

## Opis choroby \*

### Definicja

A rare demyelinating hereditary motor and sensory neuropathy characterized by prominent gait ataxia, pes cavus, tendon areflexia, distal limb weakness, tremor in the upper limbs, distal sensory loss, kyphoscoliosis, and progressive muscle atrophy. The disease becomes symptomatic in infancy or childhood, mode of inheritance is autosomal dominant.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Hereditary areflexic dystasia, Roussy-Lévy type  
Dziedziczna dystazja z brakiem odruchów, typ  
Roussy'ego i Lévy'ego

#### Kod ORPHA

3115

#### Kod OMIM

180800

#### Kod ICD10

G60.0

#### Kod ICD11

8C20.Y

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#### \*Źródło

orphanet