March will.i.can **2024**

Thank you for everyone who went "BLUE" for us on February 15th for International Angelman Day! Uncle Nick raised over $1,000 on his live Twitch Gaming Stream, houses went blue, people wore blue, people generously donated $15 (or more), and William had a blue "glow party" at school!

On March 7, 2024, Sue and Meghan joined 62 other Angelman advocates on Capitol Hill to advocate for the Federal Government to get involved because their support and funding is critical for our community to get a promising therapeutic across the finish line. Minnesota represented with the MOST advocates in D.C. where we met with both senators and 4 House representatives. This was a HISTORIC day for Angelman syndrome.

Read more: Family Advocates Urge Congress' Support During Inaugural Angelman Syndrome Congressional Advocacy Day

Upcoming EVENTS

REGISTER NOW: Sat, May 18th

Join us at **East Lake Community Park** at 10am in Lakeville to raise funds to support the everyday resources and services our AS community needs.

JOIN TEAM WILL.I.CAN TODAY

Click HERE to Register

If Will CAN, We CAN...

Bet on the Trifecta to Beat A.S.

MARK YOUR CALENDAR: Sat, Aug 3rd

Will.i.CAN & **Mighty Evie** invite you out for a memorable evening at **Canterbury Park**, one of America's last great racetracks, to raise critical research funds to support FAST's mission to **Cure Angelman Syndrome (AS).**

TICKETS GO ON SALE MAY 15th

Click HERE to Buy a Sponsorship