

Opis choroby *

Definicja

Juvenile myoclonic epilepsy is the most common hereditary idiopathic generalized epilepsy syndrome and is characterized by myoclonic jerks of the upper limbs on awakening, generalized tonic-clonic seizures manifesting during adolescence and triggered by sleep deprivation, alcohol intake, and cognitive activities, and typical absence seizures (30% of cases).

Dane

Klasyfikacja

Choroba

Synonimy

JME

JME

Juvenile myoclonus epilepsy

Kod ORPHA

307

Kod OMIM

617924

Kod ICD10

G40.3

Kod ICD11

8A61.30

*Źródło

orphanet