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

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*Źródło

orphanet