

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

[*Źródło](#)

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