

## Opis choroby \*

### Definicja

A form of limb-girdle muscular dystrophy that usually has a childhood onset (but can range from the first to third decade of life) of severe progressive proximal weakness, eventually involving the distal muscles. Some patients may remain ambulatory but most are wheelchair dependant 20 years after onset.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Autosomal recessive limb-girdle muscular dystrophy type 2J  
LGMD2J  
LGMD type 2J  
LGMD2J  
Limb-girdle muscular dystrophy type 2J  
Titin-related LGMD R10

#### Kod ORPHA

140922

#### Kod OMIM

608807

#### Kod ICD10

G71.0

#### Kod ICD11

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#### \*Źródło

orphanet