

## Opis choroby \*

### Definicja

A rare genetic, congenital, non-dystrophic myopathy characterized by neonatal or infantile-onset hypotonia and mild to severe generalized muscle weakness.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

CFTDM

CFTDM

#### Kod ORPHA

2020

#### Kod OMIM

617760

#### Kod ICD10

G71.2

#### Kod ICD11

8C72.1

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#### \*Źródło

orphanet