

## Opis choroby \*

### Definicja

A rare, autosomal dominant, hereditary, demyelinating motor and sensory neuropathy which may present either as a classic Charcot-Marie-Tooth disease phenotype with distal motor weakness and wasting, gait difficulties, paresthesias, decreased vibration and pain sensation, or as a milder, predominantly sensory form with transient paresthesias, decreased sensation and distal pain in upper or lower limbs, without significant motor weakness. Pes cavus is a common feature, and additional symptoms may include hand tremor and decreased or absent deep tendon reflexes.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

CMT1C

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#### Kod ORPHA

101083

#### Kod OMIM

601098

#### Kod ICD10

G60.0

#### Kod ICD11

8C20.0

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#### \*Źródło

orphanet