

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