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:	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
RIZIV CODE:	565456-565460
ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23

EQA:	• Myotonic Dystrophy , • Myotonic Dystrophy, • Myotonic Dystrophy
TURNAROUND TIME (MAXIMUM):	2 mois
CREATED:	31 Jul 2019 - 13:59
CHANGED:	23 Jan 2023 - 13:37
URL:	https://www.chuliege.be/jcms/c_525336/fr/myopathie-de-steinert

RELATED CONTENT

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- Steinert myotonic dystrophy

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- Centre de Génétique Humaine - CHU Sart-Tilman

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- DM1 protein kinase