Opis choroby *

Definicja

A rare, neurodegenerative disease characterized by progressive cognitive impairment, spastic tetraparesis, and cerebellar ataxia resulting from amyloid deposits in the brain. Spasticity with increased deep tendon reflexes and tone are early symptoms, muscular rigidity evolves later. Progressive mental deterioration usually starts with apathy and impaired memory with progression to complete disorientation.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Familial dementia, British type
Demencja rodzinna, typ brytyjski

Kod ORPHA
97345

Kod OMIM
176500

Kod ICD10
I68.0*

Kod ICD11
5D00.2Y

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*Źródło

orphanet