

DISEASE:
Familial hypocalciuric hypercalcemia type 1

NAME:	Familial hypocalciuric hypercalcemia type 1
ORPHACODE:	93372
SYNONYMS:	FHH type 1
XREF(S):	<u>Orphanet</u> <u>MeSH</u> <u>MedDRA</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>CASR</u>
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RELATED CONTENT

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- [Endocrine Disorders - Hyper\(Hypo\)parathyroidism \(gene panel - 24 genes\)](#)
- [Hypocalciuric Hypercalcemia, Neonatal Severe Hyperparathyroidism, Hypocalcemia](#)
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- [Hypocalciuric hypercalcemia, familial type I or Hypocalcemia or Hypoparathyroidism, familial isolated \(CASR gene\)](#)
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Related Laboratories

- [Centre de Génétique Humaine - CHU Sart-Tilman](#)
- [Centre de Génétique Humaine - Erasme ULB](#)
- [Centre de Génétique-Institut de Pathologie et de Génétique \(IPG\)](#)
- [Centrum Medische Genetica - UZ Brussel VUB](#)
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Related Analytes

- [calcium sensing receptor](#)

Related Gene Panels

- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Thyroid dysgenesis (38 genes) - VUB
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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