GENETIC TEST:
Hypocalciuric hypercalcemia, familial type I or Hypocalcemia or Hyperparathyroidism, familial isolated (CASR gene)

FULL NAME:	Hypocalciuric hypercalcemia, familial type I or Hypocalcemia or Hyperparathyroidism, familial isolated (CASR gene)
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
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565471-565482
ACCREDITATION (ISO 15189):	2022-02-24 / 2026-02-23
EQA:	• Calcium disorders
TURNAROUND TIME (MAXIMUM):	1 month

CREATED:	31 Aug 2019 - 19:40
CHANGED:	16 Jan 2024 - 15:06
URL:	https://www.chuliege.be/jcms/c_5255550/fr/genotypage-des-mutations-du-gene-casr…

Source URL: http://gentest.healthdata.be/genetic_test/853

RELATED CONTENT

Related Diseases

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

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman

Related Analytes

- calcium sensing receptor