Belgian Genetic Tests database

DISEASE:
Muscular dystrophy, Selcen type

NAME:	Muscular dystrophy, Selcen type
DESCRIPTION:	Selcen type muscular dystrophy is characterized by progressive limb and axial muscle weakness associated with cardiomyopathy and severe respiratory insufficiency during adolescence. The disease manifests during childhood and progresses rapidly.
ORPHACODE:	199340
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	BAG3
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Source URL: http://gentest.healthdata.be/disease/436

RELATED CONTENT

Related Genetic Tests

- Cardiopathies, hereditary (gene panel)
- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Peripheral neuropathy (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- BAG cochaperone 3

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Cardiopathies, hereditary (102 genes) - KUL
- Neuromuscular disorders (166 genes) - VUB
- Neuropathy (genepanel) - UZA
