Opis choroby *

Definicja

A rare infantile epilepsy syndrome characterized by infancy-onset of myoclonic seizures in otherwise neurologically and developmentally normal patients. Jerks may vary in severity, can be singular or occur in a series, and occur spontaneously or (less commonly) after sensory stimuli. Seizures are self-limiting and remit within several months to years from onset, although generalized tonic-clonic seizures or other forms of epilepsy may be seen later in life. Developmental delay and cognitive and behavioral difficulties have been reported in a considerable percentage of patients.

Dane

Klasyfikacja
Choroba

Synonimy
Benign myoclonic epilepsy of infancy
Dziecięca łagodna padaczka miokloniczna
Benign myoclonus epilepsy of infancy

Kod ORPHA
86909

Kod OMIM
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Kod ICD10
G40.3

Kod ICD11
8A61.1Y

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*Źródło

orphanet