GENETIC TEST:
Steinert myotonic dystrophy - CTG repeat expansion

FULL NAME:	Steinert myotonic dystrophy - CTG repeat expansion
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:	Targeted variant analysis
METHOD TECHNIQUE:	PCR based technique Southern blot
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• Myotonic Dystrophy , • Myotonic Dystrophy, • Myotonic Dystrophy, • Myotonic Dystrophy
TURNAROUND TIME (MAXIMUM):	6 - 8 weeks or 16 - 24 weeks
CREATED:	22 Jul 2019 - 10:14
CHANGED:	01 Mar 2023 - 16:08
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/12204

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