

Navigating the LGS Maze Community Newsletter

LGS FOUNDATION
Lennox-Gastaut Syndrome



INTERNATIONAL
LGS FOUNDATION
FAMILY & PROFESSIONAL CONFERENCE

Register Today!!

The Conference is NEXT MONTH... Don't Miss This Opportunity!

There's still time to register to attend July 12-14th in person or virtually!

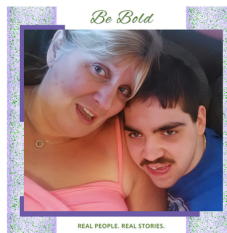
Our 9th International Family & Professional Conference is almost here! Whether you attend in person or virtually, don't let this invaluable opportunity to learn, grow, and connect pass you by. This is an event you do NOT want to miss!

- Educational Sessions and Interactive Workshops
- Building Connections to Last a Lifetime
- Exhibit Hall & Resource Center
- Dinner & Dance Party
- LGS Sibling Adventure Camp
- Memorial Celebration of Life Butterfly Release
- Butterfly Wonderland Family Field Trip

Virtual access includes live streaming and recordings of all main conference sessions and most workshops.

[Learn More & Register Now](#)

The Conference is made possible with the support of our partners at Neurelis, Jazz Pharmaceuticals, UCB, Takeda, Longboard Pharmaceuticals, LivaNova, Asserito, Marinus, and Azurity. Thank you!



BE BOLD in Raising LGS Awareness!

Each November, the LGS Foundation organizes international social media campaigns, events, and fundraisers to raise awareness and funds to support the LGS Foundation's mission to improve the lives of individuals impacted by LGS.

On **Saturday, November 2nd**, we'll host our **3rd Annual LGS Awareness Dinner in San Diego, CA**.

The evening will feature a silent auction and cocktail reception, seated dinner, inspirational program, exciting live auction, live entertainment, and more.

[Learn More](#)



Clinical Trials in LGS

LGS FOUNDATION
LENNOX-GASTAUT SYNDROME

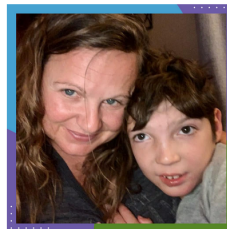
Learn About Active Clinical Trials in Lennox-Gastaut Syndrome!

Clinical trials and research studies are done to scientifically evaluate a medical or surgical intervention. They are the primary way that researchers find out if a new treatment, like a new drug, diet, surgery, or medical device, is safe and effective in reducing seizures in those with LGS.

Considering joining a trial?

Learn more about the **5 active clinical trials and research studies** in LGS happening across the United States.

[Learn More](#)



Our Community: Meet Julian

We love sharing stories and getting to know one another, so each month we will feature a member of our amazing community. This month, we would like you to meet **Julian**.

He had his first seizure at a year old, which lasted an hour and 45 minutes. He was diagnosed with LGS at 4-years-old, when he began to have daily seizures.

His Mom shares, "Despite all his struggles, Julian is a very happy boy who is constantly smiling and giving fist bumps. His positivity and perseverance are admirable, and his smile lights up a room."

[Learn More about Julian](#)

LGS RESOURCE CORNER



Learn How LGS is Treated

There are currently no cures for LGS, but numerous treatments for seizures are available.

As new treatments become available, we update the website with everything LGS Families need to know.

[Learn More](#)



Attend an Online Support Group

The LGS Foundation co-hosts several online support groups each month.

These support groups are only available to LGS caregivers who are members of the LGS Foundation's Online Caregiver Support Community.

[Learn More](#)



Connect Your Healthcare Team to Us

Do you have a healthcare professional you want to connect to the LGS Foundation?

Share our website with them today!

[Learn More](#)



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Standing Together. Stronger Together.

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