

Brunch in Blue

...for the Angelman Syndrome Crew!

The Foundation for Angelman Syndrome Therapeutics, in partnership with Will.i.CAN, invites you to sponsor our fundraiser! Critical research funds will be raised so all individuals around the world living with Angelman syndrome (AS), a rare neurogenetic disorder, can have their shot at a better future!

Sunday, October 11, 2026 from 10am-1pm @ CHS Field, Downtown St. Paul, MN

Premium Event Packages	Summit	Highland	West 7th	Midway
	\$10,000	\$5,000	\$2,500	\$1,000
Priority				
First right of refusal for 2027	X	X	X	X
Day of Event				
Complimentary event tickets	12	10	8	6
Reserved table seating	12 seats	10 seats	8 seats	6 seats
Heads Up Game Plays	12	8	4	2
Raffle Basket Tickets	150	100	50	20
Brand Recognition				
Logo listed on scrolling video screen during event	X	X	X	X
Logo on event registration website	X	X	X	X

Additional Options:

\$650: The Frogtown Table for 6 (four available)

Come with 6 of your friends and secure reserved seating!

\$500: The Phalen General Sponsor

- 4 event tickets
- Logo listed on event registration website

The Foundation for Angelman Syndrome Therapeutics (FAST) is a not-for-profit organization and is exempt from federal income tax under section 501(a) of the Internal Revenue Code as an organization described in section 501(c)(3), Federal Identification Number: 26-3160079.

If Will CAN, We CAN ...

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On October 11, 2026, the Foundation for Angelman Syndrome Therapeutics (FAST) in partnership with Will.i.CAN, will be hosting our 4th Annual fundraiser at CHS Field in Saint Paul, to raise critical research dollars to help drive forward promising therapeutics for Angelman syndrome (AS).

This fundraiser is growing each year, bringing more families around Minnesota impacted by Angelman syndrome together, so we can fight for our loved ones and all individuals living with AS around the world.

Since 2020, between peer-to-peer fundraising and annual events, Will.i.CAN's crew, with the help of other AS families in Minnesota, has raised over \$1 MILLION to benefit FAST– but we will not stop until there are approved therapeutics for all individuals living with Angelman syndrome.

Will You Help Us?

This packet will help you understand how these critical funds will support FAST and the Angelman syndrome community. We are excited to offer a variety of sponsorship opportunities in hopes there is something that fits your level of interest.

We are also seeking in-kind donations such as unique or high end alcoholic bottles (liquor, beer, wine, other), gift cards ranging from \$20-\$250, and raffle basket items and experiences.

Thank you in advance for your consideration!

Purchase a Sponsorship

Scan QR code or tap [HERE](#).

In-Kind Donation Offers

Email Meghan at will.i.can.with.angelman@gmail.com





FOUNDATION FOR
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THERAPEUTICS

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Sponsorship & In-Kind Donation Letter

The Foundation for Angelman Syndrome Therapeutics (501(c)3 non-profit, Tax ID: 26-3160079), in partnership with Edberg Family of Rosemount, MN, invites you to support our 2026 fundraiser where we will raise awareness of Angelman syndrome (AS) and funds to progress critical research aiming to one day cure Angelman syndrome and a host of other neurogenetic disorders. This event will be held on **Sunday, October 11, 2026 at CHS Field, Downtown St. Paul from 10am-1pm.**

Angelman syndrome (AS) is a rare neurogenetic disorder that affects about one in 15,000 people, or approximately 500,000 individuals worldwide. Some say it could be as frequent as one in 8,000 people, or over 1,000,000 individuals worldwide.

People living with AS have a genetic difference on chromosome 15. People with these chromosomal changes do not produce any, or remotely enough, of something called UBE3A —which is a protein involved in brain functioning that helps us walk, talk and perform many other everyday tasks.

As a result, children and adults with AS typically have balance issues, motor impairments and can have debilitating seizures. Disruptive sleep is often a serious challenge. Some never walk. Most do not speak. Individuals with AS have a normal life expectancy but require continuous care and are unable to live independently.

This is life today for people living with Angelman syndrome. But it won't be for long.

“Scientists believe that AS is now the neurogenetic disorder with the greatest potential to be cured. FAST is the reason why, and the force making it happen.”

FAST is the leading patient advocacy organization working to cure Angelman syndrome. As the largest non-governmental funder of Angelman syndrome research in the world, our goal is to drive forward transformative research and development programs as quickly as possible for those living with Angelman syndrome — regardless of age or genotype.

As of March 2026, there are over 23+ therapeutic programs in the [Angelman Syndrome Drug Development Pipeline](#), with 5 potential therapeutics currently have ongoing clinical trials.

The cost of drug development is extremely high and often falls on the shoulders of patient advocacy groups like FAST. This is why fundraising is critical. There is no denying that the numbers we need are big, but **every dollar matters**, and our family is committed to doing our part to help the community.

With that said, we are seeking sponsorships and in-kind donations for this event. 100% of the net proceeds will go to FAST to support their mission to find a cure for every person living with Angelman syndrome.

Thank you for your consideration,

Meghan Edberg - Mother of William, who lives with AS
FAST's Director of Public Engagement