



We are wrapping up 2022! This has been a year of huge growth for Cure Mito! This year we have:

- Presented our patient registry results at 8 conferences and submitted a paper to an open access and peer-reviewed journal
- Funded gene-editing project for the MT-ATP6 gene
- Funded drug repurposing project with Perlara
- Funded Rarebase on their Function platform
- Held our first International Leigh syndrome conference with over 200+ attendees from 34 countries
- Launched LS Birthday club that is currently providing birthday gifts for almost 100 kids. The children LOVE receiving their "Mito Warrior" shirts and birthday cards!
- Began work on an AboutLeighSyndrome.com - a new resource for the entire LS community
- Received a 2022 GreatNonprofits Top-Rated Award

Much more is coming up in 2023 and we have an exciting announcement to share in early 2023! Stay tuned!

NEW RESOURCE FOR OUR COMMUNITY!

We are excited to announce [AboutLeighSyndrome.com](#) a new resource for the Leigh syndrome community. This website will be dedicated to LS education, research, support, and more for both patients and physicians.

Please sign up to be notified when the site goes live and share with us what information and resources are important to you! Additionally, please reach out to us at info@curemito.org with any questions, comments, or collaboration inquiries.

A huge thank you to [PIC Therapeutics, Inc.](#) for their generous sponsorship to make this resource possible for the global Leigh syndrome community!

Coming Soon!



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:

Our posters were presented at the Mitochondrial Medicine - Therapeutic Development conference in November.

Please click on the links below to hear quick 2-minute overviews of each poster

["Building a Worldwide Community – Leigh Syndrome Patient Registry"](#)

["Interoperability of Leigh Syndrome Registry Data with Regulatory Submission Standards"](#)

We are excited to share that we have joined COMBINEDBrain - a non-profit consortium led by patient advocacy foundations, working with the clinicians, researchers and pharmaceutical firms that are developing treatments for the disorders they represent. It is devoted to speeding the path to clinical treatments for people with severe rare genetic non-verbal neurodevelopmental disorders by pooling efforts, studies and data.



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Thank you to all of you who supported us this #GivingTuesday! It is not too late to donate. 100% of your donations are tax-deductible and will go directly to research dedicated to mitochondrial diseases. Cure Mito Foundation is a 501(c)(3) nonprofit organization led by parents who volunteer their time to search for a cure.

For many ways to give, please click [here](https://www.curemito.org/ways-to-give): <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>

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