DISEASE:
Distal myopathy, Welander type

NAME:	Distal myopathy, Welander type
DESCRIPTION:	A rare distal myopathy characterized by weakness in the distal upper extremities, usually finger and wrist extensors which later progresses to all hand muscles and distal lower extremity, primarily in toe and ankle extensors.
ORPHACODE:	603
SYNONYMS:	WDM
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	TIA1 SQSTM1 TIA1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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

- sequestosome 1
- TIA1 cytotoxic granule associated RNA binding protein

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB