ANALYTE:
LRP5

NAME:	LDL receptor related protein 5
SYMBOL:	LRP5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BMND1 EVR4 HBM LR3 OPS OPTA1 VBCH2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Familial Exudative Vitreoretinopathy, autosomal dominant
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Osteogenesis Imperfecta (gene panel)
- Osteogenesis imperfecta / Osteoporose (gene panel)
- Retinal dystrophy / RETNET (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Autosomal dominant osteopetrosis type 1
- Endosteal hyperostosis, Worth type
- Familial exudative vitreoretinopathy
- Hyperostosis corticalis generalisata
- Isolated polycystic liver disease
- LRP5-related primary osteoporosis

- Osteoporosis-pseudoglioma syndrome
- Osteosclerosis-developmental delay-craniosynostosis syndrome
- Retinopathy of prematurity

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Exudative Vitreoretinopathy - UGent
- Hepatology panel - UGent
- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Osteogenesis Imperfecta (25 genes) - KUL
- Osteogenesis imperfecta and Osteoporosis (43 genes) - UGent
- Osteoporosis (3 genes)
- Panel Nephro-ULG-V1
- Retinal dystrophy - UGent
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
