

## **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	Synonimy
Podtyp kliniczny	Familial dementia, British type Demencja rodzinna, typ brytyjski
<b>Kod ORPHA</b>	<b>Kod OMIM</b>
97345	176500
<b>Kod ICD11</b>	<b>Kod ICD10</b>
5D00.2Y	I68.0*

---

### \*Źródło

orphanet