GENETIC TEST:
Endocrine Disorders - Hypothyroidism (gene panel - 42 genes)

FULL NAME:	Endocrine Disorders - Hypothyroidism (gene panel - 42 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 Whole Exome Sequencing (WES)
METHOD TECHNIQUE:	Next Generation Sequencing (NGS)
RIZIV CODE:	565493-565504
TURNAROUND TIME (MAXIMUM):	6 months
CREATED:	14 Dec 2021 - 13:03
CHANGED:	14 Dec 2022 - 13:10

RELATED CONTENT

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB

Related Analytes

- dual oxidase 2
- dual oxidase maturation factor 2
- forkhead box E1
- growth arrest specific 1
- GLIS family zinc finger 3
- G protein subunit alpha 11
- GNAS complex locus
- GNAS antisense RNA 1
- HESX homeobox 1
- immunoglobulin superfamily member 1
- iodotyrosine deiodinase
- LIM homeobox 3
- LIM homeobox 4
- NK2 homeobox 1
- NK2 homeobox 5
- orthodenticle homeobox 2
- paired box 8
- POU class 1 homeobox 1
- protein kinase cAMP-dependent type I regulatory subunit alpha
- PROP paired-like homeobox 1

- SECIS binding protein 2
- solute carrier family 16 member 2
- solute carrier family 26 member 4
- solute carrier family 5 member 5
- SRY-box transcription factor 3
- syntaxin 16
- transducin beta like 1 X-linked
- thyroglobulin
- thrombopoietin
- thyroid hormone receptor alpha
- thyroid hormone receptor beta
- thyrotropin releasing hormone receptor
- thyroid stimulating hormone subunit beta
- thyroid stimulating hormone receptor

Related Gene Panels

- Endocrine Disorders - Hypothyroidism (42 genes) - ULB
