Belgian Genetic Tests database

ANALYTE:
TRPV6

NAME:	transient receptor potential cation channel subfamily V member 6
SYMBOL:	TRPV6
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CaT1
XREF(S):	Orphanet HGNC Ensembl SwissProt OMIM
CREATED:	13 May 2019 - 01:01
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Source URL: http://gentest.healthdata.be/index.php/index.php/analyte/733

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Neurodevelopmental disorders (1300 genes)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Neonatal severe primary hyperparathyroidism

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Neurodevelopmental disorders (1300 genes) - ULB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent