Opis choroby *

Definicja

A rare non-hereditary degenerative ataxia disease characterized by a slowly progressive cerebellar syndrome (with ataxia of stance and gait, upper limb dysmetria and intention tremor, ataxic speech, and oculomotor abnormalities), presenting in adulthood (at around 50 years of age), that is not due to a known cause. Extracerebellar symptoms (e.g., decreased vibration sense and absent or decreased ankle reflexes), polyneuropathy and mild autonomic dysfunction may also be present. Mild cognitive impairment has also rarely been reported.

Dane

Klasyfikacja
Choroba

Synonimy
Idiopathic late-onset cerebellar ataxia
Idiopatyczna ataksja móżdżkowa o późnym początku
SAOA
SAOA

Kod ORPHA
247234

Kod OMIM
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Kod ICD10
G31.8

Kod ICD11
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*Źródło

orphanet