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
Choroba

Synonimy
FIME
FIME
Familial infantile myoclonus epilepsy

Kod ORPHA
352582

Kod OMIM
605021

Kod ICD10
G40.3

Kod ICD11
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*Źródło

orphanet