

## Opis choroby \*

### Definicja

A rare hereditary ataxia characterized by unusual facies (i. e. gross, rough and abundant hair, mild palpebral ptosis, thick lips, and down-curved corners of the mouth), dysarthria, delayed psychomotor development, scoliosis, foot deformities, and ataxia. There have been no further descriptions in the literature since 1985.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

1185

#### Kod OMIM

271270

#### Kod ICD10

G11.8

#### Kod ICD11

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### \*Źródło

orphanet