Opis choroby *

Definicja

A congenital, X-linked, clinical subtype of L1 syndrome, characterized by spastic paraplegia, mild to moderate intellectual disability and normal brain morphology. This subtype represents the milder end of the L1 syndrome spectrum.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
SPG1
SPG1

Kod ORPHA
306617

Kod OMIM
-

Kod ICD10
-

Kod ICD11
-

---

*Źródło

orphanet