

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

*Źródło

orphanet