Opis choroby *

Definicja

A group of human prion diseases characterized by progressive, invariably fatal neurodegeneration resulting from accidental transmission of prions. The group comprises iatrogenic Creutzfeldt-Jakob disease (CJD), which results from transmission of CJD prions in the course of medical procedures or treatments, and variant CJD (transmission via consumption of products from prion-diseased cows or via blood transfusion from an affected individual).

Dane

Klasyfikacja
Grupa fenomenów

Kod ORPHA
454700

Kod OMIM
123400

Kod ICD10
A81.0

Kod ICD11
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*Źródło

orphanet