

**ANALYTE:  
B9D2**

|                             |                                                                                                                                                                                          |
|-----------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| <b>NAME:</b>                | B9 domain containing 2                                                                                                                                                                   |
| <b>SYMBOL:</b>              | B9D2                                                                                                                                                                                     |
| <b>VERSION OF ORPHANET:</b> | 2023-06-22 14:14:43                                                                                                                                                                      |
| <b>SYNONYMS:</b>            | MGC4093<br>MKS10<br>MKSR-2                                                                                                                                                               |
| <b>XREF(S):</b>             | <a href="#">Orphanet</a><br><a href="#">Reactome</a><br><a href="#">SwissProt</a><br><a href="#">Ensembl</a><br><a href="#">Genatlas</a><br><a href="#">HGNC</a><br><a href="#">OMIM</a> |
| <b>CREATED:</b>             | 13 May 2019 - 01:01                                                                                                                                                                      |
| <b>CHANGED:</b>             | 26 Oct 2023 - 23:49                                                                                                                                                                      |

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## RELATED CONTENT

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### Related Genetic Tests

- [Ciliopathy \(gene panel\)](#)
- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophthisis / 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 \(gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)

### Related Diseases

- [Joubert syndrome](#)
- [Meckel syndrome](#)

### Related Gene Panels

- Ciliopathy (120 genes) - UGent
- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophthisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Skeletal dysplasia (genepanel) - UZA

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