

## Opis choroby \*

### Definicja

A subtype of autosomal dominant limb-girdle muscular dystrophy characterized by an adult-onset of slowly progressive, proximal pelvic girdle weakness, with none, or only minimal, shoulder girdle involvement, and absence of cardiac and respiratory symptoms. Mild to moderate elevated creatine kinase serum levels and gait abnormalities are frequently observed.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Autosomal dominant limb-girdle muscular dystrophy type 1D  
LGMD1D  
DNAJB6-related LGMD D1  
LGMD type 1D  
LGMD1D  
Limb-girdle muscular dystrophy type 1D

#### Kod ORPHA

34516

#### Kod OMIM

603511

#### Kod ICD10

G71.0

#### Kod ICD11

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#### \*Źródło

orphanet