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

FULL NAME:	Fragile X syndrome/POF/FXTAS - 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):	2021-09-11 / 2026-09-10

EQA:	• Fragile X Syndrome , • Fragile X Syndrome, • Fragile X Syndrome
TURNAROUND TIME (MAXIMUM):	6 weeks
CREATED:	19 Jul 2019 - 10:55
CHANGED:	19 May 2022 - 14:51

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 Medische Genetica - UZ Gent

Related Analytes

- fragile X messenger ribonucleoprotein 1