The Psychosocial Impact of Growing Up With a Sibling With Severe Childhood Epilepsy

Severe childhood developmental epileptic encephalopathies (DEEs), such as Dravet and Lennox-Gastaut syndromes, are rare, intractable (treatment-resistant) types of epilepsy in which seizures typically contribute to brain dysfunction. These syndromes occur in a small number of people but can affect the individual and their family in numerous and significant ways. Siblings of children with DEEs may be at risk for anxiety or depression, and many expressed fear that their sibling might die.

**≥47% of young and adult siblings reported experiencing feelings of unhappiness sometimes or a lot**

**≥53% of young siblings reported experiencing feelings of irritability or grumpiness sometimes or a lot**

**35% of adult siblings reported a history of treatment for clinical depression**

**79% of young siblings aged 9-12 expressed fear that their sibling might die**

The Siblings Voices Study, which was developed to assess the emotional impact of growing up with a sibling with an DEE, revealed that siblings often suffer from negative psychosocial effects that can impact their quality of life. Parents of these siblings may underestimate the effect growing up with a sibling with severe epilepsy has on them.

The study included 3 subgroups of siblings: The young siblings group included cohorts of siblings aged 9-12 and 13-17, and the adult siblings group included siblings aged 18 and over. Parents were also invited to participate in the study.

**Most frequently reported emotional challenges cited by young siblings**

- Easily startled
- Unhappiness
- Irritability
- Bad dreams

**Top concerns cited by young siblings**

- Worried/scared
- Less parental attention
- Parental stress
- Activities being disrupted
- More responsibilities at home
- Embarrassment
GREATER KNOWLEDGE ABOUT A SIBLING’S DEE HAD A BENEFIT (LESS PSYCHOSOCIAL IMPACT) FOR UNAFFECTED SIBLINGS

52-76% OF YOUNG AND ADULT SIBLINGS who felt knowledgeable about DEEs and/or helped during a seizure had lower depressed/anxiety mood symptom scores.

38-86% OF YOUNG AND ADULT SIBLINGS who felt knowledgeable about a sibling’s DEE reported feeling more comfortable talking with others about their sibling’s diagnosis.

ADULT SIBLINGS ARE WORRIED ABOUT TAKING ON THEIR SIBLING’S CARE

65% expressed concern over the psychological/emotional toll of caring for their affected sibling.

70% fear that something bad would happen to their sibling in their care.

67% expressed concern over being able to provide a fulfilling life for their sibling.

SIBLING SELF-REPORTING OF PSYCHOSOCIAL IMPACT DOES NOT ALIGN WITH PARENTAL REPORTS

24-58% of siblings’ responses indicated potential anxiety symptoms, yet only 14-38% of parents perceived that the siblings experienced anxiety symptoms.

42% of siblings aged 9-12 reported high scores of sadness, while only 20% of parents reported recognizing signs of this degree of sadness in siblings.

Siblings of children with severe DEEs grow up in an environment permeated by stress, anxiety, and fear. The significant strains on the family’s emotional and financial resources have the potential to significantly negatively impact the sibling’s psychological health and may lead to feelings of depressed mood, anxiety, or other psychological comorbidities.

FAMILIES ARE ENCOURAGED TO SPEAK TO A HEALTHCARE PROFESSIONAL about ways to support the challenges siblings face when having a brother or sister with a DEE.

References:
8. Bailey LD, Gammatoni AR, Galler BS, Schwartz L, Schad C. Constructive vs destructive coping methods in siblings of patients with epileptic encephalopathies: targets for intervention. Poster presented at: National Organization for Rare Disorders (NORD) Rare Diseases & Orphan Products Breakthrough Summit; October 15-16, 2018; Washington, DC.