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
Grupa fenomenów

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

Kod ORPHA
685

Kod OMIM
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Kod ICD10
G11.4

Kod ICD11
8B44.0

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*Źródło

orphanet