Belgian Genetic Tests database

ANALYTE:
FAM111A

NAME:	FAM111 trypsin like peptidase A
SYMBOL:	FAM111A
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ22794 KIAA1895
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Craniosynostosis (gene panel)
- Endocrine Disorders - Hyper(Hypo)parathyroidism (gene panel - 24 genes)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Autosomal dominant Kenny-Caffey syndrome
- Osteocraniostenosis

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Craniosynostosis (32 genes) - KUL
- Endocrine Disorders - Hyper(Hypo)parathyroidism (24 genes) - ULB
- Intellectual disability & Epilepsy - UGent

- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
