GENETIC TEST:
Fragile X syndrome/FXPOI/FXTAS - FMR1 CGG repeat expansion

FULL NAME:	Fragile X syndrome/FXPOI/FXTAS - FMR1 CGG 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
RIZIV CODE:	565375-565386
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• Fragile X Syndrome , • Fragile X Syndrome , • Fragile X Syndrome, • Fragile X Syndrome, • Fragile X Syndrome, • Fragile X Syndrome , • Fragile X Syndrome
TURNAROUND TIME (MAXIMUM):	6 - 8 weeks
CREATED:	19 Jul 2019 - 10:55
CHANGED:	04 Dec 2023 - 12:27
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB/16838

RELATED CONTENT

Related Diseases

- Fragile X syndrome
- Fragile X-associated tremor/ataxia syndrome
- Symptomatic form of fragile X syndrome in female carriers

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- fragile X messenger ribonucleoprotein 1
