CURE MITO FOUNDATION

Get to Know Us!

We are a nonprofit patient advocacy organization led by parents who volunteer their time to change the future for families affected by Leigh syndrome.

Our mission is to unite the global Leigh syndrome community to accelerate patient-centered research, treatments, and cures.

WHAT WE DO:

Largest Leigh Syndrome Global Patient Registry: curemito.org/leighsyndromeregistry

Fund research

Online resource aboutleighsyndrome.com

Empower and Inspire, Leigh Syndrome Symposium

Patient support and caregiver resources

Fibroblast repository initiatives

WHAT IS LEIGH SYNDROME?

Most common pediatric mitochondrial disease

Neurodegenerative disease causing the loss of abilities to walk, talk, swallow

Symptoms start between 3 months to 2 years of age, although earlier and later onset is possible

Caused by 110+ nuclear and mitochondrial DNA mutations

Approximately 1 in 40,000 are affected

Connect

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Cure Mito Foundation is a 501(c)(3) non-profit organization
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To learn how to be involved, please email info@curemito.org