

GENETIC TEST: Hypercholesterolemia, Familial (gene panel)

FULL NAME:	Hypercholesterolemia, Familial (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Predictive and Pre-symptomatic diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS) MLPA based techniques
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2023-11-09 / 2024-05-08
EQA:	• Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia

TURNAROUND TIME (MAXIMUM):	4 months
CREATED:	19 Jul 2019 - 13:01
CHANGED:	22 Jan 2024 - 13:50
URL:	https://labogidsmedgen.uza.be/analyses/familiale-hypercholesterolemie-chol

Source URL: http://gentest.healthdata.be/genetic_test/530

RELATED CONTENT

Related Diseases

- [Cholesteryl ester storage disease](#)
- [Heterozygous familial hypercholesterolemia \(NON RARE IN EUROPE\)](#)
- [Homozygous familial hypercholesterolemia](#)

Related Laboratories

- [Centrum Medische Genetica - UZ Antwerpen](#)

Related Analytes

- [ATP binding cassette subfamily G member 5](#)
- [ATP binding cassette subfamily G member 8](#)
- [apolipoprotein B](#)
- [apolipoprotein E](#)
- [low density lipoprotein receptor](#)
- [low density lipoprotein receptor adaptor protein 1](#)
- [lipase A, lysosomal acid type](#)
- [proprotein convertase subtilisin/kexin type 9](#)

Related Gene Panels

- Familial Hypercholesterolemia panel (8 genes) - UZA

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