

GENETIC TEST:
Lipodystrophy and/or hyperinsulinism (gene panel)

FULL NAME:	Lipodystrophy and/or hyperinsulinism (gene panel)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Carrier diagnosis, Newborn screening, Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, DNA
METHOD CATEGORY:	Sequence analysis: entire coding region Deletion/duplication analysis
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
TURNAROUND TIME (MAXIMUM):	20 to 60 days
CREATED:	14 Dec 2022 - 14:12

CHANGED:	11 Dec 2023 - 12:54
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Source URL: http://gentest.healthdata.be/genetic_test/1091

RELATED CONTENT

Related Laboratories

- [Centre de Génétique-Institut de Pathologie et de Génétique \(IPG\)](#)

Related Analytes

- [ATP binding cassette subfamily C member 8](#)
- [1-acylglycerol-3-phosphate O-acyltransferase 2](#)
- [AKT serine/threonine kinase 2](#)
- [ALMS1 centrosome and basal body associated protein](#)
- [BLM RecQ like helicase](#)
- [BSCL2 lipid droplet biogenesis associated, seipin](#)
- [caveolin 1](#)
- [caveolae associated protein 1](#)
- [cell death inducing DFFA like effector c](#)
- [glucokinase](#)
- [glutamate dehydrogenase 1](#)
- [hydroxyacyl-CoA dehydrogenase](#)
- [HNF1 homeobox A](#)
- [hepatocyte nuclear factor 4 alpha](#)
- [insulin receptor](#)
- [potassium inwardly rectifying channel subfamily J member 11](#)
- [lipase E, hormone sensitive type](#)
- [lamin A/C](#)
- [MAF bZIP transcription factor A](#)
- [pericentrin](#)

- phosphate cytidyltransferase 1A, choline
- phosphoinositide-3-kinase regulatory subunit 1
- phospholipase A and acyltransferase 3
- DNA polymerase delta 1, catalytic subunit
- peroxisome proliferator activated receptor gamma
- protein phosphatase 1 regulatory subunit 3A
- solute carrier family 16 member 1
- uncoupling protein 2
- WRN RecQ like helicase
- zinc metalloproteinase STE24

Related Gene Panels

- Lipodystrophy and/or hyperinsulinism (30 genes) - IPG

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