

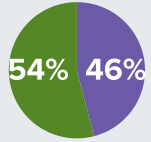
Caregiver Perspectives on Quality of Life in Lennox-Gastaut Syndrome (LGS)

Understanding Barriers to Comprehensive Care Evaluation

Survey-based study examining caregiver perspectives on behavioral, communication and quality-of-life assessments for children with Lennox-Gastaut syndrome and drug-resistant epilepsy.

STUDY AT A GLANCE

- 245 Caregivers
- 54% LGS
- 46% Non-LGS Drug-Resistance Epilepsy



WHAT CAREGIVERS VALUE



Communication



Behavior



Quality of Life

OVER 90% SAY THESE EVALUATIONS ARE IMPORTANT

THE GAP

Less than 50%

Feel current healthcare evaluations **ADEQUATELY CAPTURE** their child's:

- Behavioral challenges
- Communication needs
- Impact on daily Quality of Life

42%

Behavioral Challenges

38%

Communication Needs

45%

Quality of Life

Significant discrepancy between **perceived importance** and **satisfaction** with current assessments.

CALL TO ACTION & RECOMMENDATIONS

- Incorporate **standardized, caregiver-recommended tools** to evaluate behavior, communication, and QoL during routine doctor's visits.
- Improve **communication between doctors and caregivers** to better understand individual patient needs.
- Prioritize **comprehensive evaluations** to improve whole-person care for children with LGS and drug-resistant epilepsy.

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