GENETIC TEST:
Becker muscular dystrophy / Duchenne muscular dystrophy (deletion/duplication DMD gene)

FULL NAME:	Becker muscular dystrophy / Duchenne muscular dystrophy (deletion/duplication DMD gene)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Chorionic villi, Amniotic fluid
METHOD CATEGORY:	Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460

EQA:	• Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy
TURNAROUND TIME (MAXIMUM):	6 - 13 weeks
CREATED:	27 Jul 2018 - 11:04
CHANGED:	01 Mar 2023 - 14:42
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/12521

RELATED CONTENT

Related Diseases

- Becker muscular dystrophy
- Duchenne muscular dystrophy
- Symptomatic form of muscular dystrophy of Duchenne and Becker in female carriers

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- dystrophin

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