Opis choroby *

Definicja

A rare neonatal epilepsy syndrome characterized by seizures without specific underlying etiology, occurring during the first days of life in infants with an otherwise normal neurological state and no family history of neonatal convulsions. The most commonly partial and clonic seizures usually last for one to three minutes. Repeated seizures may lead to status epilepticus lasting up to 20 hours. Overall, remission rates are high and neurological outcome is favorable.

Dane

Klasyfikacja
Choroba

Synonimy
BINS
Benign nonfamilial neonatal seizures
BINS
Benign nonfamilial neonatal seizures

Kod ORPHA
64545

Kod OMIM
-

Kod ICD10
G40.3

Kod ICD11
8A6Y

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*Źródło

orphanet