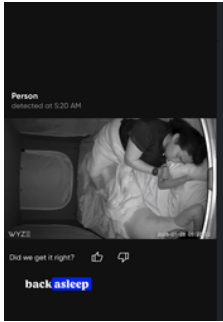


International Angelman Day *Awareness*



This year, our community spammed social media and got “real” with what it’s truly like caring for an individual living with Angelman syndrome. We shared two videos on these topics:

Our sleep schedule (waking up every night) + William’s 140 medical appts in 2025

Watch our videos on instagram then check out all the other families and their stories HERE because we all have our own version of this rare life.

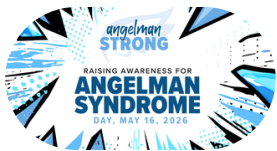
2026 AS Congressional Advocacy Day • Washington, DC

Meghan & Sue joined 138 AS advocates in Washington, DC to urge Congress to prioritize our AS legislative asks which including funding and ensuring our voices are heard when a therapeutic is ready to be approved by the FDA. **Help us and send an alert to your elected officials!**



EMAIL YOUR REPS

Our 2026 Events



Join Team Will.i.CAN for our 7th ASF Walk on **Saturday, May 16th at 9am** in Lakeville, MN - Join our virtual team!



Meghan’s running her first and last marathon representing FAST, AS, and Will.i.CAN at this year’s TPC NYC Marathon on **November 1, 2026.**

LEARN WHY!

If Will CAN, We CAN ...

Brunch in Blue

...for the Angelman Syndrome Crew!

Save the date for Sunday, October 11th at 10am

We are so excited this year to switch things up and brunch in something blue together at CHS Field’s Securian Financial Club, downtown St. Paul. We’ll enjoy a curated brunch, games, and celebrate together the progress of AS.

Sponsorships available today ★ *Tickets go on sale June 10th*
★ *Family Friendly Event for All Ages and abilities!*

LEARN MORE