

**ANALYTE:**  
**CA2**

|                             |                                                                                                                           |
|-----------------------------|---------------------------------------------------------------------------------------------------------------------------|
| <b>NAME:</b>                | carbonic anhydrase 2                                                                                                      |
| <b>SYMBOL:</b>              | CA2                                                                                                                       |
| <b>VERSION OF ORPHANET:</b> | 2023-06-22 14:14:43                                                                                                       |
| <b>SYNONYMS:</b>            | CA-II<br>CAII<br>Car2                                                                                                     |
| <b>XREF(S):</b>             | <u>Orphanet</u><br><u>Ensembl</u><br><u>Genatlas</u><br><u>HGNC</u><br><u>OMIM</u><br><u>Reactome</u><br><u>SwissProt</u> |
| <b>CREATED:</b>             | 13 May 2019 - 01:01                                                                                                       |
| <b>CHANGED:</b>             | 22 Jun 2023 - 16:14                                                                                                       |

## RELATED CONTENT

---

### Related Genetic Tests

- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Intellectual disability & Epilepsy \(gene panel\)](#)
- [Intellectual disability \(virtual gene panel\)](#)
- [Nephrocalcinosis and nephrolithiasis \(gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
- [Tubulopathy \(gene panel\)](#)

### Related Diseases

- [Osteopetrosis with renal tubular acidosis](#)

### Related Gene Panels

- [Congenital malformation \(1721 genes\) - ULB](#)
- [Intellectual disability & Epilepsy - UGent](#)
- [Intellectual disability \(gene panel\)](#)
- [Nephrocalcinosis and nephrolithiasis \(37 genes\) - IPG](#)

- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

---

Source URL: <http://gentest.healthdata.be/analyte/766>