ANALYTE:
FN1

NAME:	fibronectin 1
SYMBOL:	FN1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CIG Cold-insoluble globulin FINC GFND2 LETS MSF Migration-stimulating factor cold-insoluble globulin migration-stimulating factor

XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital structural heart defects (gene panel)
- End-stage renal disease, ESRD (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Fibronectin glomerulopathy
- Spondylometaphyseal dysplasia, 'corner fracture' type

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital structural heart defects - UGent
- End-stage renal disease (106 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1

- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA

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