

## 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

<b>Klasyfikacja</b>	Synonimy
Podtyp kliniczny	SPG1 SPG1

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
306617	-	-

**Kod ICD11**  
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### \*Źródło

orphanet