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
ACCREDITATION (ISO 15189):	2023-11-09 / 2024-05-08

EQA:	• Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy, • Muscular dystrophies
TURNAROUND TIME (MAXIMUM):	2 months
CREATED:	27 Jul 2018 - 11:04
CHANGED:	22 Jan 2024 - 11:36
URL:	https://labogidsmedgen.uza.be/analyses/duchenne-becker-spierdystrofie-stap-1-pa…

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 Medische Genetica - UZ Antwerpen

Related Analytes

- dystrophin