GENETIC TEST:
Hypercholesterolemia, Familial (4 genes)

FULL NAME:	Hypercholesterolemia, Familial (4 genes)
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)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02

EQA:	• Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia, • Familial autosomal dominant hypercholesterolemia
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	19 Jul 2019 - 13:01
CHANGED:	01 Mar 2023 - 15:11
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/12435

RELATED CONTENT

Related Diseases

- Homozygous familial hypercholesterolemia

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- apolipoprotein B
- apolipoprotein E
- low density lipoprotein receptor
- proprotein convertase subtilisin/kexin type 9

Related Gene Panels

- Familial Hypercholesterolemia panel (4 genes) - KUL