

OUR IMPACT

LGS FOUNDATION[®]
LENNOX-GASTAUT SYNDROME

The LGS Foundation is the only global nonprofit organization dedicated to improving the lives of individuals impacted by LGS through advancing research, awareness, education, and family support.

Educating,
Empowering,
& Supporting
LGS Families

7,000+ In Our Online
Private Support
Community
Caregivers

90,000+ Across All
Social Media
Channels
Followers

600+
New Family
Welcome Kits
Sent

"There is something devastating about raising a child and never knowing where they fit. The LGS Foundation has given us a place and people to love and call home. The people I've met along this journey are family to us."
- ANDEE, MOM TO KANNON

ACCELERATING LGS RESEARCH WITH LGS FAMILIES AS PARTNERS

600+
About LGS
Treatments Kits
Sent

**\$1.5
Million**

Invested in LGS
Research since
2008

500

Have joined the
LGS Collaborative
Research Network

100

Enrolled in the LGS
Learn From Every
Patient Database

19

International
Conferences
since 2008

400+
Sibling &
Caregiver Kits
Sent
in 2023

**25 ADVOCATES
FOR LGS**

"Being an advocate to me means to bring recognition, change, awareness, light and hope to a community that so desperately needs it."
-Rachel, Advocate for LGS

2023 EVENTS

**Walk 'n' Wheel for
LGS Research**

**Meeting of the Minds
Research Conference**

**International LGS
Awareness Day**

EVENT OUTCOMES

- Over 220 Attendees & Dozens of Grassroots Local Walks
- 3,000+ Press Release Views and Hits
- Over 150 Attendees
- Productive discussions on methods to measure seizure and non-seizure outcomes
- 100 Awareness Boxes Sent Across the United States
- Facebook and Instagram Reach Increased by 300% in November with Profile Visits up over 200%



**Standing Together.
Stronger Together.**



280 FAMILIES in 2023
SUPPORTED

"Being able to use what my family has experienced because of our son's diagnosis, to support others along their LGS journey is a profound gift to me."

-Tricia, Family Ambassador

50

Family
Ambassadors
and
Patient
Navigators

7

Countries

"The LGS Foundation is amazing. You become family instantly. Any questions or concerns you have, you can ask and will get responses." - LGS MOM

OUR COMMUNITY IS STRONGER WITH FACTS!

ANNUAL MYTHBUSTER CAMPAIGN

DISPELLING COMMON MYTHS AND MISUNDERSTANDINGS
SURROUNDING LGS AND TREATMENT OPTIONS

6.3%

Email CTR

69,260

Facebook Reach

19,622

Instagram Reach

1,357

Web Page Views



\$500k

103

**FAMILY MEMBERS
ATTENDED ON
SCHOLARSHIP
IN 2022**

"Attending the LGS Conference is good for the mind, body, spirit and soul." - LGS Caregiver

Platinum
Transparency
2023

Candid.



Charity
Navigator



★ FOUR-STAR ★

**In
Scholarships
Awarded to
Families to
Attend our
Biennial
Family &
Professional
Conferences.**