

Choroba ciałek Lafora o wczesnym początku

Kod Orpha: 324290 Kod OMIM: 616640

Opis choroby *

Definicja

A rare genetic progressive myoclonic epilepsy characterized by childhood onset of progressive dysarthria, myoclonus, ataxia, seizures, and cognitive decline. The disease takes a protracted course with patients surviving into adulthood, developing signs and symptoms like psychosis with outbursts of prolonged agitation and screaming, spasticity and hyperreflexia, confusion, mutism, and incontinence. There are no visual disturbances. Muscle biopsy shows numerous periodic acid-Schiff-positive inclusions, so-called Lafora bodies.

Dane

Klasyfikacja

Choroba

Kod ORPHA

324290

Kod OMIM

616640

Kod ICD10

G40.3

Kod ICD11

8A61.41

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.