

## Opis choroby \*

### Definicja

A rare hereditary demyelinating motor and sensory neuropathy characterized by slowed nerve conduction velocities, in the absence of clinically apparent neurological deficits, gait abnormalities or muscular atrophy, associated with a germline mutation in the *ARGHEF10* gene.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

140481

#### Kod OMIM

608236

#### Kod ICD10

G60.0

#### Kod ICD11

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### \*Źródło

orphanet