DISEASE:
X-linked scapuloperoneal muscular dystrophy

NAME:	X-linked scapuloperoneal muscular dystrophy
DESCRIPTION:	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.
ORPHACODE:	431272
SYNONYMS:	X-linked SPMD X-linked scapuloperoneal syndrome
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	FHL1
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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

- four and a half LIM domains 1

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB