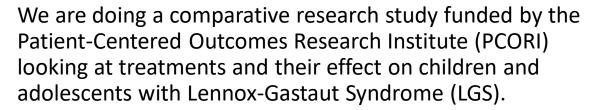
PARTICIPANTS NEEDED FOR A RESEARCH STUDY:

Do you have a Child with Lennox-Gastaut Syndrome?

LGS caregiver perspectives



We would like to look at results of treatment pathways as well as developmental outcomes such as communication, behavior, and their impact on quality of life.

We ask for your help to give feedback on your experience and thoughts in caring for people with LGS.

Please note that we plan on publishing this information; however, no private information will be used. Please know this is voluntary and you can participate if you wish.

Study Title: Comparative effectiveness of palliative surgery versus additional antiseizure medications for Lennox Gastaut Syndrome PI: Dr. Sandi Lam



Who: Caregiver to a child with Lennox-Gastaut Syndrome

What is involved: The participation consists of completing a series of standard survey assessments to measure behavior, communication, and quality of life and possible follow up including LGS patient history, recent seizures (severity and frequency), medications since onset. You can complete surveys at a time and place convenient for you. Your family will receive a gift card as a thank you for your time and participation.

To learn more or to participate, please visit lgsresearch.org and email study coordinator for guidance-info@LGSResearch.org