

Opis choroby *

Definicja

A genetically and clinically heterogeneous group of slowly progressive neurological disorders which in the pure form is characterized by pyramidal signs (weakness, spasticity, brisk tendon reflexes, and extensor plantar responses) predominantly affecting the lower limbs and with possible association of sphincter disturbances and deep sensory loss; and in the complex form by the addition of variable neurological or non-neurological features.

Dane

Klasyfikacja	Synonimy
Grupa fenomenów	Familial spastic paraplegia Choroba Strümpella i Lorraina HSP Rodzinna paraplegia spastyczna SPG HSP Hereditary spastic paraparesis SPG Strümpell-Lorrain disease

Kod ORPHA	Kod OMIM	Kod ICD10
685	-	G11.4

Kod ICD11
8B44.0

*Źródło

orphanet