Opis choroby *

Definicja

A rare, genetic, movement disorder characterized by early-onset, very slowly progressive choreiform movements that may involve variable parts of the body, typically aggravated by stress or anxiety, in various members of a family. Additional variable manifestations include hypotonia, often resulting in psychomotor delay (including gait disturbances) and dysarthria, as well as myoclonus, dystonia, behavioral symptoms (ADHD, obsessive-compulsive disorder), learning difficulties (particularly in writing) and spasticity with hyperreflexia and/or flexor/extensor plantar reflexes.

Dane

Klasyfikacja

Synonimy

Choroba

BHC

Rodzinna łagodna pląsawica

Benign familial chorea

Kod ORPHA

Kod OMIM

Kod ICD10

1429

215450

G25.5

Kod ICD11

8A01.0

*Źródło

orphanet