



Dear Will.i.CAN Crew,

Happy Holidays! What a year it's been – because of you! We raised an astonishing amount of money through donations, events, and t-shirt sales to support a cure. William's fantastic caregivers, teachers, and therapists pushed him to be his best self. Because of that, William has made so many strides this year! Our support system continued to show up for us, which we so appreciate. After watching William make huge progress in his development and hearing the exciting updates in the science and research, the **Edberg's are DREAMING BIG heading into 2023!** We hope you continue to join us and dream big with us as we continue to fight. *We will not stop until we have a cure, which for us means the freedom from the unknown, the anxieties, and worries that come from having a non-verbal and developmentally delayed child.*

Thank you from the bottom of our hearts, *Jesse, Meghan, William, Henry, and Puck*



“There are many disorders that will not be cured or treated in our lifetime, but I'm positive that Angelman syndrome will not be one of them! Because this is not only possible, but it is probable.”

Dr Allyson Berent
FAST Chief Science Officer
Mother of Quincy who lives with AS



Meghan & Sue are getting involved in a BIG way in 2023! Sue has joined the FAST Advisory Council to help advocate for AS

at the state and national level. Meghan left Thomson Reuters and will now be the **Director of Community Engagement** for the Foundation for Angelman Syndrome Therapeutics (FAST)! Watch out AS, the Edberg ladies are coming!

2022 Fundraising Total: \$246,201
3-year total: \$508,201



2nd Annual Will.i.CAN Bowling Event!
Raised \$4,434!

2022 FAST Gala Paddle Raise



Our table gave \$62,600!



Ride with William
Raised \$22,820!