GENETIC TEST:
Steinert myotonic dystrophy - DMPK gene CTG repeat expansion

FULL NAME:	Steinert myotonic dystrophy - DMPK gene CTG repeat expansion
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Mutation confirmation, Post-natal Diagnosis, Pre-implantation genetic diagnosis, Predictive and Pre-symptomatic diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi, Cell culture
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique Southern blot
RIZIV CODE:	565456-565460

ACCREDITATION (ISO 15189):	2021-10-07 / 2026-06-14
EQA:	• Myotonic Dystrophy , • Myotonic Dystrophy, • Myotonic Dystrophy , • Myotonic Dystrophy, • Myotonic Dystrophy, • Myotonic Dystrophy, • Myotonic Dystrophy
TURNAROUND TIME (MAXIMUM):	3 months (10 working days for prenatal diagnosis)
CREATED:	31 Jul 2019 - 13:59
CHANGED:	09 Mar 2023 - 16:31
URL:	https://laboguide.uzbrussel.be/laboguide#Analyses:ziekte%20van%20Steinert&&&&&1…

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