

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

*Źródło

orphanet