## Opis choroby *

Definicja

A rare hereditary neuromuscular disorder characterized by multiple cores on muscle biopsy and clinical features of a congenital myopathy.

Dane

**Klasyfikacja**
Choroba

Synonimy
MmD
Choroba multiminicore
MmD
Multiminicore disease

**Kod ORPHA**
598

**Kod OMIM**
602771

**Kod ICD10**
G71.2

**Kod ICD11**
8C72.0Y

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*Źródło

orphanet