Opis choroby *

Definicja

A rare, genetic, infantile epilepsy syndrome disease characterized by neonatal- to infancy-onset myoclonic focal seizures occurring in various members of a family, associated in some with mild dysarthria, ataxia and borderline-to-moderate intellectual disability.

Dane

Klasyfikacja Synonimy Choroba FIME

FIME

Familial infantile myoclonus epilepsy

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 352582
 605021
 G40.3

Kod ICD11

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*Źródło

orphanet