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