

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 61 (SPG61) is a rare, complex form of hereditary spastic paraplegia characterized by an onset in infancy of spastic paraplegia (presenting with the inability to walk unsupported and a scissors gait) associated with a motor and sensory polyneuropathy with loss of terminal digits and acropathy. SPG61 is due to a mutation in the *ARL6IP1* gene (16p12-p11.2) encoding the ADP-ribosylation factor-like protein 6-interacting protein 1.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG61

SPG61

#### Kod ORPHA

401780

#### Kod OMIM

615685

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.01

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#### \*Źródło

orphanet