

DISEASE:
Classic multiminicore myopathy

NAME:	Classic multiminicore myopathy
ORPHACODE:	324604
SYNONYMS:	Classic MmD Classic multiminicore disease
XREF(S):	<u>Orphanet</u> <u>ICD-10</u> <u>OMIM</u>
ANALYTE(S):	<u>SELENON</u> <u>TTN</u> <u>MYH7</u>
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/1688>

RELATED CONTENT

Related Genetic Tests

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

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- myosin heavy chain 7
- selenoprotein N
- titin

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB

Source URL: <http://gentest.healthdata.be/disease/1688>