

# LGS FOUNDATION LENNOX-GASTAUT SYNDROME

## Navigating THE LGS MAZE



### 12 Days of LGS Giving Stories Because You Give, We Can Give To Others!!

Thanks to you, we are changing the world of LGS!

As we wrap up 2022, we would like to extend our deepest gratitude and thank you for all the ways you support and empower the LGS Foundation and those impacted by this horrible disease.

We invite you to follow and share the 12 Days of Giving throughout December to celebrate important moments made possible because we are Standing Together and Stronger Together!

Please share these special moments with people you know – Because *Every Moment Matters!*

[Give Now](#)

[Read the 12 Days of Giving Stories](#)



### The LGS Research Roundtable Was A Huge Success!

Every year, the LGS Foundation hosts the LGS Research Roundtable at the American Epilepsy Society (AES) meeting. This year's meeting, which was in Nashville, filled us with so much hope as we learned from medical experts about exciting research going on to find better treatments and cures for LGS. Topics included:

- The Natural History of LGS
- An Update on Medical Treatments for LGS
- About a New PCORI-Funded Study Comparing Surgery vs. Medication in LGS
- Comparing Corpus Callosumotomy to Vagus Nerve Stimulation (VNS) in LGS
- Responsive Neurostimulation (RNS) in LGS
- Deep Brain Stimulation (DBS) in LGS
- Clinical Trial Design in LGS
- Measuring the Evolution from Infantile Spasms to LGS

These talks set the stage for our upcoming 2023 LGS Meeting of the Minds: Advancing Clinical Research in LGS, which will take place in September at the University of Maryland and help guide the research funding strategy for the LGS Foundation in 2023 and beyond. We hope you will take a moment to learn more about these important updates in LGS Research and plan to join us for the Meeting of the Minds.

[Watch The Recordings](#)

#### We Are Committed to Funding LGS Research!

Since 2014, the LGS Foundation has funded over **one million dollars** in LGS research.

The Foundation's Cure LGS 365 Research Grant program has provided funds for fourteen projects around the world, and we are not done yet!

The LGS Foundation accepts unsolicited proposals **year-round** to seed new basic, translational, and clinical research projects on LGS.

[Learn More](#)

#### Save the Date

### 2023 Meeting of the Minds: Advancing Clinical Research in LGS

The LGS Foundation is hosting its second Meeting of the Minds to discuss how we can **Advance Clinical Research in LGS**. For this hybrid conference, we will bring together patients, families, doctors, researchers, and partners to understand how we can learn from every patient and brainstorm how to find the best treatments for LGS.

The Family Science Day will be on **September 16, 2023**. We hope you will join us!

[Learn More](#)

## SEPTEMBER 2023

## LGS RESOURCE CORNER

#### Planning an LGS Event or FUNdraiser?

You can now submit your upcoming LGS events to our Calendar of Events so we can help spread the word! Check it out.

[Submit Your Event](#)

#### Clinical Trial Opportunity for Those with LGS

The PACIFIC Study is a Phase 1b/2a clinical study for adult participants with DGS. This study will assess the safety & tolerability of LP352.

[Learn More](#)

#### Has your Loved One had Epilepsy Surgery?

Please take a moment to complete this short survey for one of our scientists. Answers will remain anonymous.

[Take the Survey](#)

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