



Cure Mito Foundation, in collaboration with five of the industry's prominent rare disease companies - PTC Therapeutics, Astellas, Saol Therapeutics, Standigm, and Abliva, announced the launch of a unique Corporate Advisory Council (CAC).

Please learn more in the press release:  
<https://www.eurekalert.org/news-releases/976914>



## Corporate Advisory Council



Leigh Syndrome Conference is back again this year!

### **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](#)

If you are interested in sponsoring, please reach out to us at [info@curemito.org](mailto:info@curemito.org)

#### PROJECTS

You can now find our current projects on one page - we hope it makes it easier to follow! All our projects are done for the entire community WITH the community - please reach out at [info@curemito.org](mailto:info@curemito.org) if you'd like to be involved.



#### PARENTAL WELL-BEING GUIDE

How can parents take care of themselves while also caring for their child with a life-limiting condition, such as Leigh syndrome? Please find some tips in our new guide. Thank you, Dr. Rachel Kramer for writing it.  
<https://www.curemito.org/caregiver-wellness-guide>



#### NEW RESOURCE FOR OUR COMMUNITY!

We are continuing to work on [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](mailto:info@curemito.org) with any questions, comments, or collaboration inquiries.

A huge thank you to [PTC 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](#)**



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**

