GENETIC TEST:
Familial hypercholanemia (3 genes)

FULL NAME:	Familial hypercholanemia (3 genes)
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:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2022-12-22 / 2027-12-21
TURNAROUND TIME (MAXIMUM):	3 months

CREATED:	06 Nov 2019 - 12:04
CHANGED:	24 Jan 2023 - 15:06

Source URL: http://gentest.healthdata.be/genetic_test/923

RELATED CONTENT

Related Diseases

- Familial hypercholanemia

Related Laboratories

- Centre de Génétique Médicale UCL

Related Analytes

- bile acid-CoA:amino acid N-acyltransferase
- epoxide hydrolase 1
- tight junction protein 2

Related Gene Panels

- Familial hypercholanemia (3 genes) - UCL