

LGS FOUNDATION LENNOX-GASTAUT SYNDROME

Navigating THE LGS MAZE



Celebrating 15 Years of Supporting Those Impacted by Lennox-Gastaut Syndrome

It all started with a sibling... a sibling who watched their family struggle through what felt like overwhelming obstacles as they navigated life with LGS.

15 years later, the LGS Foundation, an organization formed from the love of a family member, has transformed into an unstoppable force. It has been an exciting decade and a half of growth and success for our LGS Community. As a part of this great community, YOU helped make this happen.

You're Invited!



We invite you to join us for an evening of dance and fun at the 15th Anniversary Dance Party on August 12, 2023. This private event is only available to members of the LGS Foundation Caregiver Support Community. Join to get connected and learn more about the dance.

[Join the LGS Caregiver Support Community](#)

Swing FORE a Cure is Next Month!

It's that time of year again... Join LGS Foundation's Vice President, Karen Graft, and her family for the 2nd Annual Swing FORE a Cure for LGS.

Karen's son Danny, age 37, began having seizures in his first year of life and was eventually diagnosed with LGS when he developed multiple seizure types.

Karen shares, "My husband Jeff and I are inspired to help raise funds for LGS research. We know this is the best way to ensure better outcomes for those suffering from this disease."

[Learn More About Swing FORE a Cure](#)



Save the Date: International LGS Awareness Day is November 1, 2023!

This November, we'll kick off Epilepsy Awareness Month with International LGS Awareness Day!

Stay tuned for more details on all the exciting things we have planned:

- LGS Awareness T-Shirts
- LGS Awareness Boxes
- LGS Foundation Anniversary Dinner: 15 Years of Supporting Those Impacted by LGS in San Diego, CA
- Social Media Toolkit and Awareness Campaign
- Buildings and Monuments Illuminate for LGS
- ... and More!

[Learn More About LGS Awareness](#)

15th Anniversary - Fun Fact!

Did You Know?

Our First Meeting of the Minds Research Conference was in 2021!

In September 2021, over 250 researchers and family members from 7 different countries joined the LGS Foundation for a 2-day discussion focused on finding disease-modifying therapies for LGS.

This September, the LGS Foundation is hosting its second Meeting of the Minds to discuss how we can advance Clinical Research in LGS.

[Learn More](#)



About the LGS Community of Support

The LGS Foundation has grown exponentially in the last 15 years, so we're taking some time to reintroduce you to our amazing community of support. Each month we will feature someone who has made a significant impact on the lives of those affected by LGS.

This month we would like to you meet...

LGS Family Ambassador Mandy Graham

Mandy, her husband Keith, and two children, Brook (16 with LGS) and Gabriel, live in Kansas. Mandy shares, "I believe it is vital for us to share information to further grow, educate, and raise awareness about LGS. So much has changed since Brook's diagnosis. I feel it is important for us to share our journey with others. The more knowledge we can openly share can lead to advances in treatments, caregiver education, and quality of life for those with LGS."

[Learn About the Patient Navigator & Family Ambassador Program](#)

LGS RESOURCE CORNER

This month we'd like to present to you,

KIT-Palooza!

Below are three carefully curated Resource Kits to help caregivers and siblings understand and navigate their journey with LGS. These kits offer a wealth of information and are available to limited free-of-charge.

Shipping in the United States only. Electronic downloads available.



NEW FAMILY WELCOME KITS

Whether you're new to LGS or new to the LGS Foundation, we're here to support you. The New Family Welcome Kits provide educational materials to those who want to learn more about LGS and the programs the Foundation offers.

[Request Yours Now](#)



VIP SIBLING KITS

Living with a chronic illness impacts everyone in the family. Our VIP Sibs Kits offer support and resources for families who are impacted by new epilepsies.

[Request Yours Now](#)



ABOUT LGS TREATMENTS KITS

This comprehensive kit contains information on current treatments, clinical trials, and tips on maximizing your time with your loved one's healthcare team.

[Request Yours Now](#)

Follow the LGS Foundation on Social Media



[Check Out Our Calendar of Events](#)

[Give Now to Help Families Impacted by LGS](#)

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