Opis choroby *

Definicja

A rare epilepsy syndrome characterized by onset of epileptic spasms in infants between 2 and 12 months of age, and rarely up to 24 months. Infants may have no antecedent history, or a history reflecting the underlying cause. The classical triad of epileptic spasms, hypsarrhythmia and developmental stagnation or regression is historically referred to as West syndrome.

Dane

Klasyfikacja
Zespół kliniczny

Synonimy
West syndrome
Drgawki dziecięce
Niepełnosprawność intelektualna - hipsarytmia

Kod ORPHA
3451

Kod OMIM
617065

Kod ICD10
G40.4

Kod ICD11
8A62.0

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*Źródło

orphanet