## Opis choroby *

Definicja

A rare, genetic, muscular dystrophy disease characterized by the co-occurrence of late onset scapular and peroneal muscle weakness, principally manifesting with distal lower limb and proximal upper limb weakness and scapular winging.

Dane

**Klasyfikacja**
Choroba

Synonimy
X-linked SPMD
SPMD sprzężony z chromosomem X
Zespół łopatkowo-strzałkowy sprzężony z chromosomem X
X-linked scapuloperoneal syndrome

**Kod ORPHA**
431272

**Kod OMIM**
300695

**Kod ICD10**
G71.0

**Kod ICD11**
8C70.5

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*Źródło

orphanet