

GENETIC TEST: Hypercholesterolemia, Familial (9 genes)

FULL NAME:	Hypercholesterolemia, Familial (9 genes)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal 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):	2022-02-24 / 2026-02-23
EQA:	• Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia

TURNAROUND TIME (MAXIMUM):	3 months
CREATED:	19 Jul 2019 - 13:01
CHANGED:	01 Dec 2023 - 13:43
URL:	https://www.chuliege.be/upload/docs/application/pdf/2019-04/mq.a11.24_-_genetiq...

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RELATED CONTENT

Related Diseases

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

Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)

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](#)
- [signal transducing adaptor family member 1](#)

Related Gene Panels

- Familial Hypercholesterolemia panel (9 genes) - ULG

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