National Ataxia Foundation

2025 ANNUAL REPORT

Accelerating Progress.
Building Community.
Advancing Hope.

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ABOUT NAF

Since 1957, NAF has been at the forefront of efforts to find treatments and a cure, providing hope and vital resources to the Ataxia community.

VISION

A world without Ataxia

MISSION

To accelerate the development of treatments and a cure while working to improve the lives of those living with Ataxia.

CORE VALUES

- Accountable
- Caring
- Collaborative
- Impactful
- Motivated

NAF
National Ataxia Foundation

LETTER FROM THE CEO

Andrew Rosen

Chief Executive Officer

Every so often, an organization reaches a moment that changes what it believes is possible.

For NAF, that moment came in 2025 in the form of an extraordinary anonymous gift. We don't know the donor's name. We don't know their story. What we do know is that they placed an incredible amount of trust in our mission and in our ability to make that generosity count. Rather than asking, "What can we spend this on?" we asked a different question: **How can we honor this gift in a way that creates the greatest possible impact for people living with Ataxia?**

The answer wasn't immediate.

Over the course of many months, our staff and Board of Directors came together in an ambitious planning process unlike any we've undertaken before. Guided by feedback from our community and organized around four strategic priorities – research, support, advocacy, and philanthropy – we challenged ourselves to think bigger than ever before. Together, we explored hundreds of ideas, debated priorities, and ultimately built a three-year strategic plan designed to accelerate treatments while improving lives today.

The result is more than a plan on paper. It is a commitment to strengthen the programs our community depends on while investing in bold new initiatives, from enhancing our support network with professional social work services to building research resources like a centralized open-access cell line repository and expanding our advocacy efforts so the voices of people living with Ataxia are heard where decisions are made.

As I look ahead, I'm struck by something I've never been able to say before: **we have an opportunity today that would have been difficult to imagine just five years ago.**

Research is advancing at an unprecedented pace. Our community continues to grow stronger and more connected. More partners than ever are working alongside us to move treatments forward. And now, thanks to this extraordinary act of generosity and the thoughtful planning that followed, we are better positioned than ever to help shape what comes next.

Progress has never been the work of one person or one organization. It belongs to every researcher pursuing a new discovery, every volunteer leading a support group, every advocate sharing their story, every fundraiser and donor investing in hope, and every family refusing to give up.

Thank you for believing in this mission and for being part of this remarkable community. The work ahead is ambitious, but so is this community. Thank you for your trust, your partnership, and your unwavering commitment to a future with better treatments and, one day, a cure.

With gratitude,
Andrew Rosen
Chief Executive Officer

Accelerating Treatments. Improving Lives.

Community Voices → Thoughtful Stewardship → Strategic Investment

SO WHAT IS THE PLAN?

In 2025, NAF launched a new three-year Strategic Plan that will guide our work through 2028. Built around four strategic pillars, the plan provides a clear roadmap for accelerating treatments and a cure while improving the lives of people affected by Ataxia today.

SUPPORT

Empowering Ataxia Families

Goal: Deliver a modern, scalable, and inclusive support system that meets the evolving needs of the Ataxia community.

RESEARCH

Driving Scientific Progress

Goal: Accelerate research progress by building the infrastructure necessary to fast-track the discovery of treatments and a cure.

ADVOCACY

Championing the Patient Voice

Goal: Strengthen NAF's advocacy and influence to drive awareness and policy for Ataxia.

DEVELOPMENT

Fueling Mission Through Philanthropy

Goal: Build a funding "flywheel" capable of sustaining NAF's mission.

These priorities strengthen our ability to invest in research, expand support and education, advance advocacy, and grow the resources needed to create lasting impact for the Ataxia community.

2025 AT A GLANCE

$3.5
Million invested in research

$1.3
Million invested in support programs

41
Clinicians trained at ACT

674
Support Group meetings

23
Research studies funded

78
Educational sessions

43
NAF Ataxia Centers of Excellence

99
Congressional meetings

OUR MEMBERS

Every person impacted by Ataxia has a role in moving our mission forward. From individuals newly navigating a diagnosis to longtime advocates, caregivers, researchers, and healthcare professionals, our members help build connections, advance research, and create a stronger future for the Ataxia community.

20,751
Members

12,279 Living with or at-risk of Ataxia

5,642 Family or Friend of person with Ataxia

826 Clinicians

621 Researchers

279 Industry

IN THEIR OWN WORDS

The voices of the Ataxia community are at the heart of our work. Throughout the year, members share their personal stories, offering honest reflections, hope, and encouragement to others navigating a similar journey. Explore more inspiring stories at www.ataxia.org/members.

KAREN W

COMMUNITY MEMBER

As a child, in the 1960s–1970s, I can remember my paternal grandfather walking with a wide, somewhat unbalanced gait. He was born in 1896 in rural Michigan, and never diagnosed with a specific disorder. I just thought his problems were due to normal aging. My father, born in 1924, starting showing similar problems with balance and gait in his 50's. It was not until the mid-1990's that my father's condition was identified as a specific disease: Spinocerebellar Ataxia. I was caregiver to my father, along with a brother, during his later years. It never once occurred to me that I would one day be diagnosed with this same condition.

"It is good to connect with a community of patients with a similar diagnosis and have regular access to experts in the field, as well as be kept aware of new developments in treatment."

COMMUNITY SERVICES

Living with Ataxia is easier when no one has to face it alone. Through education, support, and meaningful connections, NAF's Community Services programs help individuals and families build community, access trusted resources, and navigate their journey with confidence.

$1.3M

Spent on Community Services

674

Support Group Meetings

99

Congressional Meetings

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SUPPORT IMPACT

Connection can make all the difference when living with Ataxia. NAF's Support Groups bring individuals and families together to share experiences, learn from one another, and find encouragement along the way. These welcoming communities are made possible by our dedicated volunteer Support Group Leaders, whose passion and commitment help ensure no one has to navigate Ataxia alone.

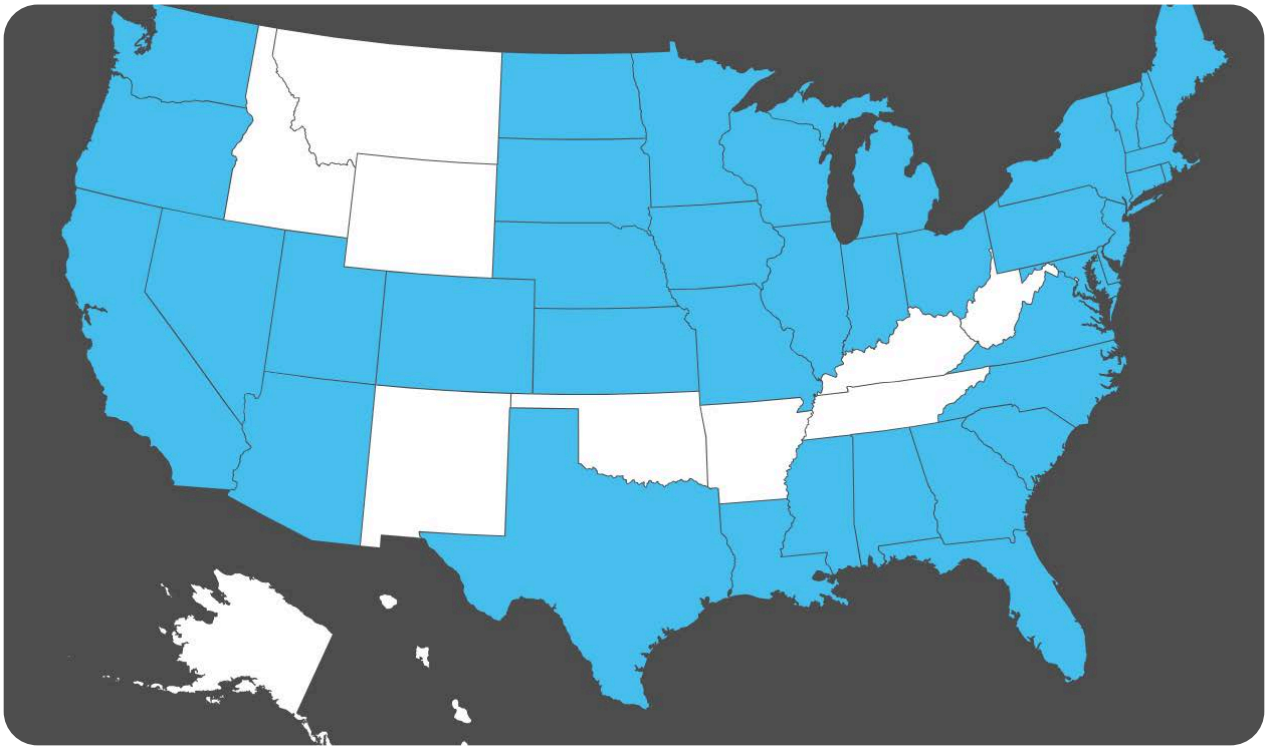
Shannon had this to say about a meeting...

> The day was full of laughter, good conversation, and a reminder of how special it is to spend time together in person.

— **Shannon Dunphy Lazo,**
St. Louis Ataxia Support Group Leader

THANK YOU TO OUR VOLUNTEER SUPPORT GROUP LEADERS!

52 SUPPORT GROUPS IN 41 STATES



13 INTERNATIONAL OR DEMOGRAPHIC GROUPS

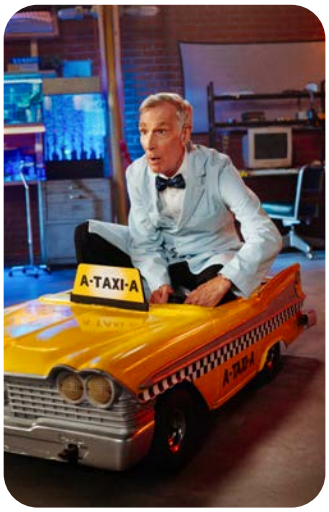
- Ataxia Support Group Canada
- Global Ataxia Support Group Leader
- Ataxia Resources and Discussion Group
- 18-24 Years Old with Ataxia Support Group
- 25-35 Years Old with Ataxia Support Group - The Dominoes
- 35-55 Years Old with Ataxia Support Group
- 55+ Years Old with Ataxia Support Group
- African Americans with Ataxia Support Group
- Spouses and Partners of Loved Ones with Ataxia Support Group
- Parents with Ataxia Support Group
- Spanish Speaking Ataxia Support Group
- U-MATTER, Unbroken - Military Members with Ataxia Talking Team Room
- Women with Ataxia International Support Group

ADVOCACY

Strong advocacy helps create a better future for the Ataxia community. NAF works alongside individuals, families, researchers, and partners to elevate the voices of those affected by Ataxia, advance policies that support research and access to care, and ensure the needs of our community are represented at every level of government.

2025 ADVOCACY ACHIEVEMENTS

- September 25th designated as National Ataxia Awareness Day
- 99 congressional meetings to advocate for the Ataxia community
- 80 House and Senate requests supporting continued inclusion of Hereditary Ataxia in the FY26 PRMRP/CDMRP
- Bill Nye was *Back in the Lab* with four impactful Friedreich's Ataxia awareness videos.



HILL DAY - SEPTEMBER 17, 2025

United Against Ataxia Hill Day connected advocates from across the country to Capitol Hill to share their stories and champion the needs of the Ataxia community. Together, participants met with members of Congress and their staff to advocate for policies that support research, improve access to care, and accelerate the development of new treatments.

ANNUAL ATAXIA CONFERENCE (AAC)

NAF's Annual Ataxia Conference continues to bring the global Ataxia community together to learn, connect, and inspire one another. In 2025, hundreds of individuals living with Ataxia, family members, healthcare professionals, researchers, and industry partners gathered to explore the latest advances in research and care while building meaningful connections. Through expert-led education, shared experiences, and a strong sense of community, the conference remains a cornerstone of hope, collaboration, and progress.

> The presenters did a very good job going over the different assessments that Ataxians have to complete for be able to receive the different agencies services. There is hope that research continues to develop medications to help folks that already have had Ataxia symptoms for a number of years.
>
> -2025 AAC Attendee

HIGHLIGHTS

838 Attendees

223 First Timers

27 Countries

60 Travel Grants
= $66,000

128 Virtual Scholarships
= $3,300

RESEARCH

Research is the foundation of progress. By investing in innovative science, supporting researchers, and fostering collaboration, NAF is helping advance discoveries that bring us closer to better treatments and, ultimately, a cure for Ataxia.

$3.5M

Spent on Research

23

Research Studies Funded

440

Natural History Study Research Visits

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RESEARCH IMPACT

Behind every research breakthrough is a dedicated scientist working to better understand Ataxia. Our *Faces of Ataxia Research* series introduces the researchers whose work is made possible through the generosity of our community, highlighting their passion, discoveries, and commitment to developing better treatments for people affected by Ataxia.

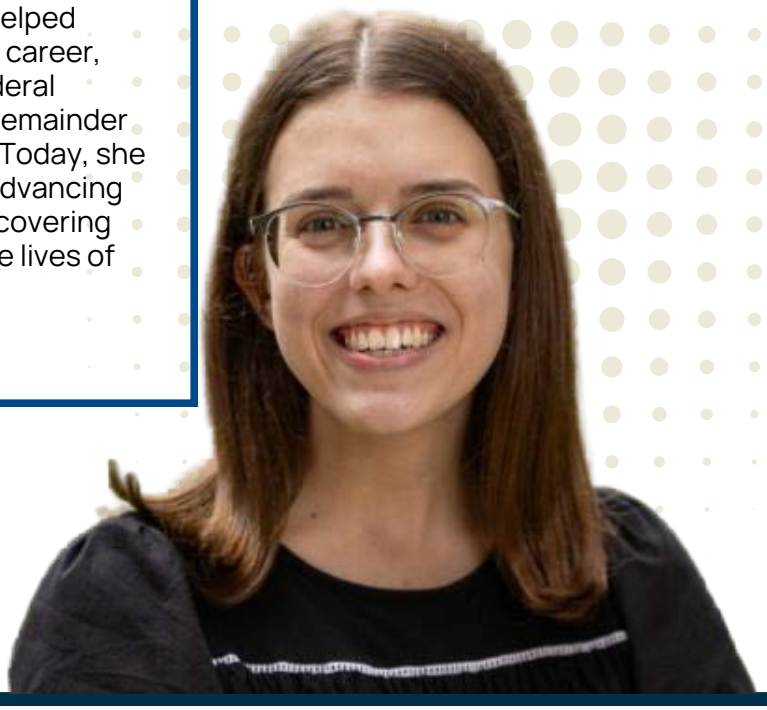
ALEXANDRA PUTKA

ATAXIA RESEARCHER

When Alexa Putka joined Dr. Hayley McLoughlin's laboratory at the University of Michigan, she set out to answer important questions about Spinocerebellar Ataxia Type 3 (SCA3). With support from an NAF research grant, Alexa identified important biological changes that could help guide future treatment development, leading to the first large-scale study of brain lipids in SCA3. The findings were published in the journal Neurobiology of Disease and opened new opportunities for collaboration and continued research.

NAF's investment also helped launch Alexa's research career, leading to additional federal funding to support the remainder of her doctoral studies. Today, she remains committed to advancing Ataxia research and discovering new ways to improve the lives of people living with SCA3.

Alexa hopes her research can provide new answers to patients on SCA3 pathology. She hopes her work may give patients more information on the biology of SCA3 and potential therapeutics being researched.



Research Grants

Advancing research begins with investing in the ideas and people shaping the future of Ataxia care. In 2025, NAF funded innovative research across six grant categories, supporting investigators at every stage of their careers—from graduate students and early-career scientists to established researchers developing potential therapies. These investments are helping expand scientific knowledge, accelerate treatment development, and build the next generation of Ataxia researchers.

Impact By the Numbers

$910,000
Research Investment

18
Institutions Funded

7
Countries Represented

12
Ataxia Types

13
Early-Career Investigators

Ideas into Research

170 Letters of Intent

50 Full Applications

23 Funded Projects

New in 2025 - Supporting Innovative MSA-C Research

In 2025, NAF launched the **Michael Lundquist Grant to Advance MSA-C Research**, a new $50,000 grant supporting innovative research focused on Multiple System Atrophy – Cerebellar subtype (MSA-C). Established in honor of Michael Lundquist, the grant reflects NAF's ongoing commitment to advancing research across the Ataxia spectrum and supporting discoveries that improve the lives of those affected by MSA-C.

NATURAL HISTORY STUDY

The Clinical Research Consortium for the Study of Cerebellar Ataxia (CRC-SCA) brings together leading clinicians and researchers across the United States and Canada to strengthen Ataxia research and accelerate the development of new treatments. Through its natural history study, the consortium follows how Ataxia changes over time, creating essential data that helps researchers design stronger clinical trials and understand whether potential therapies are making a meaningful difference.

This information is also an important resource for pharmaceutical companies developing new therapies. Natural history data helps inform clinical trial design, provides critical context for interpreting trial results, and can support the evidence submitted to regulatory agencies as part of the treatment approval process.

We are deeply grateful to the individuals and families who participate in this long-term research. Their commitment is helping build the foundation for tomorrow's treatments and bringing hope to the entire Ataxia community.

HIGHLIGHTS

1298 Enrolled Participants

146 Baseline Visits

297 Follow-Up Visits

> Two family members participating in the CRC-SCA declined their participant stipend at the end of the research visit. They do this every year- they just want the money to go back towards research.
>
> - CRC-SCA Site Coordinator

Thank you to the leadership and clinical sites of the CRC-SCA for your commitment to advancing Ataxia research. Together with the individuals and families who participate in these studies, you are helping build the foundation for tomorrow's treatments.

CRC-SCA
Clinical Research Consortium for the Study of Cerebellar Ataxia

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DRUG DEVELOPMENT COLLABORATIVE

Partnerships are essential to advancing new treatments for Ataxia. Through the NAF Drug Development Collaborative (DDC), we work alongside leading pharmaceutical companies to accelerate research and therapy development while strengthening the resources that support the Ataxia community. These collaborations help advance critical initiatives, including the CRC-SCA Natural History Study, free genetic counseling, and expanded access to research opportunities—bringing us closer to better treatments.

Thank you to our DDC members for your continued collaboration and investment in advancing research and improving the future for everyone affected by Ataxia.

arrowhead pharmaceuticals · Biogen · biohaven · LoQus23 THERAPEUTICS · PTC THERAPEUTICS measured by moments · REPROCELL · SERVIER moved by you · SKYHAWK THERAPEUTICS · Solaxa · VICO Therapeutics

Free Genetic Counseling and Testing

Access to genetic information can provide clarity, guide care, and help individuals make informed decisions about their future. Through NAF's Genetic Counseling and Testing Initiative, eligible U.S. adults with a family history of SCA1, SCA2, SCA3, SCA6, SCA7, or SCA27B can receive free virtual genetic counseling and testing. Supported by NAF's DDC, this program helps expand access to expert guidance while connecting more individuals and families to the resources they need.

2025 HIGHLIGHTS

191 Genetic Tests

199 Pre-Test Counseling Sessions

134 Post-Test Counseling Sessions

CLINICAL SERVICES

Access to expert care is essential for people living with Ataxia. NAF's clinical initiatives help expand specialized care, support healthcare professionals, and improve access to the resources and expertise individuals and families need throughout their Ataxia journey.

41

Healthcare Providers Trained in Ataxia Care

43

Ataxia Centers of Excellence

826

Clinician Members

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ATAXIA CLINICAL TRAINING

Expanding access to expert care starts with educating the next generation of healthcare providers. NAF's Ataxia Clinical Training (ACT) program offers an immersive, hands-on learning experience for neurology residents, fellows, and clinicians, helping them build the knowledge and confidence needed to diagnose and care for individuals with Ataxia.

As more clinicians receive this training, individuals and families can benefit from earlier diagnoses, improved care, and greater access to healthcare professionals who understand the complexities of Ataxia.

HIGHLIGHTS

41 Attendees

25 Fellows

4 Returning Attendees

In 2025, ACT was dedicated in memory of Carol Tate, a beloved member of the Ataxia community, former co-leader of the St. Louis Ataxia Support Group, and longtime contributor NAF. Carol generously shared her wisdom, compassion, and lived experience to help others navigate life with Ataxia. Her legacy lives on through ACT, inspiring the education of future healthcare providers and improving care for the Ataxia community.

What Attendees are Saying

"Thanks for doing this - the mix of pharma, clinicians, and patients was superb. This is one of the best conferences that I have attended."

"I feel more comfortable recognizing ataxias and just better knowing about NAF as a resource to help when I have questions."

"Very educational and I plan to start an academic center ataxia clinic."

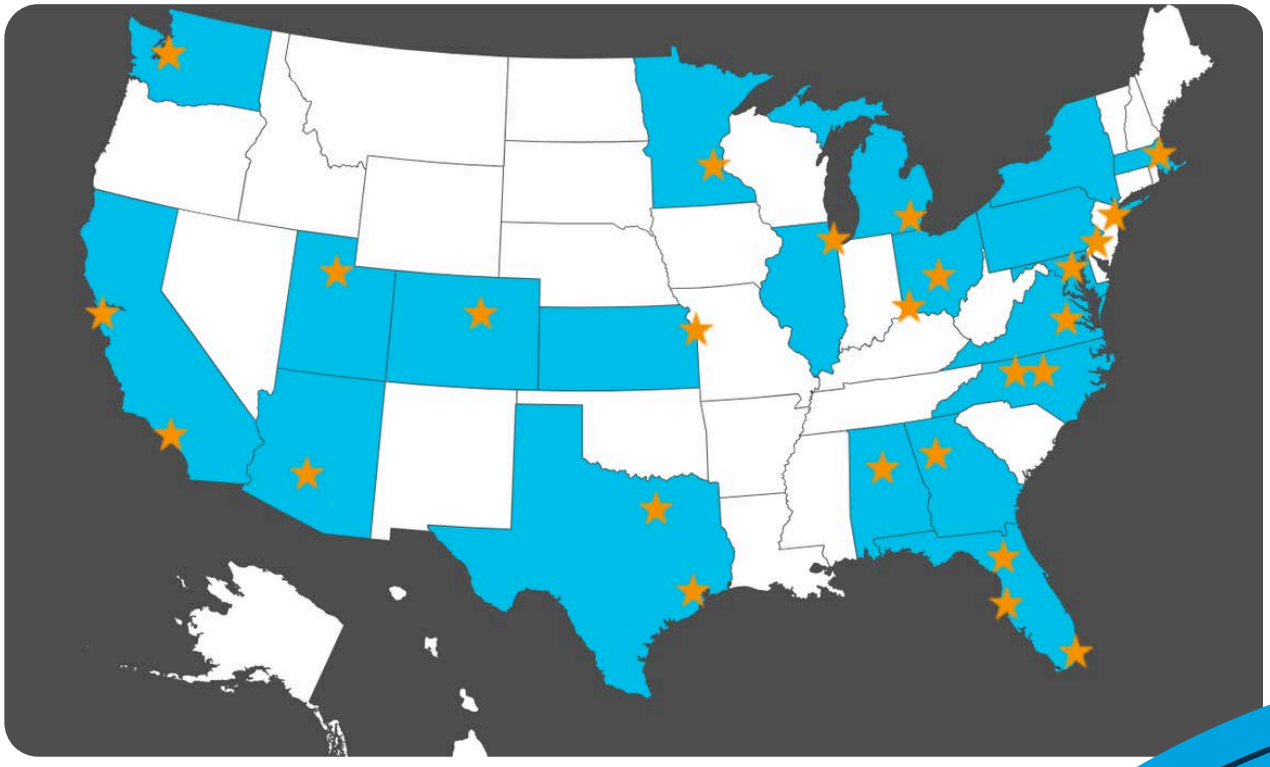
Ataxia Clinical Training

NAF ATAXIA CENTERS OF EXCELLENCE

Access to expert care can make a meaningful difference for people living with Ataxia. NAF's Ataxia Centers of Excellence (ACE) program recognizes leading clinics that provide comprehensive, multidisciplinary care through teams of neurologists, genetic counselors, rehabilitation specialists, social workers, and other experts working together to meet the unique needs of each individual.

NAF National Ataxia Foundation
ATAXIA Centers of EXCELLENCE
Experts in Care & Collaboration

In 2025, NAF's growing ACE network included 32 centers across the United States and 11 international sites. Beyond delivering specialized care, these centers advance research, support clinical trials, educate healthcare professionals, and help expand access to high-quality Ataxia care around the world.



HIGHLIGHTS

32 U.S. Designations

11 International Designations

9 New Sites Added

EDUCATION

Knowledge empowers our community. NAF provides trusted educational programs and resources that help individuals, families, caregivers, and healthcare professionals better understand Ataxia, make informed decisions, and navigate every stage of the journey.

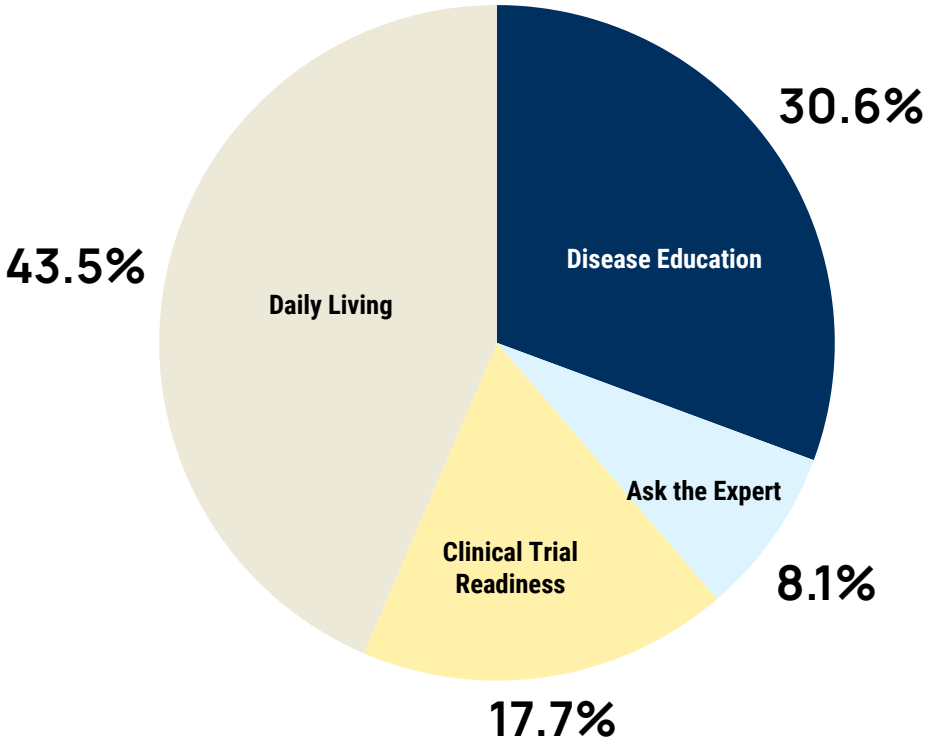
HIGHLIGHTS

18,968 People Reached

81 Webinars/ AAC Sessions

44 Articles

Webinar Topics Covered in 2025



COMMUNITY FUNDRAISING

Meaningful progress begins with a community that cares. Thanks to the generosity and dedication of our supporters, NAF continues to invest in research, expand vital programs, and build a brighter future for everyone affected by Ataxia.

15

Walk N' Rolls

14

Passion Fundraisers

$359K

Raised

NAF
National Ataxia Foundation

WALK N' ROLL TO CURE ATAXIA

Walk N' Roll events bring communities together to raise awareness, celebrate connection, and support NAF's mission. Organized by dedicated local volunteers, these events unite individuals, families, and supporters in making a meaningful difference for the Ataxia community. Through every walk, roll, and fundraiser, participants help advance research, expand support programs, and inspire hope for a future without Ataxia.

HIGHLIGHTS

- **15** Walk N'Rolls
- **1628** Donations
- **$245K** Raised

2025 Walk Locations

Atlanta	Baton Rouge	Central Ohio	Dallas
Denver	Jacksonville	Las Vegas	Maryland
Massachusetts	Milwaukee	Minnesota	Pittsburgh
St. Louis	Tampa Bay	Texas DIY	Treasure Coast

THANK YOU TO OUR WALK N' ROLL ORGANIZERS WHO VOLUNTEER THEIR TIME!

WALK N'ROLL TO CURE ATAXIA

PASSION FUNDRAISERS

Across the country, passionate volunteers are turning their interests into meaningful support for the Ataxia community. From comedy shows and poker nights to tea parties and community barbecues, these fundraising events raise critical funds, increase awareness, and bring people together in support of NAF's mission. We are grateful for the creativity, generosity, and dedication of everyone who makes these events possible.

HIGHLIGHTS

14 Events

659 Donations

$114K Raised

2025 Events

Artisans for Ataxia

Bowl for a Cure - Wisconsin

Chuck & Duck - Western NY

Cooking with Troy - Ontario

Florida 500 - Jackson

Krewe of Ataxia - Baton Rouge

Pedal for Progress - Berkley

Pickleball Tournament - St. Louis

Serving Up a Cure - Rhode Island

Stand-Up for Ataxia - Atlanta

Stand-Up for Ataxia - Minneapolis

Stand-Up for Ataxia - Philadelphia

XTREME HIKE 2026: GRAND CANYON

HIGHLIGHTS

21 Hikers

$347K Raised

> "Watching my dad's challenges with SCA3 motivates me to do everything I can to push research forward — from participating in studies to taking on new physical challenges. Knowing that I also carry the gene makes this fight deeply personal. The Xtreme Hike is my way of pushing back, showing strength, and turning hope into action."
>
> **-Jessica Oberlin/Hiker**

In 2025, NAF's inaugural Xtreme Hike brought together 21 hikers to take on the challenge of crossing the Grand Canyon in support of the Ataxia community. After months of training, each participant committed to raising at least $10,000 before the hike, and together they surpassed their fundraising goal—raising more than $300,000 to support Ataxia research, education, and community programs. This remarkable achievement was made possible by the hikers, donors, and supporters who rallied behind the event every step of the way. We look forward to building on this success with future Xtreme Hikes.

The Hikers

- David Brunnert
- Lisa Frei
- Joe Frei
- Cathy Gokie
- Jeff Gokie
- Mary Hogan
- Seth Johnson
- Peter Kurachek
- Dana Mauro
- Mark Minkin
- Lauren Moore
- Jessica Oberlin
- Eddie O'Leary
- Pam Perault
- Randy Perault
- Andrew Rosen
- Matt Sefcovic
- Joel Sutherland
- Tanoa Thome
- Karin Westgate
- Craig Westgate

XTREME HIKE 2025
GRAND CANYON SOUTH RIM

FUNDRAISING IMPACT

MARK MINKIN

COMMUNITY MEMBER / XTREME HIKE PARTICIPANT

When Leah Minkin was diagnosed with SCA6, her husband, Mark, turned their determination into action. Together, they have become dedicated champions for the Ataxia community, raising tens of thousands of dollars through local fundraising events, including bowling tournaments and community walks. They also helped lead NAF's inaugural Accelerate! Annual Fund campaign, which has raised nearly $1 million to support NAF's mission.

In 2025, Mark continued that commitment by participating in NAF's inaugural Xtreme Hike, crossing the Grand Canyon after months of training and fundraising. Inspired by Leah and everyone affected by Ataxia, Mark embraced the challenge to raise awareness, inspire others, and help advance research, support, and hope for the entire Ataxia community.

"Ever since my wife was diagnosed with Ataxia 6 years ago, I have made it my mission to raise awareness about this terrible disease while also raising funds for research and education."

REVENUE & PHILANTHROPY

Living with Ataxia is easier when no one has to face it alone. Through education, support, and meaningful connections, NAF's Community Services programs help individuals and families build community, access trusted resources, and navigate their journey with confidence.

5,942

Donors

$11.1M

Donations

$13.2M

Total Revenue

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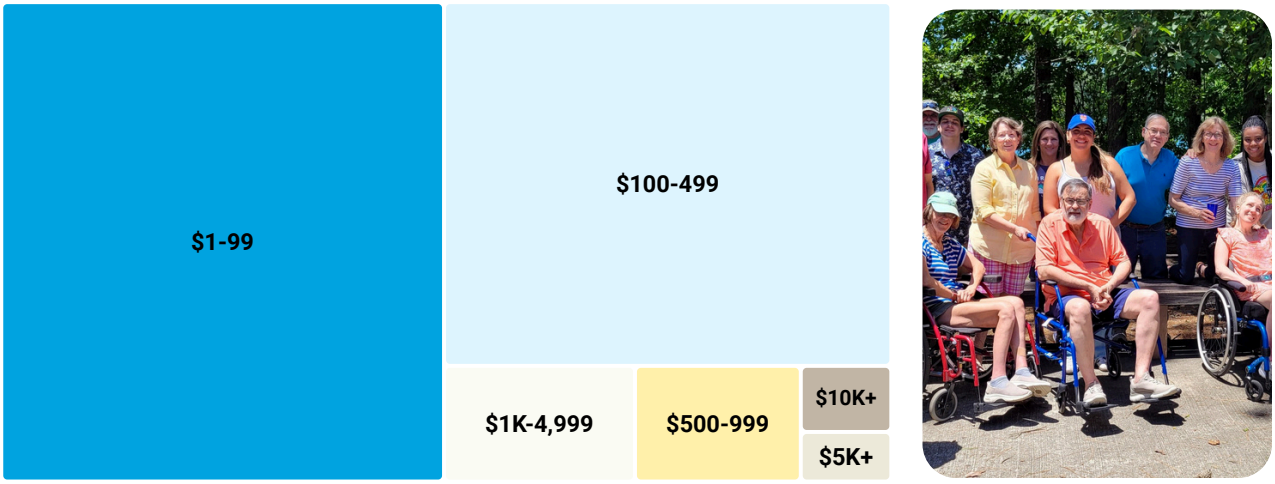
Donations by Gift Size

Progress is powered by people. Every gift—no matter the size—helps advance research, expand support programs, and bring us closer to better treatments and, ultimately, a cure for Ataxia. Together, the generosity of our community creates lasting impact.



CORPORATE PARTNERS

Thank you to our corporate partners for your generosity and commitment in advancing NAF's Mission.

$25K+
Arrowhead Pharmaceuticals
Biogen
Biohaven Pharmaceuticals
Columbia University
Google Inc.
LoQus23 Therapeutics Ltd
Morgan Stanley Gift Fund
PTC Therapeutics
ReproCELL Inc.
Servier Pharmaceuticals

$10K-$24,999
Epic Games
Mass General Brigham
Massachusetts Mutual Life Insurance Company
Microsoft
Mirum Pharmaceuticals, Inc
National Financial Services, LLC
Pershing Advisor Solutions, LLC
Skyhawk Therapeutics
Solaxa Pharmaceuticals
Transcend Medical Communications, LLC
Vanguard Corporation

$10K-$24,999 (continued)
Variantyx
Vico Therapeutics B.V.

$5K-$9,999
Ameriprise Financial
Charter West Bank
Exokinetics, Inc
Goldman Sachs & Co
IntraBio
Lenovo
TIAA
TJX Companies, Inc.
Torres Electrical Supply Co, Inc.

$2,500-$4,999
Abbott Laboratories
CeGaT
Coca-Cola
Farm Bureau Insurance
Los Alamos National Security LLC
Newmark
Tony's Seafood
UBS Financial Services

$2,500-$4,999
Arch Painting Inc
Assured Settlement Solutions

$1K-$2,499 (continued)
Bienvenu Foco Viator
Breadsmith
C3 Presents
Caterpillar Foundation
Cigna
Construction General Laborers & Mtrl Handlers
Cruisin Cajun Country, Inc.
Diane Short Volleyball Inc
Don Vance Auto Group
Exelon Foundation
First National Bank of Omaha
Gartner, Inc.
Greater Horizons
Grossberg Company LLP
Lexeo Therapeutics
Mal Warwick & Associates, Inc.
Material Motion, Inc.
Mobility Works
Rare Patient Voice, LLC.
RunSignUp
Simply Delicious Catering & Management
Theravance Biopharma
Thrivent Choice
UPMC

COMMUNITY GIVING PLATFORMS

We are grateful to everyone who supported NAF through third-party giving platforms. Although individual donor information is not shared with us, we sincerely appreciate every gift.

$25K+

Bank of America Charitable Gift Fund
Benevity Community Fund
Community Foundation for the Land of Lincoln
Fidelity Charitable
Paypal Giving Fund
Raymond James Charitable
Schwab Charitable/DAFGiving 360

$10K-$24,999

Community Foundation of the Holland/Zeeland Area
JPMorgan Chase Foundation

$10K-$24,999 (continued)

Mississippi Ataxia Support Group
Nationwide Life Insurance
The Minneapolis Foundation

$2,500-$4,999

American Endowment Foundation
Burnt Hills - Ballston Lake School District
Chase Online
Renaissance Charitable Foundation
Seattle Foundation
United Life Insurance Company
YourCause

$1K-$2,499

Edward Jones Charitable Gift Fund
E-Trade Morgan Stanley
Give 270, Inc
Hirtle Callaghan Charitable
Jewish Communal Fund
Network for Good
Northern Trust
PSEG Foundation
Transamerica Life Insurance

FOUNDATIONS

We are grateful to the following foundations for their generosity and partnership in advancing NAF's mission.

$25K+

A Stone Legacy Fund
Schmidt Family Fund of the Community Foundation for the Land of Lincoln
Steele Family Foundation
Susan Harris Family Trust
The Clementz Foundation

$10K-$24,999

Don and Jodi Heeringa Family Fund of the Community Foundation of the Holland/Zeeland Area
Haider Quality Freedom Foundation
Leader Family Foundation
Vivian Klassen Giving Fund

$5K-$9,999

Bob & Cappa Woodward Charitable Fund
Buzz & Carolyn Pierce Family Fund
Chip & Susan Carlisle Charitable Fund

$5K-$9,999 (continued)

Donna Harris Family Giving
Everence Foundation Inc
K2 Adventures Foundation
Luthy and Tate Charitable Fund
Mike & Toni Rosen Fund of The Minneapolis Foundation
The Shannon Foundation
Trudy Nye Family Trust
Karen Brown & Michael Peffer Giving Fund

$2,500-$4,999

Alabama Power Foundation
Community Foundation for a Greater Richmond
David C. Copley Foundation
Hankins Family Fund
Leigh and Louise Rabel Foundation at Seattle Foundation
Parker Family Giving Fund
Pledgeling Foundation
The Thomas & Suzanne Harris LEP
Vitrano Family Giving Fund

$1K-$2,499

Allison S. Och Donor Advised Fund
ALMS Fund
Anne & Jeffrey Sternberg Charitable Fund
Ayco Charitable Foundation
Baird Foundation Inc
Brian and Sheila Jellison Family Foundation
Community Health Charities
Cybergrants 3M Foundation
Daniel & Violet Harrell Charitable Fund
DLMC Foundation
Gates Foundation
Hass/Cowan Charitable Gift Fund
Liou Family Fund
Matrix Medical Foundation
Mays Giving Fund
Morris-Goodspeed Giving Fund
National Christian Foundation Carolinas

FOUNDATIONS (CONTINUED)

$1K-$2,499 (continued)

O'Leary Giving Fund
Pamela Graber Donor Advised Account
Paulsen-Hoffman Fund
Peter Skordilis Memorial Fund
Pinekenstein Family Fund
Pinkston Charitable Gift Fund
Potnick Giving Fund

$1K-$2,499 (continued)

Richard James Cohen Giving Fund
Saint Paul & Minnesota Foundation
Ted & Debra Durdel Charitable Fund
The Evelyn S & Jim Horne Hankins Foundation
The Fairlady Fund of the St Paul & Minnesota Foundation

$1K-$2,499 (continued)

The Ho Hing Dai Family Foundation Inc
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INDIVIDUALS

We are grateful to the generous donors whose support makes NAF's mission possible. Their contributions help advance research, expand education and support programs, and bring us closer to better treatments and, ultimately, a cure for Ataxia.

$25K+

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Susan Huffman
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Gregory Klassen
Nina Meister Estate
Marcia Neugebauer
Robert Perkins
Nancy & Rob Schultz
Manoj Shekar
Steven Thomas Jr. Estate

$10K-$24,999

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Lawrence Klotz
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$5K-$9,999 (continued)

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Cherie Nixon
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Tom Risinger
Andrew Schwartz
Sylvia Shelstad
Michael & Nancy Sortum
Marcia Stewart

INDIVIDUALS

$2,500-$4,999 (continued)

Bryan Tabery
Karen Whittle
Paul Wilner

$1K-$2,499

Lawrence Adair
Katharine Adams
Marcie Anthone
Chris Arnold
Bob & Virginia Atwell
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$1K-$2,499 (continued)

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Christopher Hodges
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Richard Lavery
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$1K-$2,499 (continued)

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Anthony & Robyn Zeller

ADVANCING TOWARD A CURE

Every gift to NAF helps move Ataxia research and support services forward. Through careful stewardship and strategic investment, we maximize the impact of every dollar entrusted to us. The summary below shows how your generosity is helping accelerate research, strengthen our community, and move us closer to better treatments—and ultimately, a cure.

Support & Revenue

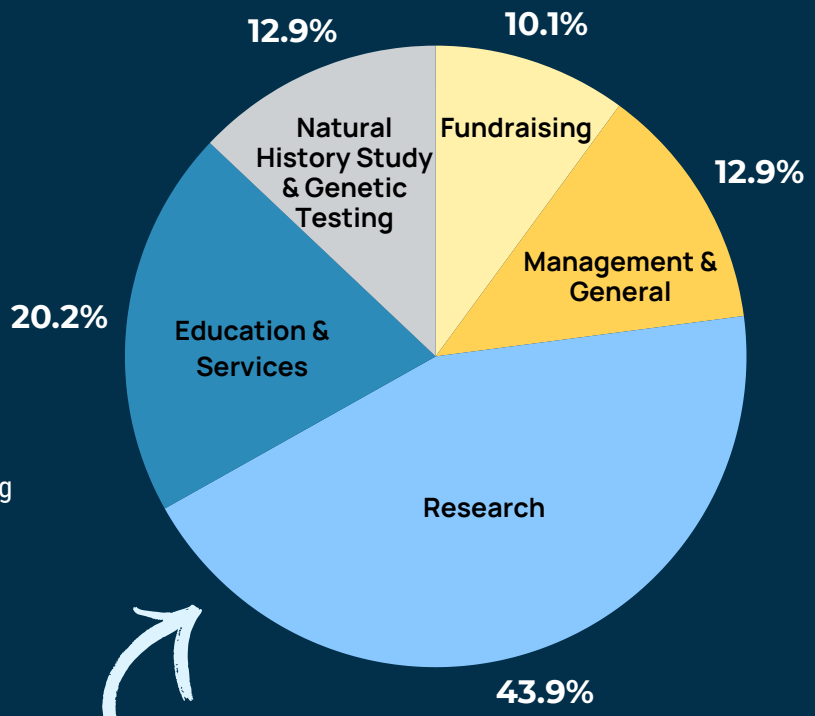
- Contributions: $11,093,572
- Drug Development Collaborative: $897,000
- Conference Income: $653,080
- Other Income: $520,588

Expenses

- Research: $2,745,747
- Education and Services: $1,262,194
- CRC-SCA Natural History Study & Genetic Testing Programs: $809,201
- Management and General: $803,917
- Fundraising: $628,750

Net Assets

- Change in Net Assets: $6,914,432
- Net Assets, beginning: $4,307,152
- Net Assets, ending: $11,221,584
- Total Liabilities: $630,748
- Total Assets: $11,852,332



HOW WE DELIVER ON OUR MISSION

NAF
National Ataxia Foundation

Organization Leadership

NAF STAFF

Senior Leadership

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Andrew Rosen, MBA
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Chief Development Officer
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Meet the Team

Sr. Research Coordinator
Aimee Alcott

Senior Accountant
Sue Baker

Community Services Associate
Courtney Bockhop

Community Services Coordinator
Kari Brooks

Community Services Manager
Barbara Cox

Sr. Clinical Research Coordinator
Laura Crespo

Communications Coordinator
Derick Franklin

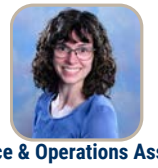
Events Manager
Marianne Johnson

Development Manager
Rich McCutchen

Controller
Patricia Morel

Communications Associate
Mary Ann Peterson

Research Associate
Carrie Taylor



Finance & Operations Associate
Julia Taylor

Clinical Services Director
Kelsey Trace

Patient Engagement Director
Celeste Suart, PhD

VP, Development
Joel Sutherland

Communications Manager
Mollie Utting

Development Manager
Jon Wegman

Communications Associate
Hannah Weldon

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NAF's Board of Directors provide strategic leadership and governance to help advance our mission. Through thoughtful oversight, long-term planning, and responsible stewardship of resources, the Board helps ensure NAF remains financially strong and well-positioned to serve the Ataxia community.

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NAF's Medical and Research Advisory Board brings together leading experts in Ataxia research and clinical care. Their knowledge and guidance help inform our research priorities, educational programs, and strategic initiatives, ensuring our work is grounded in the latest scientific and medical advances.

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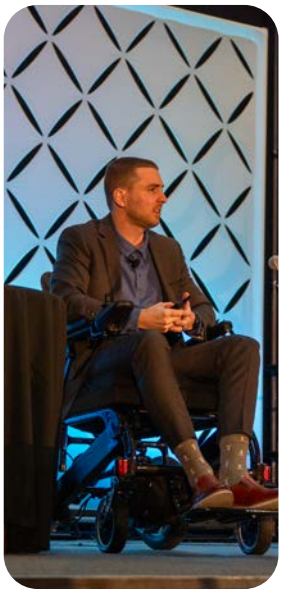
THANK YOU

As we look back on another year of progress, we are grateful to everyone who makes NAF's mission possible. To our donors, volunteers, healthcare professionals, researchers, advocates, and members—thank you. Your generosity, expertise, and commitment help advance research, expand support and education programs, and bring hope to individuals and families affected by Ataxia. Every contribution strengthens our community and moves us closer to a future with better treatments and, ultimately, a cure. We look forward to continuing this journey together in the year ahead.



National Ataxia Foundation

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