

**ANALYTE:**  
**CRB2**

<b>NAME:</b>	crumbs cell polarity complex component 2
<b>SYMBOL:</b>	CRB2
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	FLJ16786 FLJ38464
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Ciliopathy \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [End-stage renal disease, ESRD \(gene panel\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Nephrotic syndrome, Focal Segmental Glomerulosclerosis \(FSGS\) , Alport syndrome and podocytopathy \(gene panel\)](#)

### Related Diseases

- [Familial idiopathic steroid-resistant nephrotic syndrome with focal segmental hyalinosis](#)
- [Genetic steroid-resistant nephrotic syndrome](#)
- [Ventriculomegaly-cystic kidney disease](#)

### Related Gene Panels

- [Ciliopathy \(120 genes\) - UGent](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [End-stage renal disease \(106 genes\) - IPG](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)

- [Nephropathy panel - UGent](#)
  - [Nephrotic syndrome, FSGS, Alport syndrome \(76 genes\) - IPG](#)
  - [Panel Nephro-ULG-V1](#)
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