

FAST Auction Donation Form

FAST (*Foundation for Angelman Syndrome Therapeutics*) will host its annual Global Summit & Gala Dec. 1-2, 2017. The star-studded event draws more than 1,000 attendees and is recognized as Chicago's biggest celebrity fundraiser. The gala features an auction that helps generate significant revenue for FAST. One hundred percent of the money raised through FAST's auction will go immediately into grants that fund research focused on finding treatments and a cure for Angelman syndrome. Scientists believe finding treatments for AS in the near future is not only possible it's inevitable. FAST is the largest nongovernmental funder of Angelman syndrome (AS) research.

"Compared to 30 other pediatric neurological disorders, I would make the case that Angelman syndrome is at the top - it is the single-most optimistic possibility for a cure."

- Arthur L. Beaudet, M.D.
Responsible for identifying the gene that causes Angelman syndrome

By supporting FAST's auction you are not only making an investment in finding treatments for Angelman syndrome but also a host of other disorders. Researchers and pharmaceutical companies are interested in Angelman syndrome because it is a single-gene disorder. They believe that finding treatments for AS will help pave the way for breakthroughs with multi-gene disorders like autism and Alzheimer's disease.



GALA DONATION FORM (PLEASE PRINT)

name

phone

email

name of person or company who donated this item

We will contact you when we receive your item(s).

Is this donation in honor of a special child and their family? Y N

please tell us who

description of item

approximate retail value

\$

Please send item(s) with a copy of this form to: FAST, c/o Barb Baldassarre, 504 S. Madison Ave., La Grange, IL 60525-2801 **by Nov. 8, 2017.**

Keep a copy of your completed form for your records.

(If item(s) are being shipped internationally, please email Barb Baldassarre at FASTGala@CureAngelman.org for special instructions.)



FAST

FAST is the largest, non-governmental funder of Angelman-specific research in the world. Paula Evans, an Illinois mother whose daughter was diagnosed with Angelman syndrome, founded FAST in 2008. FAST raises money to fuel cutting-edge research and takes an active role in drug development to treat, and ultimately cure, the disorder.

FAST is a registered nonprofit, tax-exempt charitable organization under Section 501(c)(3) of the Internal Revenue Code. Donations are tax deductible as allowed by law. Please consult your tax advisor for specific guidance. FAST's Employer Identification Number (EIN) is 26-3160079.

WHAT IS ANGELMAN SYNDROME?

Angelman syndrome (AS) is a rare neurogenetic disorder that affects approximately one in 15,000 people – about 500,000 individuals worldwide. Children and adults with AS typically have balance issues, motor impairment and debilitating seizures. Some individuals never walk. Most do not speak. Disrupted sleep cycles also can be a serious challenge to the individual and caretaker(s). Individuals with AS require continuous care and are unable to live independently. They have a normal life expectancy. In most cases, Angelman syndrome isn't inherited. Angelman syndrome was named after Harry Angelman, an English pediatrician, who first described the syndrome in 1965.

CureAngelman.org/Gala

If you have questions,
please email Barb Baldassarre at
FASTGala@CureAngelman.org

Thank you for your support!