

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

*Źródło

orphanet