



## Patient Registry Update

We are so proud to share that just a few months since starting a patient registry, our poster "[Leigh Syndrome Global Patient Registry - Cure Mito Foundation](#)" will be presented at the Mitochondria-Targeted Drug Development summit this February! We will also share the poster with the entire community at that time! Please stay tuned!

After that we're planning a bigger, more detailed publication. If you or your loved one has Leigh Syndrome and have not enrolled into the registry yet, please enroll soon to contribute to the results! Register here: <https://www.curemito.org/registry>



## Finding Your Family Path

### Insights and perspective from other parents and resources from Courageous Parents Network

As Rare Disease Day approaches please join us for a discussion with Blyth Lord, founder of [Courageous Parents Network](#) exploring the spectrum of the parent experience grounded in qualitative research about parent beliefs, values, and priorities when caring for a medically complex child.

Register: <https://us02web.zoom.us/meeting/register/tZckcOtpzMiEt02BBgcHYmdCz2gfDsZDyC>



## 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

Copyright © 2021 Cure Mito Foundation, All rights reserved.

Want to change how you receive these emails?  
You can [update your preferences](#) or [unsubscribe from this list](#).

