PARTICIPANTS NEEDED FOR A RESEARCH STUDY:

Do you have a Child with Lennox-Gastaut Syndrome?
LGS caregiver perspectives

We are doing a comparative research study funded by the Patient-Centered Outcomes Research Institute (PCORI) looking at treatments and their effect on children and adolescents with Lennox-Gastaut Syndrome (LGS).

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.

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

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

This study is Lurie Children’s IRB #2022-5454. The content of this flier/brochure/e-mail/etc. has been approved by the Lurie Children’s IRB.