Belgian Genetic Tests database

DISEASE:
Symptomatic form of muscular dystrophy of Duchenne and Becker in female carriers

NAME:	Symptomatic form of muscular dystrophy of Duchenne and Becker in female carriers
DESCRIPTION:	A rare, genetic muscular dystrophy affecting female carriers and characterized by variable degrees of muscle weakness due to progressive skeletal myopathy, sometimes associated with dilated cardiomyopathy or left ventricle dilation.
ORPHACODE:	206546
XREF(S):	Orphanet
ANALYTE(S):	DMD
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Source URL: http://gentest.healthdata.be/disease/1314

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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- Becker muscular dystrophy / Duchenne muscular dystrophy (deletion/duplication DMD gene)
- Becker muscular dystrophy / Duchenne muscular dystrophy (deletion/duplication DMD gene)

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 Menselijke Erfelijkheid - KUL

Related Analytes

- dystrophin