

ANALYTE:
FGFR1

NAME:	fibroblast growth factor receptor 1
SYMBOL:	FGFR1
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SYNONYMS:	BFGFR CD331 CEK FLG H2 H3 H4 H5 N-SAM Pfeiffer syndrome

XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Related Gene Panels

- [Brain malformations \(34 genes\) - ULB](#)
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- [Congenital malformation \(1721 genes\) - ULB](#)
- [Congenital malformation gene panel - VUB](#)
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- [Craniosynostosis \(32 genes\) - KUL](#)
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- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Hypogonadotropic hypogonadism (33 genes) - VUB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Overgrowth & vascular anomalies (65 genes) - KUL
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
- Sturge-Weber syndrome (65 genes) - KUL

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