Mortimer
Marianne Muise
Domenick Muro
The National Mah Jongg League Foundation, Inc.
New Jersey State Department of Health
New York State Department of Health
New York Yankees Foundation
The William and Anita Newman Foundation
Jacqueline Nieman
Nonna's Garden
NuMotion Foundation
NYU Langone Medical Center
Susan Ochs-Scher
The Eleanor R. Orberg Fund
Paralyzed Veterans of America Education Foundation
Patrizia's
John Pjeternikaj
David Pelton
Plandome Country Club
Plandome Ladies' Golf Association
Rita Plush
Allen and Emily Popowitz
Judith Pulig
Jeff Pulley
John and Mary Pulley
Tom and Jill Pulley
QurAlis Corporation
Cynthia Rimpo
Rockefeller Group International, Inc.
John V. and Susan Roos
Richard Rothman and Melissa Salten
Donna and Mark Rubin
William and Ashley Rue
Estate of Ronald J. Ryan
Jin Ryu
The Milton and Marilyn Safenowitz Family Foundation
Daniel and Sara Sapadin
Ellen Sarna
Jeremy Sarna
Estate of Francis M. Schaefer
Judythe and Bennet Schonfeld
Linda and Jesse Singh
Tim Sloan
SS&C Intralinks
Thomas F. Staley Foundation
Alan L. Stange and Gloria M. Trillo
Thomas Stemwedel
Holly Stitley
Richard Stravitz and Kathy Thompson
Tanabe Pharma America, Inc.
Textile Import LLC
The TJX Foundation, Inc.
The Tow Foundation
The Trayner Group Ltd
Carol Tschupp and Linda Getman
Rob and Amy Tuchman
Alexandra and Johan Utterman
Marene and Robert Van Farowe
York Wang
Arthur and Joanne Weinbach
Nigel and Christine Weston
Wise Music Corporation
Melissa and Ronnie Wise
Bill Wolf Petroleum Corp.
Xena Capital
Yorktel
David and Lynette Zinberg

Thank you to everyone who contributed to the mission of ALS United Greater New York in 2025! Without your support, our programs and services would not otherwise be possible. For more information on how you can expand your mission-based support through various ways of giving, such as estate planning, IRA rollovers, or gifts of stock and securities, please contact: Vivian Banks, Director of Individual and Institutional Giving | 212-720-3051 or vbanks@als-ny.org

2026 EVENTS

SUNDAY, MARCH 15
United Airlines NYC Half Marathon

SUNDAY, APRIL 26
NYCRuns Brooklyn Experience
Half Marathon

SATURDAY, MAY 2
Walk ALS New York City
South Street Seaport
Pier 16, NYC

SUNDAY, MAY 3
TD Five Boro Bike Tour

MAY 14 –15
Education and Resource Summit
Virtual

SATURDAY, MAY 30
Walk ALS Westchester
Manhattanville University
Purchase, NY

MONDAY, JUNE 2
Lou Gehrig Day
Throughout Major League Baseball

SUNDAY, JUNE 7
Walk ALS North Jersey
Saddle River Park
Saddle Brook, NJ

MONDAY, AUGUST 10
ALS Iron Horse Golf Classic
Sands Point, NY

SATURDAY, SEPTEMBER 19
Walk ALS Long Island
Eisenhower Park
East Meadow, NY

SATURDAY, SEPTEMBER 26
Walk ALS Jersey Shore
Long Branch, NJ

SUNDAY, OCTOBER 18
Walk ALS Hudson Valley
Walkway Over the Hudson
Highland, NY

SUNDAY, NOVEMBER 1
TCS New York City Marathon

TUESDAY, NOVEMBER 10
The Lou Gehrig Legacy Gala
Pier Sixty, NYC

Our Mission is to support and empower the ALS community through a collaborative approach to fostering bold research initiatives, advancing national and state advocacy, and providing comprehensive care and support to individuals and families affected by ALS.

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**in memoriam*

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