

Non-Seizure Symptoms of Lennox-Gastaut Syndrome

In addition to the seizures that are the hallmark of Lennox-Gastaut syndrome (LGS), there are numerous non-seizure symptoms that occur in most patients. These symptoms can have a major impact on patient well-being and the quality of life of both patients and their caregivers.¹ Read more about the non-seizure symptoms of LGS below.



COGNITIVE^{1,2}

- Intellectual disability occurs in 75% to 95% of children with LGS, with impairment worsening over time
- Some level of cognitive impairment is required for diagnosis
- Communication difficulties (both verbal and non-verbal)



BEHAVIORAL AND MOOD^{1,3,4}

- Hyperactivity (ADHD), aggression, and autistic traits
- Behavioral issues worsen with age, particularly in adolescence
- Anxiety/depression
- Sleep disorders exacerbated by nocturnal seizures



MOTOR AND MOBILITY⁵

- Staggered gait, difficulty walking
- Some individuals require mobility support, such as a wheelchair
- Challenges in feeding, dressing, and caring for self
- Dysphagia increasing with age



DEATH^{6,7}

- Sudden unexpected death in epilepsy (SUDEP) is the main contributor to early mortality
- Nearly 14-fold increase in death compared to peers
- True extent of mortality may be underestimated due to underdiagnosis

References

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