Belgian Genetic Tests database

ANALYTE:
GNA11

NAME:	G protein subunit alpha 11
SYMBOL:	GNA11
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FBH FBH2 FHH2
XREF(S):	Orphanet Reactome SwissProt Ensembl Genatlas HGNC OMIM
CREATED:	13 May 2019 - 01:01
CHANGED:	26 Oct 2023 - 23:49

Source URL: http://gentest.healthdata.be/analyte/2736

RELATED CONTENT

Related Genetic Tests

- Congenital hemangioma (2 genes)
- Congenital malformation (gene panel - 1721 genes)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Endocrine Disorders - Hypothyroidism (gene panel - 42 genes)
- Genetic disorders of Calcium and Phosphate metabolism (gene panel)
- Hypocalciuric hypercalcemia, familial type II
- Inherited Kidney Diseases (Gene Panel)
- Maffucci syndrome (gene panel)
- Nephrocalcinosis and nephrolithiasis (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Overgrowth & vascular anomalies (gene panel)
- Sturge-Weber syndrome (gene panel)
- Thyroid disgenesis (38 genes)
- Tubulopathy (gene panel)
- Vascular malformations (somatic)

Related Diseases

- Autosomal dominant hypocalcemia
- Cutis marmorata telangiectatica congenita
- Familial hypocalciuric hypercalcemia type 2
- Phakomatosis cesioflammea
- Phakomatosis cesiomarmorata

- Uveal melanoma

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Endocrine Disorders - Hypothyroidism (42 genes) - ULB
- Genetic disorders of Calcium and Phosphate metabolism (31 genes) - KUL
- Maffucci syndrome (65 genes) - KUL
- Nephrocalcinosis and nephrolithiasis (37 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Overgrowth & vascular anomalies (65 genes) - KUL
- Panel Nephro-ULG-V1
- Sturge-Weber syndrome (65 genes) - KUL
- Thyroid disgenesis (38 genes) - VUB
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
- Vascular malformations (somatic) (19 genes) - UCL
