Opis choroby *

Definicja

A rare, genetic, neurodevelopmental disorder characterized by early-onset of recurrent, transient episodes of hemiplegia (including quadriplegia), which typically disappear upon sleep.

Dane

Klasyfikacja
Choroba

Synonimy
AHC
AHC

Kod ORPHA
2131

Kod OMIM
614820

Kod ICD10
G98

Kod ICD11
MB53.0

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*Źródło

orphanet