

## GENETIC TEST: Dyslipidemia (gene panel)

<b>FULL NAME:</b>	Dyslipidemia (gene panel)
<b>TEST TYPE:</b>	Clinical
<b>TEST SPECIALTY:</b>	Molecular Genetics
<b>TEST PURPOSE:</b>	Mutation confirmation, Post-natal Diagnosis
<b>SPECIMEN:</b>	Peripheral (whole) blood on EDTA
<b>METHOD CATEGORY:</b>	Sequence analysis: entire coding region Deletion/duplication analysis
<b>METHOD TECHNIQUE:</b>	Next Generation Sequencing (NGS)
<b>RIZIV CODE:</b>	565471-565482
<b>TURNAROUND TIME (MAXIMUM):</b>	3 months
<b>CREATED:</b>	13 Dec 2019 - 10:56
<b>CHANGED:</b>	17 Aug 2023 - 13:38
<b>URL:</b>	<a href="https://ulbgenetics.be/compendium-des-analyses/#GM33">https://ulbgenetics.be/compendium-des-analyses/#GM33</a>

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Source URL: [http://gentest.healthdata.be/index.php/index.php/genetic\\_test/970](http://gentest.healthdata.be/index.php/index.php/genetic_test/970)

## RELATED CONTENT

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### Related Diseases

- [Cholesteryl ester storage disease](#)
- [Familial GPIHBP1 deficiency](#)
- [Familial apolipoprotein A5 deficiency](#)
- [Familial apolipoprotein C-II deficiency](#)
- [Familial lipase maturation factor 1 deficiency](#)
- [Familial lipoprotein lipase deficiency](#)
- [Homozygous familial hypercholesterolemia](#)
- [Sitosterolemia](#)
- [Wolman disease](#)

### Related Laboratories

- [Centre de Génétique Humaine - Erasme ULB](#)

### Related Analytes

- [ATP binding cassette subfamily G member 5](#)
- [ATP binding cassette subfamily G member 8](#)
- [apolipoprotein A5](#)
- [apolipoprotein B](#)
- [apolipoprotein C2](#)
- [apolipoprotein E](#)
- [glycosylphosphatidylinositol anchored high density lipoprotein binding protein 1](#)

- low density lipoprotein receptor
- low density lipoprotein receptor adaptor protein 1
- lipase A, lysosomal acid type
- lipase maturation factor 1
- lipoprotein lipase
- proprotein convertase subtilisin/kexin type 9

## Related Gene Panels

- Dyslipidemia ( 13 genes) - ULB

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