

Opis choroby *

Definicja

A rare hereditary ataxia characterized by progressive truncal and limb ataxia resulting in gait instability. Dysarthria, dysphagia, nystagmus, spasticity of the lower limbs, mild peripheral sensory neuropathy, cognitive impairment and accelerated ageing have also been associated.

Dane

Klasyfikacja	Synonimy	
Choroba	SCAR16	
	Autosomalna recesywna ataksja rdzeniowo-mózdkowa typu 16	
	SCAR16	
	Spinocerebellar ataxia autosomal recessive type 16	
Kod ORPHA	Kod OMIM	Kod ICD10
412057	615768	G11.1
Kod ICD11	-	

*Źródło

orphanet