

Rodzinny mioklonus korowy

Kod Orpha: 319189 Kod OMIM: 614937

Opis choroby *

Definicja

Familial cortical myoclonus is a rare, genetic movement disorder characterized by autosomal dominant, adult-onset, slowly progressive, multifocal, cortical myoclonus. Patients present somatosensory-evoked, brief, jerky, involuntary movements in the face, arms and legs, associated in most cases with sustained, multiple, sudden falls without loss of consciousness. Seizures or other neurological deficits, aside from mild cerebellar ataxia late in the course of the illness, are absent.

Dane

Klasyfikacja

Choroba

Kod ORPHA

319189

Kod OMIM

614937

Kod ICD10

G25.3

Kod ICD11

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[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.