

## Opis choroby \*

### Definicja

A rare subtype of autosomal dominant limb-girdle muscular dystrophy characterized by slowly progressive proximal muscular weakness initially affecting the lower limbs (and later involving the upper limbs), hypotrophy of upper and lower limb-girdle muscles, hyporeflexia, calf hypertrophy, and increased serum creatine kinase. There is no involvement of oculo-facial-bulbar muscles and cardiac muscle.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Choroba	LGMD1H LGMD1H

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
238755	613530	G71.0

**Kod ICD11**  
-

---

### \*Źródło

orphanet