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, Amniotic fluid, Chorionic villi
METHOD CATEGORY:	Deletion/duplication analysis
METHOD TECHNIQUE:	MLPA based techniques
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23

EQA:	• Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy, • Duchenne / Becker Muscular Dystrophy
TURNAROUND TIME (MAXIMUM):	2 weeks
CREATED:	27 Jul 2018 - 11:04
CHANGED:	23 Aug 2023 - 15:49
URL:	https://www.chuliege.be/jcms/c_525333/fr/myopathie-de-duchenne

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RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

Related Analytes

- dystrophin