DISEASE:
Duchenne muscular dystrophy

NAME:	Duchenne muscular dystrophy
DESCRIPTION:	A rare, genetic, muscular dystrophy characterized by rapidly progressive muscle weakness and wasting due to degeneration of skeletal, smooth and cardiac muscle.
ORPHACODE:	98896
SYNONYMS:	DMD Severe dystrophinopathy, Duchenne type
XREF(S):	Orphanet OMIM MeSH MedDRA ICD-10
ANALYTE(S):	DMD LTBP4
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Becker muscular dystrophy / Duchenne muscular dystrophy (Full sequencing DMD gene through Myopathy gene panel)
- Becker muscular dystrophy / Duchenne muscular dystrophy (deletion/duplication DMD gene)
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- Myopathy (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- dystrophin
- latent transforming growth factor beta binding protein 4

Related Gene Panels

- Myopathy (332 genes) - IPG
- Neuromuscular disorders (166 genes) - VUB
