



Maddie's Mission is a private, non-profit 501c3 philanthropic organization founded in February 2022.

Maddie's Mission is affiliated with The Shaw Family Foundation

Maddie's Mission is committed to raising awareness, promoting education, and generating funding for research and clinical trials that will provide meaningful treatment for Angelman syndrome and other rare diseases.

Donate Today!

Maddie's Mission will also support several Kansas City organizations that have been instrumental in promoting health and development of children living with different needs.



Meet Maddie!



On January 2nd, 2020, Patrick and Laura Sargent received life-changing news. They were informed that their beautiful 15-month-old daughter, Maddie, had a rare condition called Angelman syndrome (AS).

AS is a rare neuro-genetic disorder occurring in 1 in 15,000 births or 500,000 people worldwide resulting from a loss of function of the UBE3A gene on the 15th chromosome. Several characteristics of AS generally include developmental delays, speech impairment, recurrent seizures, severe sleep disturbances, gastrointestinal issues, and problems with balance and movement. For Maddie, she has daily challenges with gross and fine motor function, communication, GI issues, vision, and sleep.

A typical week for Maddie includes a multi-faceted 40 hours of therapy a week to help Maddie reach unexpected milestones.



OUR NEXT EVENT

Maddie's Mission First Annual Block Party

Join us for a fun filled evening with **Fritz's KC** meats, ice cream food truck featuring **Betty Rae's** and singing and dancing alongside **Atlantic Express!**

Date: 6/12/2022 **Time:** 5-8pm

Location: The JKC Main Entrance- Follow Signs
5801 W. 115th Street, Overland Park, KS 66211

We need sponsors for this event and offer many options:

Platinum (\$5,000) Advertising on event merchandise, printed materials in event goody bag, logo featured in 3 local magazines thanking organization for support, 6 social media posts thanking for sponsorship, 10 free tickets to block party

Gold (\$2,500) Advertising on event merchandise, printed materials in event goody bag, logo featured in 3 local magazines thanking organization for support, 4 social media posts thanking for sponsorship, 8 free tickets to block party

Silver (\$1,500) Advertising on event merchandise, printed materials in event goody bag, logo featured in 3 local magazines thanking organization for support, 2 social media posts thanking for sponsorship, 6 free tickets to in block party

Bronze (\$1,000) Advertising on event merchandise, printed materials in event goody bag, logo featured in 3 local magazines thanking organization for support, 1 social media post thanking for sponsorship, 4 free tickets to block party

Contributor (\$750) Advertising on day of signage, printed materials in event goody bag, 2 social media posts thanking for sponsorship, 2 free tickets to block party

*If the packages above do not meet your needs, please contact us about custom opportunities or in-kind donations. We'd be happy to work with you!

Sponsorship & Volunteering

We have many ways you can get involved with Maddie's Mission. We are always in need of volunteers for our events; to help with simple fundraisers and just spread awareness of our mission on social media.

We need monetary donations as well to help fund research, support local community programs that have had such great impact on Maddie's progress. Please refer to the back of this brochure for specific opportunities.

To learn more, go to:
www.themiraculousmissmaddie.com



A CURE FOR ANGELMAN

The Foundation for Angelman Syndrome Therapeutics (FAST) is committed to finding meaningful treatment for all individuals living with AS as quickly as possible.

"Angelman syndrome is a model for many other genetic diseases including some autism spectrum disorders, and hundreds of other monogenetic disorders. If we can cure AS, these other conditions will benefit; it's only a matter of time."

"We have 9 therapeutic platforms we are working rigorously to de-risk for human application. This process is expensive but unbelievably promising. The only thing standing between us and the potential to have a meaningful treatment for all individuals living with AS is the money—not the science. Your support will bring this all to fruition."

- Allyson Berent, FAST Chief Science Officer, mother of a child living with AS

Our Community



Donate Today!

