

Short-form Article

# Classifying Neonatal and Infant Epilepsy Syndromes

The ILAE Task Force on Nosology and Definitions updates their 2017 guidelines for the classification of epilepsy syndromes in neonates and infants

## **Background**

The incidence of epilepsy is remarkably high in children up to 2 years of age, and often syndromic, with a significant mortality rate. However, the study and licensing of syndrome-specific anti-epileptics and development of gene therapies require the identification of electroclinical phenotypes of these conditions.

## What this study addresses

A revised classification for epilepsy syndromes with seizure onset before 2 years of age is proposed, with pointers for their effective diagnosis and clinical management.

# **Main findings**

The proposed classification discusses the emerging class of aetiology-specific epilepsies, which have genetic, structural, metabolic, immune, or infectious causes. These include self-limited epilepsy syndromes, in which there is a likelihood of spontaneous remission, and developmental and epileptic encephalopathies (DEEs), which involve developmental impairment or brain lesions resulting in epileptiform activity. Some other changes include:

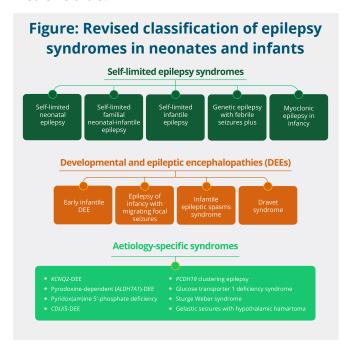
- 1. Incorporation of transparent clinically descriptive terms and merging of clinically related syndromes
- 2. Suggested changes to syndrome names that contain the terms "severe," "malignant," or "benign"
- 3. Replacing "partial seizures" with "focal seizures" and "convulsions" with "epilepsies" in some places
- 4. Certain newly coined nomenclatures

Lastly, the document tabulates seizure presentation, MRI, EEG, and neurological

examination features for common syndromes, and outlines the minimum criteria for syndrome diagnosis in underdeveloped areas.

### **Key takeaway**

The new classification aims to improve early diagnosis, optimal treatment, and long-term prognosis of epilepsy syndromes in neonates and infants, while ensuring readiness for precision medicine trials.



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