Belgian Genetic Tests database

ANALYTE:
GCM2

NAME:	glial cells missing transcription factor 2
SYMBOL:	GCM2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	hGCMb
XREF(S):	Orphanet HGNC OMIM SwissProt Ensembl Genatlas Reactome
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Source URL: http://gentest.healthdata.be/analyte/1765

RELATED CONTENT

Related Genetic Tests

- Brain malformations (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Epilepsy gene panel
- Genetic disorders of Calcium and Phosphate metabolism (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Familial isolated hyperparathyroidism
- Familial isolated hypoparathyroidism due to agenesis of parathyroid gland

Related Gene Panels

- Brain malformations (34 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Epilepsy gene panel - VUB
- Genetic disorders of Calcium and Phosphate metabolism (31 genes) - KUL
- Skeletal dysplasia (genepanel) - UZA