Is your child suspected to have a genetic condition but the doctors don't know what it is?



Does your child have medical, physical or learning disabilities which are currently unexplained?

Has your child had genetic tests that have come back negative or as a 'variant of unknown significance'?

SWAN UK supports families affected by a syndrome without a name – a genetic condition so rare it often remains undiagnosed.

- Connect with other families who understand the unique challenges of raising a child affected by a syndrome without a name
- Attend our events such as virtual coffee meets and family discos
- Receive the latest news and updates that affect the SWAN UK community

To find out more about joining visit geneticalliance.org.uk/swanuk















'When we entered the undiagnosed world it was daunting and exhausting and terrifying. Being able to speak to others who literally know what we are trying to navigate was so comforting and reassuring.'



SWAN UK, run by Genetic Alliance UK, is the only dedicated support network for families affected by a syndrome without a name. Find out more about our work at geneticalliance.org/swanuk

SWAN UK is the only dedicated community for UK based families of children aged 0-25 years old who are searching for a genetic diagnosis.

We bring families together in a supportive community who understand the unique challenges of life without a diagnosis. We also:

- Provide high quality information for families
- Offer members of the SWAN **UK community opportunities** to develop personally and professionally
- Raise awareness of syndromes without a name by working to increase public and professional understanding of the unique challenges that families face

Scan the QR code to donate today or text SWANUK followed by the amount 1-20 to 70085. For example, text SWANUK10 to donate £10. (Text messages cost the donation value, plus your standard network message charge).







