ANALYTE:
B9D1

NAME:	B9 domain containing 1
SYMBOL:	B9D1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	B9 EPPB9 MKS9 MKSR-1 endothelial precursor protein B9
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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Source URL: http://gentest.healthdata.be/analyte/485

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ciliopathy (gene panel)
- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Joubert syndrome
- Meckel syndrome

Related Gene Panels

- Ataxia (348 genes) - ULB
- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Intellectual disability & Epilepsy - UGent

- Intellectual disability (gene panel)
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia - UGent
