

Choroba podobna do choroby Huntingtona typu 1

Kod Orpha: 157941 Kod OMIM: 603218

Opis choroby *

Definicja

A rare, genetic, human prion disease characterized by adult-onset neurodegenerative manifestations associated with a movement disorder and psychiatric/behavioral disturbances. Patients typically present personality changes, aggressiveness, manias, anxiety and/or depression in conjunction with rapidly progressive cognitive decline (presenting with dysarthria, apraxia, aphasia, and eventually leading to dementia) as well as ataxia (manifesting with gait disturbances, unsteadiness, coordination problems), Parkinsonism, myoclonus, and/or chorea. Additional features may include generalized spasticity, seizures, urine incontinence and pyramidal abnormalities.

Dane

Klasyfikacja

Choroba

Synonimy

Early-onset prion disease with prominent psychiatric features

Choroba prionowa o wczesnym początku z wyraźnymi objawami psychicznymi

HDL1

HDL1

Kod ORPHA

157941

Kod OMIM

603218

Kod ICD10

G10

Kod ICD11

8A01.11

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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