

Opis choroby *

Definicja

A rare, neurodegenerative disease characterized by progressive cataracts, hearing loss, cerebellar ataxia, paranoid psychosis and dementia. Neuropathological features are diffuse atrophy of all parts of the brain, chronic diffuse encephalopathy and the presence of extremely thin and almost completely demyelinated cranial nerves.

Dane

Klasyfikacja	Synonimy
Podtyp kliniczny	Familial dementia, Danish type Rodzinna demencja, typ duński
Kod ORPHA	
97346	
Kod OMIM	
	117300
Kod ICD10	
	I68.0*
Kod ICD11	
5D00.2Y	

*Źródło

orphanet