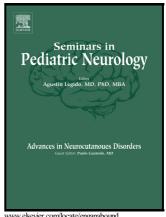
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Impact of Epilepsy on Families, Communities and Society

Adam P. Ostendorf, Satyanarayana Gedela



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Adam P. Ostendorf MD and Satyanarayana Gedela MD, MRCP (UK)

Affiliations: Neurology Division, Department of Pediatrics, The Ohio State University, Columbus, OH and Nationwide Children's Hospital, Columbus, OH

Address correspondence to: Adam Ostendorf, M.D., Neurology Division, Department of Pediatrics, Nationwide Children's Hospital, 700 Children's Drive, Columbus, OH 43205. Phone: 614.722.4625; Fax 614.722.4633. Email: adam.ostendorf@nationwidechildrens.org.

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Abstract

The effect of epilepsy extends beyond those with the diagnosis and impacts families, communities and society. Caregiver and sibling quality of life is often negatively affected by frequent seizures, comorbid behavioral and sleep disorders and stigma surrounding the diagnosis. Furthermore, the negative impacts can be magnified by individual coping styles and resources available to families of those with epilepsy. Beyond the family and immediate caregivers, epilepsy impacts local communities by drawing additional resources from education systems. The direct costs of caring for an individual with epilepsy and the indirect costs associated with decreased productivity place financial strain on individuals and healthcare

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