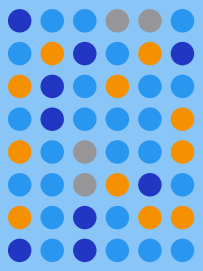




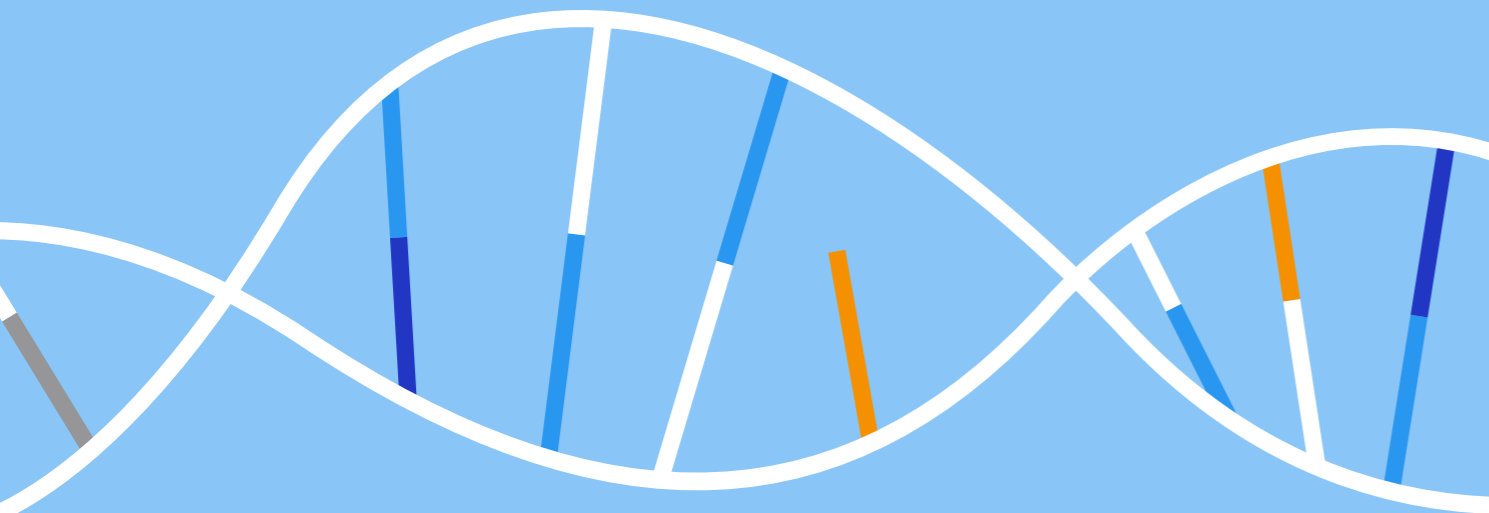
**YOUR SUPPORT MEANS A CURE
FOR ANGELMAN SYNDROME
IN OUR LIFETIME**

fast 
cureangelman.org.uk

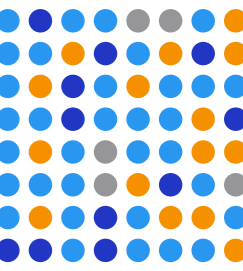
CORPORATE
PARTNERSHIP 2022



WHAT IS ANGELMAN SYNDROME?



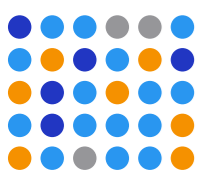
Neave, 6 years



Angelman Syndrome is a complex genetic disorder that primarily affects the nervous system. It affects around 1 in every 15,000 births and is currently incurable. Most people have never heard of this condition despite the estimated 500,000 individuals living with it.

People with Angelman Syndrome don't usually talk and may never walk. They have severe intellectual and physical disabilities as well as seizures. They also develop serious problems with sleep, anxiety and communication.

Based on what scientists know today, there is so much hope that the cure for Angelman Syndrome can be found in our lifetime. Clinical trials are being performed in the UK and worldwide.



OUR STORY MEET EMILY

Angelman Syndrome is not routinely checked for during the pregnancy in the UK, and **there is nothing that can prepare new parents for this diagnosis for their child.**

Diagnosing Angelman Syndrome remains a challenge due to being very rare and, in some cases, due to children not having any apparent symptoms before the age of 6 months.

Since Emily was born, her mother knew that something was not right with her little baby. Still, **it took Emily's family to visit every specialist doctor and run countless tests to finally have a diagnosis.** The one that splits any family's life into before and after.

Emily was diagnosed with Angelman Syndrome at 11 months which is earlier than average. **Most parents only receive the diagnosis for their child between 1-3 years** after doctors start investigation upon missed developmental milestones.

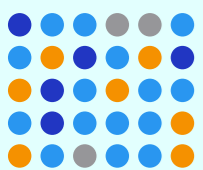
Emily's parents were devastated when they first learnt about the diagnosis. However, **researching about potential treatments and understanding what FAST UK is doing gave them hope.** A hope that one day their daughter may live a more fulfilling and independent life.



Emily, 1 year

“Even reading the news about the recent success in clinical trials makes me cry these days. I can't imagine what I will feel if the cure for Angelman Syndrome is found one day. If I ever get to experience it, I will be the happiest human being on Earth.”

- Emily's mum



WHO ARE FAST?

FAST UK is a UK operation of the Global FAST family, founded in 2016. It focuses primarily on domestic initiatives, including clinical trials for Angelman Syndrome. FAST UK also does all it can to raise awareness about Angelman Syndrome in the UK and across the globe.

The Foundation for Angelman Syndrome Therapeutics (FAST) is an organisation of families and professionals dedicated to finding a cure for Angelman Syndrome and related disorders through funding research, education, and advocacy.

FAST was founded in 2008 in the USA. Today, with operations in Australia, Canada, France, Italy, Spain, and the UK, FAST has become the largest nongovernmental funder of Angelman Syndrome research. And we take this responsibility seriously.



“I was told as a medical student that Angelman Syndrome is one out of a large number of untreatable diseases. Thanks to the hard work of an entire community that needs your support, we are now able to say that we can make this disease disappear.”

- Laurent Servais
*Professor of Paediatric Neuromuscular Diseases,
University of Oxford*

\$20M

Funded by FAST in research towards a cure for Angelman Syndrome

60

FAST-funded grants for Angelman Syndrome research and advancement of therapeutics

21

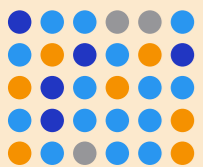
Pharmaceutical companies interested in developing a potential therapeutic for Angelman Syndrome

9

FAST-funded potentially disease-modifying therapeutics

4

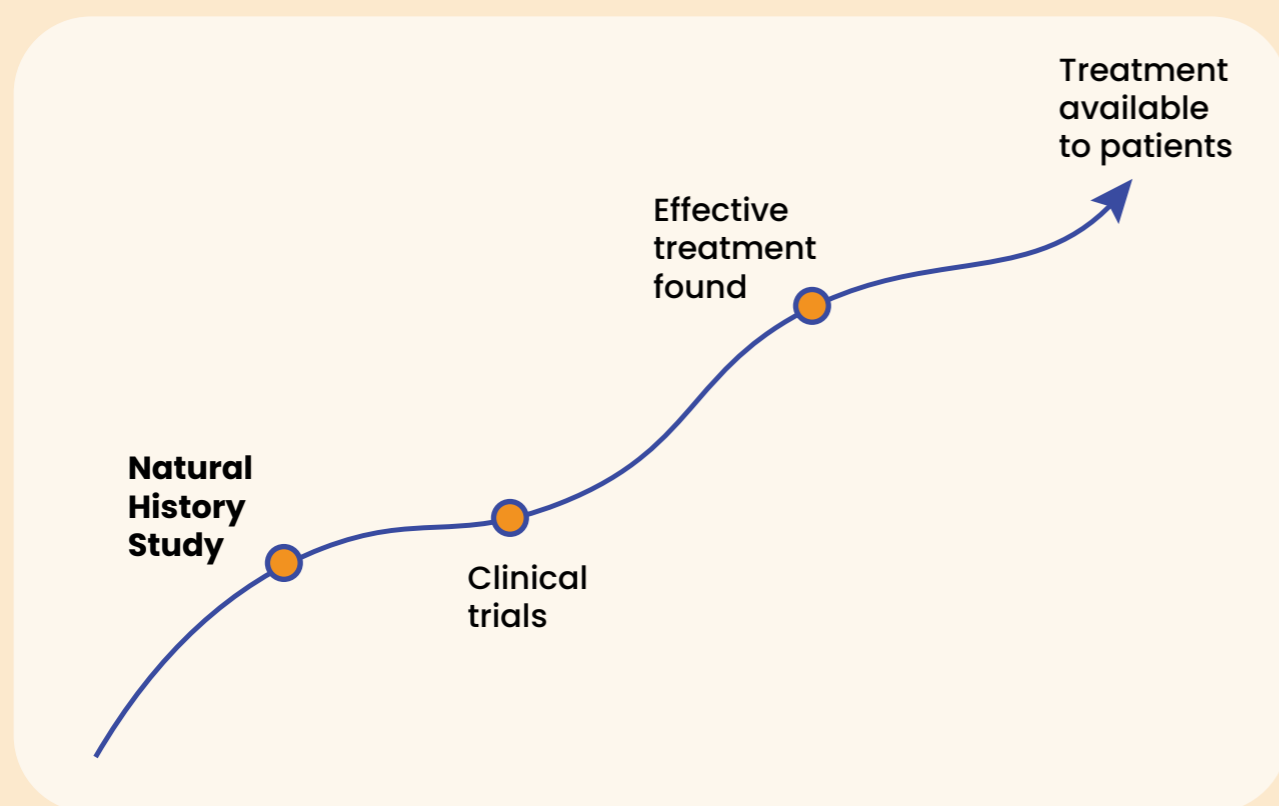
Clinical trials for Angelman syndrome



HOW CAN YOU HELP FIND A CURE IN OUR LIFETIME?

We need your help to raise awareness about Angelman Syndrome, and we need a lot of help to raise funds for the research into a cure for this rare condition.

Our current flagship project is The Natural History Study (The Study) which is a significant step towards bringing more clinical trials into the UK.



Performing The Study will cost £1m over 4 years.

Your contribution can lead to a cure.



£2k covers an innovative ultra-sensitive wearable device for one patient enabling valuable data collection

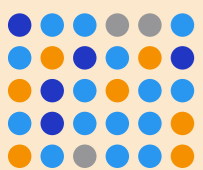


£10k allows one patient to participate in the Study and to line them up for future trials



£250k pays for 1 year of The Study completely

We would love to discuss any options on how your company can support our cause and make a currently incurable disease curable in the near future.



GET INVOLVED



Neve, 11 years

Corporate Donation

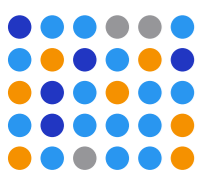
We will show you how your money helps

Strategic Partnership

Join us and we will cure Angelman Syndrome together

Employee Fundraising

We will enable team bonding for an important cause



OUR EVENTS



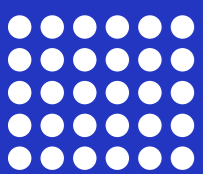
We have had some fantastic fundraising events all over the country, from people holding galas to others running marathons for FAST UK.

One of the most memorable fundraisers was Art for Angelman, where we had a raffle for over 200 donated pieces of art from professional artists. People bought tickets to come, and as part of the ticket, they were entered into the raffle. We had 400 people at the reception, and as a result, 200 people won a piece of art. The event raised just over £60,000.

Another noteworthy event was run in July 2021 by Cecilia's parents. It included a golf event, a cycling challenge and a raffle. More than £8,000 was raised thanks to the monumental efforts of one family.

“As well as raising money for scientific research, we feel that it is as important to be raising as much awareness as we possibly can about AS in the UK. Fundraisers like Art for Angelman not only raise large amounts of money, but they also do so much for raising awareness.”

- Tom Keogh,
FAST UK Chairperson



CONNECT WITH US

We'd love to hear from you.

Email: info@cureangelman.co.uk

We would love to partner with your business. Please get in touch with us to discuss options and opportunities, whether it's employee fundraising and engagement, commercial opportunities or one-off events.

#CureAngelmanNow



