Opis choroby *

Definicja

A rare form of genetic prion disease characterized by typical CJD features (rapidly progressive dementia, personality/behavioral changes, psychiatric disorders, myoclonus, and ataxia) with a genetic cause and sometimes a family history of dementia.

Dane

Klasyfikacja
Choroba

Synonimy
Inherited CJD
Dziedziczna CJD

Kod ORPHA
282166

Kod OMIM
123400

Kod ICD10
A81.0

Kod ICD11
8E02.0

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*Źródło

orphanet