angelman SYNDROME FOUNDATION

Fall 2026

FAMILY NEWSLETTER

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A MESSAGE

from Our CEO and Board Chair

Dear Angelman Community,



As we look back on this past year, one theme keeps rising to the surface: Lift Them Up.

We saw it in the parent who drove hours for a clinic appointment because access to expert care mattered enough to make the trip. We saw it in the sibling who spoke up on Capitol Hill, becoming an advocate before they were old enough to drive. We saw it in every donor who gave, and gave again, because they believed this community was worth lifting.

This year, ASF grew in record ways: our largest Family Conference yet, an expanding clinic network, deepened family support programs reaching more families than ever before, and a growing LADDER database.

This year also asked something harder of us. The news from the ASPIRE trial was a profound disappointment. To every family who enrolled a loved one in that trial, who gave your time, trust, and hope: we see you, and we are endlessly grateful. Your courage lifted this field forward, even when the outcome wasn't the one we all prayed for.

If this year taught us anything, it's that lifting each other up was never contingent on a single trial or headline. It lives in the parents who show up to every walk, the siblings who become lifelong advocates, the researchers and clinicians who keep pushing forward, and the staff and board who build an organization capable of carrying this mission for decades to come.
As you turn through this newsletter, we hope you feel it: every story and every milestone here is a small act of lifting someone up, and together, this community lifted higher than ever this year.

We are so proud of you, and so grateful for you. Here's to lifting each other higher still in the year ahead.

Amanda Moore
CEO, Angelman Syndrome Foundation
Mom to Jackson (age 9, AS deletion)

John Sugden
Board Chair, Angelman Syndrome Foundation
Father to Sebastian

Angelman Strong:

Your Community. Your Event. Your Impact.



Angelman Strong is more than a walk or 5K. It's an opportunity for the Angelman syndrome community to come together, raise awareness, build connections and support ASF's mission.

And what a year 2026 has been! More than 9,000 community members registered for Angelman Strong events across the country, raising over $1.53 MILLION! Thanks to every participant, donor, volunteer and event coordinator who helped make this incredible impact possible. Funds raised support critical research, programs and resources for the Angelman community.

Interested in Getting Involved?

Angelman Strong events don't have to be a traditional walk or 5K. You can create an event that reflects what you and your community are passionate about. A cornhole tournament, car show, pickleball tournament, family fun day, trivia night . . . . there are endless possibilities!

And you don't have to figure it all out on your own. ASF will be there with resources, guidance, ideas, and support to help you bring your event to life. Whether you are an experienced event planner or simply someone who wants to create something meaningful in honor of a loved one with Angelman syndrome, **there is a place for you in Angelman Strong.**

Tell us a little about yourself and your idea by completing our interest form:
Angelman Strong 2027 Host Interest Form

To learn more about Angelman Strong and find an event near you, visit Angelman.org/Strong.

Turn What You Love Into Impact

Host an ASF Community Event

Our community has gotten incredibly creative! ASF Community Events have included cornhole tournaments, golf outings, fashion shows, galas, block parties, bar crawls, trivia nights, game nights, wine nights, runs, walks, car shows, gymnastics tournaments, pickleball tournaments, bowling events, benefit concerts, restaurant nights, bake sales, community festivals, and so much more. Have a completely different idea? We want to hear it—the possibilities really are endless!

We are incredibly grateful to the many dedicated fundraisers who give their time, energy, and creativity to host events benefiting ASF. Thank you for turning your passion into impact!

Sutton Strong
The annual Sutton Strong Golf Outing in Macomb, MI, hosted by Nicole Smith in honor of Sutton, raised $17,000 to support ASF and the Angelman syndrome community.

Infinity Ice Invitational
Infinity Athletics raised $3,200 in their second year hosting the Infinity Ice Invitational gymnastics competition in Hopewell Junction, NY, benefiting ASF.

Chris Hanson | Microsoft Give Campaign
Chris raised over $10,000 for ASF by offering professional headshots to colleagues in exchange for donations, with additional support from volunteer hour matching and Microsoft's company match.

BMX Race
Tiffany Hartnett and the Saber Racing Team raised $1,250 for ASF through a BMX race in Port Orchard, WA.







Every event—whether it brings together 25 people or 500—creates another opportunity to introduce people to Angelman syndrome, strengthen connections, and support the programs, resources, and research that matter to our community.

Have an idea? Start with what you love and make it count. Complete our Community Event Interest Form and tell us what you're thinking:
ASF Community Event Interest Form

Miles With a Mission

Meet Our ASF Running Teams



There is something pretty incredible about crossing a finish line knowing every mile has helped move the ASF mission forward. We have **two amazing teams** getting ready to do just that!

Celebrating 10 Years of Windy City Angels!

On October 11, 2026, 68 Windy City Angels (WCA) took on the 2026 Bank of America Chicago Marathon, celebrating 10 years of ASF participation in this iconic race! Together, these incredible runners represented the Angelman syndrome community through 26.2 miles of Chicago streets, raising awareness and over $300,000 in critical funds to support ASF's mission.

Recruitment for the 2027 team has begun. If running the Chicago Marathon is on your bucket list, complete our interest form to receive information about joining our team.

Windy City Angels 2027 Interest Form

The Magic Begins in 2027!

We are also counting down to the inaugural year of our newest running team, the Magical Miles Angels (MMA)! In February 2027, 100 runners will represent ASF at Walt Disney World during runDisney Princess Half Marathon Weekend, participating in the 5K, 10K, Half Marathon, and Fairytale Challenge.

The response to this new team has been incredible, and our 100-member Magical Miles Angels team is currently full! We can't wait to watch our runners train, fundraise, support one another, and bring the ASF community and mission along for every magical mile.

Be Part of the Magic as a Partner

While our running teams are full, there is still another way to join us! We are currently looking for corporate and community partners who want to be part of the magic and support our Windy City Angels and Magical Miles Angels teams.

Partnership provides a unique opportunity to stand alongside 168 runners collectively representing ASF while helping make every mile even more meaningful. Companies and organizations interested in supporting our inaugural Magical Miles Angels team can learn more about the available partnership opportunities here:

Magical Miles Angels Partnership Agreement

To our 68 Windy City Angels and 100 Magical Miles Angels runners: thank you for choosing to put your miles behind the ASF mission. From Chicago to Walt Disney World, we will be cheering for you, celebrating you, and sharing the incredible impact you are making with every step!



There’s a Place for You at ASF

Volunteer at an event. Become an advocate. Serve on a committee. Help with a special project.

Whether you have a few hours to give, a special skill to share, or an idea you’re excited to bring to life, your involvement could have a meaningful impact on the Angelman community.

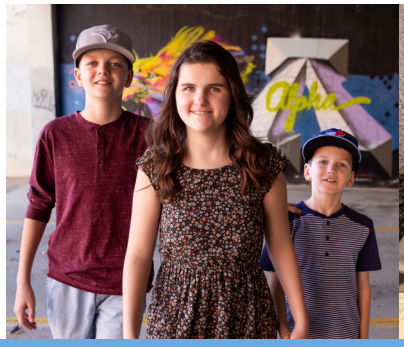
There are opportunities to get involved locally and nationally and you can find something that fits your interests and availability. Volunteers are an important part of what makes what we do possible.

Ready to explore the possibilities?

Visit the Make an Impact section of our website to learn more about current volunteer opportunities and the many ways you can put your time and talents to work for ASF: Explore ASF Volunteer & Get Involved Opportunities

angelman SYNDROME FOUNDATION

nextgen







Introducing ASF NextGen: Lead. Create. Impact.

When we think about the future of the Angelman Syndrome Foundation, we know that siblings have an important role to play—and ASF NextGen is giving them a place to make that role their own. We're excited to introduce ASF NextGen, our newest initiative designed to engage and empower siblings ages 15–22. Here, the next generation of Angelman supporters can explore how their individual interests, talents, and experiences can make a difference within the ASF community.

ASF NextGen isn't about telling siblings how they should be involved. It's about helping them discover how they want to be involved. Whether they are passionate about advocacy, fundraising, volunteering, raising awareness, or creating something entirely new, we want to give them the tools and opportunities to turn their ideas into action.

If you're between the ages of 15 and 22 and looking to make an impact within the Angelman community, please reach out to Alisa Tammaro atammaro@angelman.org to learn more about becoming involved with ASF NextGen. We'd love for you to find a way to use your voice, interests, and talents to make a difference.

The next generation is ready to make its mark.
Lead. Create. Impact.
ASF NextGen.

Adulthood & Transition Resources

Committed To Supporting Adults & Their Caregivers

If you care for an adult with AS, having a strong support system will help you overcome challenges. Each month, our Adult Task Force meets to discuss ways we can try to help enrich the lives of adults living with AS. We recognize that the adult caregivers have paved the way for all our families, and it's part of our mission to help provide support and assistance.

AS Grows Up

The Angelman Syndrome Foundation is committed to supporting every individual, every step of the way. As individuals with Angelman syndrome age, new needs arise for housing, programs, community, care, and more. We are taking action to close the gaps in essential support and expand the vital resources and services families desperately need. Watch for more and expanded resources for adults as we continue to work to close the gap.

Visit Angelman.org/asgrowsup

ASF Family Champions with Adults

ASF Family Champions are family members who have volunteered to be available to other families to share their knowledge, experience and lend an ear. There is a group of champions who are parents of adults with AS. These families are a great resource for questions, concerns and advice on various aspects of adults with AS transitioning and thriving.

Visit Angelman.org/resources/asf-family-champions/#adult-champs

Adult Housing Champions

Adults with Angelman syndrome live in various settings. These include homes of their own, living with family members or host families, and provider-owned or -operated settings. The Adult Housing Champions have navigated living situations for their loved one and are available to guide and share their experience with others.

Visit Angelman.org/adult-housing-champions

Future Care Planning Series with Protected Tomorrows

Planning for the future can feel like standing at the edge of a very big map with no clear starting point.
The Future Care Planning Series is designed to give you a place to begin, a path to follow, and the confidence to move forward.

Through a four-part series, you'll gain the tools and clarity needed to start building a Future Care Plan that reflects your family's unique needs, priorities, and goals.

Whether you're just starting to think about long-term planning or ready to take action, this series is built to meet you where you are and guide you step by step.

Visit Angelman.org/resources/future-care-planning

Behavior Resources

With AS comes unique and challenging behaviors and increasing anxiety. ASF is happy to offer assistance and information about dealing with and understanding these behaviors, at no cost to you.

Visit Angelman.org/resources/filter/meta/topics!is_-custom_checkbox:Behavior

Printable Resources

The ASF Adult Task Force created several documents that are beneficial for situations that could arise with your adult with AS. Examples include Transition to Adulthood, Adult FAQs, Comprehensive Care Binder, Facility Tour Checklist, All About Me Template and more.

Visit Agelman.org/asgrowsup

Research Recap

Where Things Stand with Clinical Trials

Even in a season marked by disappointment, the pursuit of better treatments for Angelman syndrome hasn't slowed, it's evolving, and the community's resilience continues to drive this field forward. Here's where things stand.

The Angelman research landscape has shifted significantly. Ultragenyx's Phase 3 ASPIRE trial of GTX-102 (apazunersen) has concluded, and the company announced the trial did not meet its primary endpoint. Ultragenyx has since discontinued development of GTX-102 and its Angelman syndrome program entirely, ending dosing as of late October.

IONIS

Meanwhile, two other ASO therapies remain in active development. Ionis Pharmaceuticals' obudanersen (ION582) is progressing through its Phase 3 REVEAL study: the pediatric cohort (ages 2–<18) finished enrollment this past July with 136 participants, and the adult cohort (ages 18–50) is expected to complete enrollment by the end of this year. Topline results are expected in the second half of 2027. REVEAL's primary endpoint focuses on improvement in expressive communication. Ionis also plans to launch a related trial, CHAMPION, before the end of 2026, specifically for individuals with UPD or imprinting-defect genotypes.

Oak Hill Bio

Oak Hill Bio dosed its first participant this past July in BEACON, its own Phase 3 trial of rugonersen (an ASO originally developed by Roche, building on encouraging early results from the Phase 1 TANGELO trial). BEACON's primary endpoint looks at improvement in cognition and/or expressive communication, and the company expects topline results in early 2029.



Skylark Bio

Just this month, Skylark Bio, a clinical-stage biotechnology company, announced the addition of SKY-AS to its pipeline. SKY-AS is an investigational gene therapy designed to deliver a functional copy of the UBE3A gene to neurons in hopes of restoring UBE3A protein expression. This program is currently advancing through IND-enabling studies.

LADDER Database
Linking Angelman & Dup15q Data for Expanded Research

Behind these headline trials, quieter but essential work continues. Natural history studies keep building a clearer picture of how Angelman syndrome unfolds over time, and ASF's LADDER database continues connecting data from clinics and registries so future trials can move faster and reach more families.

But one trial does not define the future of Angelman syndrome research and treatment. The data collected through ASPIRE will add to the growing body of knowledge about the biology of Angelman syndrome, clinical trial design, and how potential treatments are evaluated. That information can help researchers ask better questions and design stronger trials in the future, including for the two other ASO therapies, from Ionis and Oak Hill Bio, that remain actively in development today.

Newly Diagnosed Corner

Newly Diagnosed? You Are Not Alone.

If you've recently received an Angelman syndrome diagnosis, we want you to know—you are not alone. While this news can feel overwhelming, it's also the beginning of a journey filled with support, connection, and hope. The Angelman Syndrome Foundation is here to walk alongside you with resources, information, and a community that understands.

Our close-knit community is made up of people who truly get it—parents, caregivers, medical professionals, and advocates who are all working together to make life better for individuals with Angelman syndrome. We invite you to explore, ask questions, and lean on us for help. Whether you're looking for medical guidance, educational tools, or just someone who understands, we're here for you.

FIND NEWLY DIAGNOSED RESOURCES

FIND NEWLY DIAGNOSED RESOURCES:

The First 100 Days Journey

The First 100 Days Journey is a guide, which was created by Angelman parents and caregivers who have walked this path before you, to help you navigate this new diagnosis. Let us stress that you do not have to do everything on this list. Remember to give yourself grace as you are processing the road that lies ahead. This is simply a support tool to help you start to think about important steps you can take to help your child thrive and the resources available to assist you along the way.

AS 101 Webinars

These informative online events are hosted every other month by our CEO, Family Resource Director and Chief Medical Officer. We encourage you to attend and invite family members. These webinars offer a wealth of information that can help guide you during this time.

Family Champions

Connect with others who understand. Please reach out to our Family Champions in your area! If at any point you would like to connect one-on-one or have any questions, please reach out to Kathryn Ely at kely@angelman.org

For more information visit: Angelman.org/newly-diagnosed

ASF Family Support Funds

Every Angelman family has different needs, challenges, dreams, and goals. That's why there are three Family Support Funds designed to provide meaningful support in different ways.

Whether you need help getting a medical device, creating time to rest and recharge, or having an experience that can make a lasting impact, there's a support fund that can help.

Applications are open annually October 1–14.

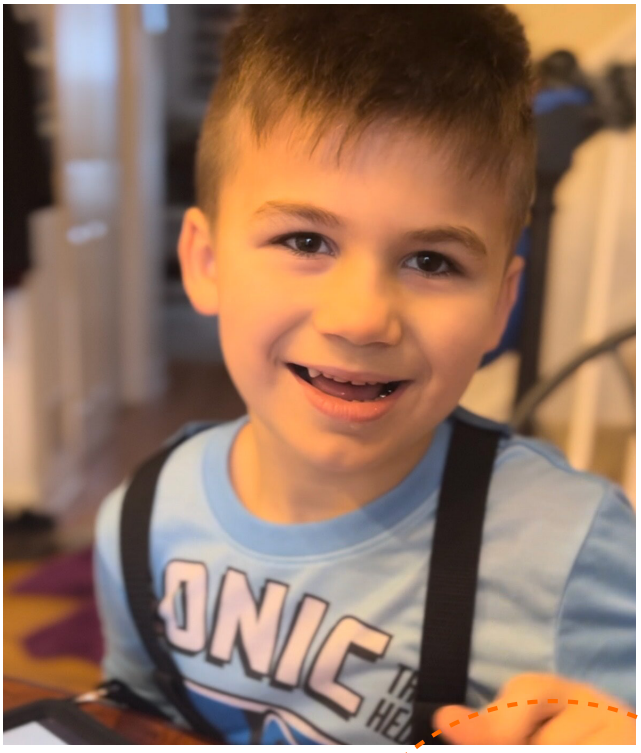
ASF Family Fund
Apply for funding to gain access to something that will improve the quality of life of an individual with AS. Commonly funded items include: Safety beds, iPads, adaptive bikes & strollers, medical equipment, travel costs to an ASF Clinic, and various therapies.

ASF Respite Fund
The Respite Fund provides financial support to help create a meaningful respite opportunity for caregivers. Funding can allow you to step away from your caregiving responsibilities, spend time with your partner or other children, pursue something that brings you joy, or simply take time to rest and recharge.

Sophie Burdine Schwartz Impact Award
This fund is meant to bring joy-driven experiences and everyday pleasures. Apply for something that makes a meaningful impact in the life of your individual with Angelman syndrome. Not a clinical need, but something that would make their life happier.

Find details, eligibility criteria, funding amounts, FAQs and more on our website: Angelman.org/resources/family-support-funds





2026 ASF Family Conference and Research Symposium Recap



ASFConference.org

Living with Angelman syndrome brings unique challenges and questions at every stage of life. Because AS is rare, finding others who truly understand can be difficult.

The 2026 ASF Family Conference & Research Symposium brought together nearly 1,600 attendees from 45 states and 14 countries, making it the largest gathering of the Angelman syndrome community in the world.

From newly diagnosed families to those who have navigated Angelman syndrome for decades, attendees came together for four days of learning, connection, resources and hope for the future.

That is what the ASF Family Conference is built to provide.

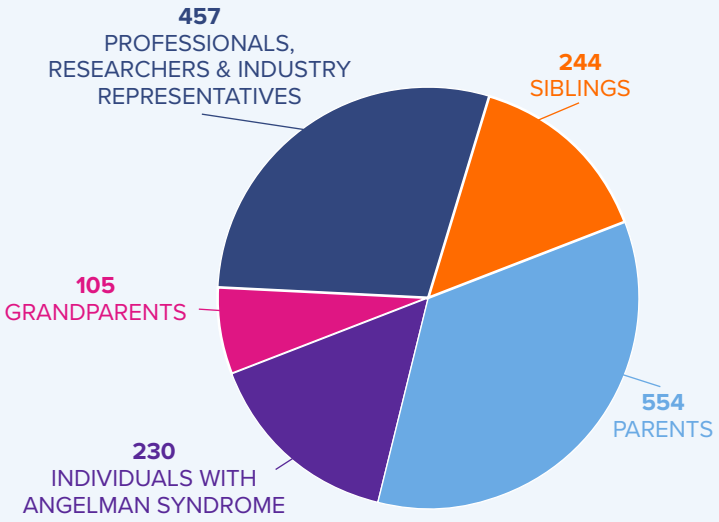
During the conference, ASF hosted 170 opportunities for learning, consultation, connection and community. Families learned from 117 speakers, sat down one-on-one with experts through private consultations, met 43 exhibitors offering resources across the full spectrum of life with AS, and spent time with other people who understand their lives.

I would tell someone else going to the Conference to...

"...play and make friends. And learn, actually."

FELIX (OKLAHOMA, AGE 6)
BROTHER TO VIVIAN, AGE 10, LIVING WITH ANGELMAN SYNDROME





1,590 Total Attendees

117 Speakers
including researchers, clinicians, professionals, advocates and community voices

43 Exhibitors
connected families to resources across the full spectrum of life with Angelman syndrome.

12 Disability Service Providers
13 Product & Technology Solutions
7 Family Support & Resource Representatives
11 Clinical care and research organizations

7 SIBLING ACTIVITIES

13 SOCIAL GATHERINGS

47 PRIVATE CONSULTATIONS

170 PROGRAMMED LEARNING, CONSULTATION, CONNECTION & COMMUNITY EXPERIENCES.

And we know those experiences made a difference. Eighty-eight percent of surveyed families said they learned something they could apply at home, 75% left feeling more confident caring for their family member with AS, and 92% left with a stronger sense of belonging in the Angelman community.

But some of the most important things that happen when we bring this many people together cannot be captured in a schedule.

Families learn from experts. Experts learn from families. Researchers find collaborators. Organizations connect directly with the people they are working to serve. Families find other families who understand. And conversations that may never have happened otherwise can begin.

Thirty partners helped make all of this possible, representing research and therapeutic development, products and technology, services and family support, and families and individuals who believe in ASF's mission and the importance of bringing this community together. Their investment helped us create something that is difficult to replicate anywhere else.

- A place where a family can learn about the latest Angelman research in the morning and ask a clinician a personal question that afternoon.
- A place where a parent can discover a product, service or resource they didn't know existed.
- A place where siblings are included and find people who understand their experience.
- A place where researchers, clinicians and families share the same hallways, meals and conversations.
- A place where a family who has never met another person with Angelman syndrome can suddenly find themselves surrounded by hundreds of people who understand.



"Coming back home from the Angelman conference, I can honestly say something in me changed.

For years, I loved my daughter with everything in me, advocated for her, cared for her, fought for her, and celebrated every little win. I never gave up on her. But somewhere along the way, I think I accepted certain limits without even realizing it.

Then I went to this conference.

I saw other children and adults with Angelman syndrome doing things I had never personally seen before. Communicating. Learning. Becoming more independent. Using their AAC devices. Participating in their communities. Growing, progressing and surprising the people around them.

And all I could think was... why not my baby too?

Not because every child with Angelman is the same, because they aren't. Their journeys are all different. But for the first time in a long time, I stopped focusing so much on what my daughter might never do and started thinking about everything she still has the opportunity to learn.

I came home with HOPE.

And I came home with a completely different mindset.
We're removing the limits and we're going to see just how far my girl can soar."

GABRIELLE, MOM, MISSOURI

And we are still listening. Families told us they want more support around adulthood and independence, behavior and anxiety, family relationships, siblings and caregiver well-being. They want more opportunities to connect with one another. What we learned in 2026 will help us build an even stronger experience for the community in 2028.

Families came looking for answers. They went home with information they could use, resources they could access, experts they could turn to, people they could call, greater confidence in caring for someone they love, and a stronger sense that they belong to a community that will continue walking alongside them.

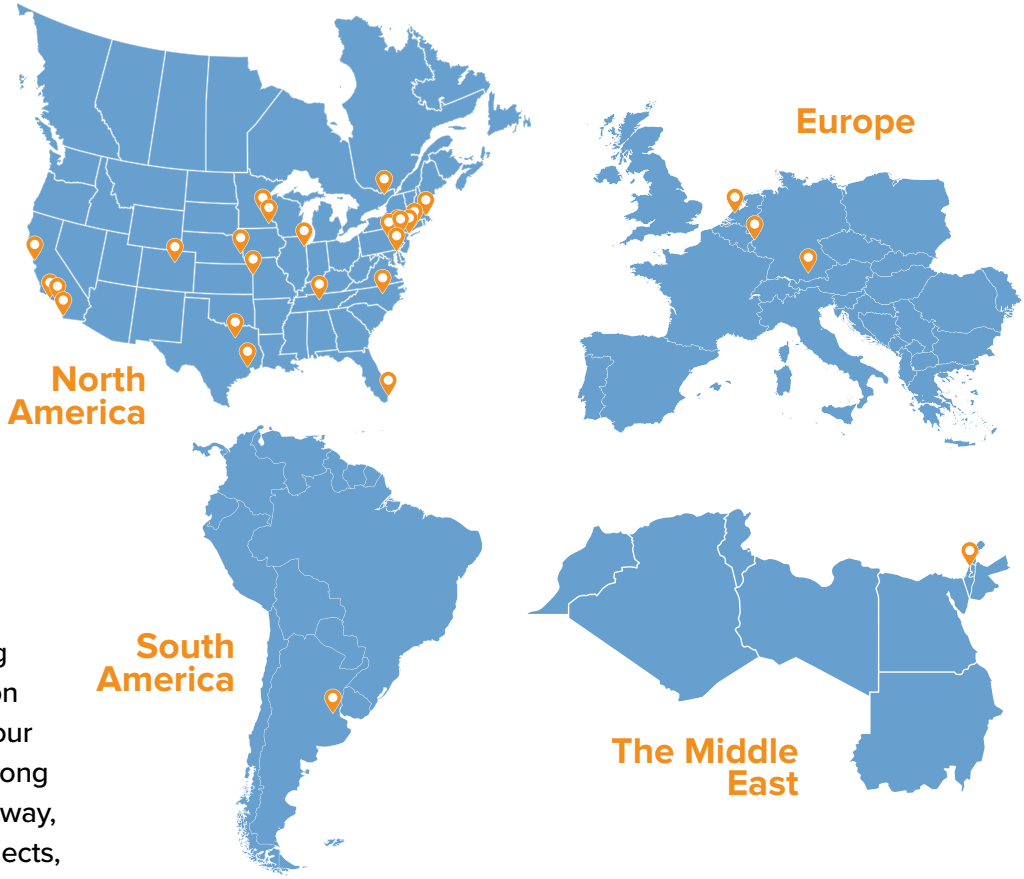
That is why we bring the Angelman community together. The conference may last four days. Its impact lasts much longer.

Save the Date!

We hope you will be part of it in June 2028 at the new Kalahari Resort in Woodford, VA.

View the photo gallery from the 2026 ASF Family Conference.

ASF Clinics Update



To learn more visit: angelman.org/as-clinics

The ASF is dedicated to increasing access to expert Angelman care around the world.
Our Clinical Network has seen some incredible growth over the last few years, and 2026 has been no different. This year, we have focused on assessing our resources, gathering feedback on what is working—*and what isn't*—for our families and clinicians, and building strong foundations for the future. Along the way, we have also created new projects, platforms, and pathways to make it easier for clinicians and caregivers to connect with expert Angelman care. Read the updates below to learn more about a few key achievements this year and some of our plans for the future.

Increasing Access to Expert Angelman Care
Please join us in welcoming new and returning teams to the ASF Clinical Network!

UCLA has officially reopened its Chromosome 15 Clinic, kicking off the return of the program with an opening ceremony on August 30. Families interested in making an appointment or learning more can contact Chromosome15clinic@mednet.ucla.edu.
In Texas, UTSW has expanded its Angelman care at Children's Health Dallas with the opening of an adult care transition clinic led by Dr. Irfan Sheikh. This new service expands access to expert Angelman care across the lifespan for families in the Dallas/Fort Worth area.

We are also working with an amazing team at Seattle Children's Hospital to meet the care needs of families in the US Northwest Region. Stay tuned for more information as we prepare to officially launch this new clinic site in early 2027!

Expanding Care Across the Globe
Aside from welcoming many new faces into our clinical network around the world, including teams in Tbilisi, Georgia and Prague, Czech Republic–the ASF has also made important progress in strengthening our International Clinical Network by finalizing the structure of ASF Europe. This partnership brings our network of clinic sites, industry partners, and advocacy organizations across Europe together under the leadership of the Erasmus team in the Netherlands.

Find an ASF Clinic Near You
Did you know the ASF Clinical Network has an online clinic map? Visit the ASF Clinic Map to explore our network of multidisciplinary clinics and find expert Angelman care in your area. You can browse the map or

ASF Clinics Update, cont.

use the search tool to locate a clinic by region. We are continually adding clinics as they come online, making this an easy, self-serve resource for families looking for multidisciplinary care.

More Than Clinics: Bringing Expert Care to You
Expert Angelman care doesn't only happen within multidisciplinary clinics. The ASF Provider Registry recognizes individual healthcare providers, specialty providers, and private practices with an interest in and expertise in Angelman syndrome who practice outside of an established ASF clinic.

Help us identify experts in your area! If you know a healthcare provider or specialty practice that has experience caring for individuals with Angelman syndrome, you can nominate them for the ASF Provider Registry. Nominations are reviewed by the ASF Clinical Programs team, and providers who meet program criteria will be invited to complete a formal application to join our network.





Expanding Clinical Expertise Through ASF ECHO
ASF ECHO is a virtual, case-based learning program that connects clinicians around the world with Angelman syndrome experts to expand knowledge, identify gaps or barriers, and improve care. Through live expert-led educational presentations and real-world case discussions, clinicians can learn from AS specialists while connecting with other providers learning to care for individuals with Angelman syndrome.

Our ongoing series, Caring for Adults with Angelman syndrome, focuses on the unique clinical considerations of caring for adults with AS and is facilitated by Wilfreda Lindsey, MD, Director of the Angelman Syndrome Clinic at Kennedy Krieger Institute. Two sessions have already taken place.

Looking ahead, ASF ECHO will also feature series focused on speech-language pathology & communication and Angelman syndrome genetics in early 2027. Learn more about ASF ECHO and request to join the invitation list by clicking here.

BEACON

THE PHASE 3 STUDY OF RUGONERSEN FOR ANGELMAN SYNDROME

Advancing Rare Disease Therapies Across the Finish Line

New hope for families living with Angelman syndrome

A clinical study that could make a difference

BEACON is testing rugonersen, an investigational medicine that targets the genetic root cause of Angelman syndrome — with the potential to improve communication, cognition, and daily life.

Ask your doctor if BEACON may be right for your loved one.

Oak Hill Bio is committed to carrying rugonersen forward — a promising investigational product that could change the course of Angelman syndrome.

Oak Hill Bio works alongside families, researchers, and the Angelman community to give rugonersen the chance it deserves.



www.oakhillbio.com

Oak Hill Bio

Proud partner of the Angelman Syndrome Foundation

Rugonersen is investigational and not approved by any regulatory authority.

OAK HILL BIO · OHB-724-301

Going beyond every day.®

Ultragenyx is honored to sponsor the 2026 Angelman Syndrome Foundation's Research Symposium and Family Conference, and stands proudly with the Angelman syndrome (AS) community.

Together with ASF, we are committed to advancing AS research that is grounded in science and insights from families.

We are grateful for our partnership with ASF as we support individuals, family members, and loved ones living with and caring for those with Angelman.

Learn more at **Ultragenyx.com**

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Scan to get started

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safeplacebedding.com

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Thank You To Colin & Kellie Sargent and Maddie's Mission for supporting the 2026 ASF sibling programs.

Maddie's Mission Foundation

"Turning Can't Into Can - One Miracle at a Time"



Our science is creating an unprecedented opportunity **to change the course of human health.**

Ionis is proud to sponsor the 2026 ASF Research Symposium and Family Conference. For more than three decades we've united groundbreaking science and technology with relentless passion to discover and deliver medicines that enable better futures for people living with serious diseases. Learn about our RNA-targeted medicines at **ionis.com**.

angelman SYNDROME FOUNDATION

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JOIN OUR COMMUNITY

DONATE NOW

The mission of the Angelman Syndrome Foundation is to advance the awareness and treatment of Angelman syndrome through education and information, research and support for individuals with Angelman syndrome, their families and other concerned parties. We exist to give all of them a reason to smile, with the ultimate goal of finding a cure.

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