ANALYTE:
MEGF8

NAME:	multiple EGF like domains 8
SYMBOL:	MEGF8
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	FLJ22365 HBV pre s2 binding protein 1 SBP1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- Craniosynostosis (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Primary ciliary dyskinesia (PCD) Heterotaxyies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Carpenter syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- Craniosynostosis (32 genes) - KUL
- Heterotaxie PCD - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA

- Skeletal dysplasia - UGent
