Choroba podobna do choroby Huntingtona 2
Kod Orpha: 98934 Kod OMIM: 606438

Opis choroby *

Definicja
A rare severe neurodegenerative disorder that is considered one of the phenocopies of Huntington Disease (HD) affecting patients of African descent and characterized by a triad of movement (chorea, oculomotor, parkinsonism), psychiatric (prominently sadness, irritability and anxiety), and cognitive abnormalities (early cognitive decline and subcortical-like dementia).
Dane

Klasyfikacja
Choroba

Synonimy
HDL2
HDL2

Kod ORPHA
98934

Kod OMIM
606438

Kod ICD10
G10

Kod ICD11
8A01.11

---

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.