sciensano **Belgian Genetic Tests database**

GENETIC TEST:
Hypocalciuric Hypercalcemia, Neonatal Severe Hyperparathyroidism, Hypocalcemia

FULL NAME:	Hypocalciuric Hypercalcemia, Neonatal Severe Hyperparathyroidism, Hypocalcemia
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)
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2021-07-08 / 2026-02-02
TURNAROUND TIME (MAXIMUM):	4 - 6 months
CREATED:	19 Jul 2019 - 15:54
CHANGED:	01 Mar 2023 - 15:13
URL:	https://laboboeken.nexuzhealth.com/pboek/internet/GHB%20CME/13388

RELATED CONTENT

Related Diseases

- Autosomal dominant hypocalcemia
- Familial hypocalciuric hypercalcemia type 1
- Neonatal severe primary hyperparathyroidism

Related Laboratories

- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- calcium sensing receptor