

## Opis choroby \*

### Definicja

Charcot-Marie-Tooth disease type 4A (CMT4A) is a subtype of Charcot-Marie-Tooth disease type 4 characterized by early-onset (infancy to early childhood) of severe, rapidly progressing demyelinating, axonal, or intermediate sensorimotor neuropathy usually affecting first, and more severely, the distal lower extremities and later the proximal muscles and upper extremities. Nerve conduction velocities range from very slow to normal. Apart from the typical CMT phenotype (distal muscle weakness and atrophy, sensory loss, frequent pes cavus foot deformity), patients commonly present delayed motor development, vocal cord paresis, mild sensory loss, abolished deep tendon reflexes, and skeletal deformities.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

CMT4A

CMT4A

#### Kod ORPHA

99948

#### Kod OMIM

214400

#### Kod ICD10

G60.0

#### Kod ICD11

8C20.0

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#### [\\*Źródło](#)

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