DISEASE:
MYH7-related late-onset scapuloperoneal muscular dystrophy

NAME:	MYH7-related late-onset scapuloperoneal muscular dystrophy
ORPHACODE:	437572
SYNONYMS:	MYH7-related late-onset SPMD MYH7-related late-onset scapuloperoneal syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	MYH7
CREATED:	13 May 2019 - 01:02
CHANGED:	01 May 2022 - 06:55

RELATED CONTENT

Related Genetic Tests

- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- myosin heavy chain 7

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB