



Rare disease Day is a week away from today! Below are some ways you can make a difference and support Leigh syndrome community!

**DONATE**

Share our posts on social media

Join Leigh syndrome [registry](#).

Shop at CURE MITO [bonfire store](#)

For more ideas visit: [CUREMITO.ORG](#)



[RARE DISEASE DAY.ORG](#)

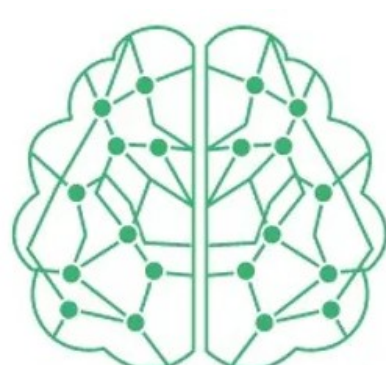
Cure Mito Foundation is led by parents like you!  
We volunteer our time to raise funds for a cure,  
work on a patient registry  
and create resources and support for families



In the honor of Rare Disease Day, we made a fun 1-minute doodle video. Please watch and share!  
[https://youtu.be/GTR32\\_BeSYo](https://youtu.be/GTR32_BeSYo)

We will be supporting #RareDiseaseDay and raising awareness of mito for the global community by participating in a "Tweetstorm" taking place on 28<sup>th</sup> February 2023 and organized by Wellcome Centre for Mitochondrial Research!

Please follow us on twitter [@cure\\_mito](#)



**We are getting ready for the 2nd Annual Leigh Syndrome Symposium!**

Empower & Inspire:  
Understanding and Accelerating Research for Leigh Syndrome

TUESDAY, SEPTEMBER 19, 2023

Brought to you by:  
Cure Mito Foundation  
&  
integrative Cardiovascular Metabolism and Pathophysiology Laboratory (iCaMP) at Boston University

**REGISTER NOW**

To view conference 2022 agenda and recordings, please click [HERE](#)  
If you are interested in speaking, please complete [speaker interest form](#)  
Please click [HERE](#) to learn more about sponsorships.



**Leigh Syndrome Families - Please Join the Patient Registry and Be Counted**

Largest LS patient registry in the world  
Accessible and available data  
Results are always reported back to the community

[CLICK HERE TO LEARN MORE](#)

**LATEST PATIENT REGISTRY NEWS**

Preprint of our paper: "Leigh Syndrome Global Patient Registry: Uniting Patients and Researchers Worldwide" is available! Please find it at:  
<https://www.researchsquare.com/article/rs-2280399/v1>

**ABOUTLEIGHSYNDROME.COM**

We are continuing to work on ABOUTLEIGHSYNDROME.COM - a new website and a landing place for patients, families, providers, and the entire global Leigh syndrome community. If you are interested in being involved in this project please contact us at [info@curemito.org](mailto:info@curemito.org)

A huge thank you to [PTC Therapeutics, Inc.](#), for their generous sponsorship that is making this resource possible for our community.

Cure Mito Foundation is a 501(c)(3) nonprofit organization led by parents who volunteer their time to search for a cure. 100% of your donations are tax-deductible and will go directly to research dedicated to mitochondrial diseases.

For many ways to give, please click here:

<https://www.curemito.org/ways-to-give>

**Get in touch and join us!**

Please visit our website to learn about our research projects, resources, and more:

<https://www.curemito.org/>

We invite you to join us in our efforts! For ways to get involved please visit:

<https://www.curemito.org/get-involved>

**DONATE NOW**

