ANALYTE:
FGFR2

NAME:	fibroblast growth factor receptor 2
SYMBOL:	FGFR2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CD332 CEK3 Crouzon syndrome ECT1 K-SAM Pfeiffer syndrome TK14 TK25
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Craniosynostosis (gene panel)
- Craniosynostosis / Apert syndrome (hot spot mutations - exon 7)
- Craniosynostosis syndromes (Apert, Crouzon)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Disorders of sex development - Primary Ovarian insufficiency - Hypogonadotropic Hypogonadism (gene panel)
- Epidermal nevus syndrome (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Maffucci syndrome (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Overgrowth & vascular anomalies (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Sturge-Weber syndrome (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Antley-Bixler syndrome
- Antley-Bixler syndrome without genital anomaly or disorder of steroidogenesis
- Apert syndrome
- Crouzon syndrome
- Cutis gyrata-acanthosis nigricans-craniosynostosis syndrome
- FGFR2-related bent bone dysplasia
- Familial scaphocephaly syndrome, McGillivray type
- Jackson-Weiss syndrome
- Lacrimoauriculodentodigital syndrome
- Pfeiffer syndrome type 1
- Pfeiffer syndrome type 2
- Pfeiffer syndrome type 3
- Saethre-Chotzen syndrome

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
- Congenital structural heart defects - UGent
- Craniosynostosis (32 genes) - KUL
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Disorders of Sex Development - Primary Ovarian Insufficiency - Hypogonadotropic Hypogonadism - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Maffucci syndrome (65 genes) - KUL
- 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
- Sturge-Weber syndrome (65 genes) - KUL
- epidermal nevus syndrome (65 genes) - KUL
