

## Opis choroby \*

### Definicja

A rare group of inherited neuromuscular disorders characterized by clinical features of a congenital myopathy and centrally placed nuclei on muscle biopsy. The clinical picture and other histologic features varies according to gene involved and mode of inheritance.

### Dane

<b>Klasyfikacja</b>	Synonimy
Grupa fenomenów	CNM CNM

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
595	-	G71.2

**Kod ICD11**  
8C72.01

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### \*Źródło

orphanet