



LGS FOUNDATION®

LENNOX-GASTAUT SYNDROME



Accelerating LGS Research



180+

Families enrolled in Phase 1 of the LGS Learn from Every Patient Database



\$125K

In Research Grants Awarded to New Cure LGS 365 Grant Recipients

Support for LGS Families

3,000+ LGS Educational Kits Mailed to Families

"As a parent dealing with this new diagnosis, the welcome kit helped me better understand and explain to family and teachers more about my child's condition."

-LGS Caregiver



"I am excited by the prospect of contributing to LGS research in a manner that can ultimately lead to improved treatments for patients."

- Colleen Carpenter, PhD
Cure LGS 365 Grant Recipient



Increasing Awareness & Education

400+ Global leaders, healthcare professionals, and families united in Phoenix, AZ for 3 days of educational sessions, treatment updates, and workshops at the 9th International Family & Professional Conference.

\$15K+

Awarded to patient-families to assist with essential durable medical equipment

800+

LGS Caregivers received individualized one-on-one support from our team

4 Monthly Support Groups

Newly Launched this year for Spanish speaking families: LGS Grupo de Esperanza

7.5K+

Members in our private Global Online Caregiver Support Community

133

Financial scholarships provided to patient-families who would otherwise be unable to attend.

"Leaving the conference, I felt a mix of emotions, but most prominently, I felt hopeful and empowered. The wealth of information and support available provided a renewed sense of optimism."

-LGS Caregiver



120+

locations across 6 countries Illuminated for LGS during Epilepsy Awareness Month

43 Advocates for LGS

Advocates attended Rare Disease Day on Capitol Hill and Rare Across America



9th International Family & Professional Conference

