



**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/registry](https://curemito.org/registry)

Drug Repurposing Research

Gene Replacement Therapy Research

Empower and Inspire: Leigh Syndrome Symposium

Patient Support and Education

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



[curemito.org](https://curemito.org)



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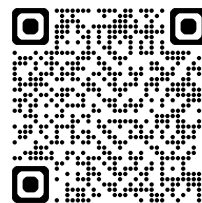
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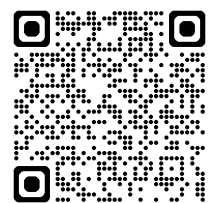
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Cure Mito Foundation is a 501(c)(3)  
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