

**ANALYTE:**  
**CFHR1**

<b>NAME:</b>	complement factor H related 1
<b>SYMBOL:</b>	CFHR1
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	CFHL FHR1 H36-1 H36-2
<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> <a href="#">Reactome</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

---

### Related Genetic Tests

- [Atypical Hemolytic Uremic Syndrome \(aHUS\) \(gene panel\)](#)
- [Hereditary Hemolytic Anemias due to unknown or doubtful origin \(gene panel - 52 genes\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)
- [Primary immune deficiencies \(gene panel\)](#)

### Related Diseases

- [Atypical hemolytic uremic syndrome with anti-factor H antibodies](#)
- [C3 glomerulonephritis](#)
- [Dense deposit disease](#)

### Related Gene Panels

- [Atypical Hemolytic Uremic Syndrome \(aHUS\) and Complement disorders \(17 genes\) - IPG](#)
- [Hereditary Hemolytic Anemias \(52 genes\) - ULB](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)
- [Nephropathy panel - UGent](#)
- [Panel Nephro-ULG-V1](#)
- [Primary immune deficiencies \(444 genes\) - KUL](#)
- [Primary immune deficiencies - UGent](#)

---

Source URL: <http://gentest.healthdata.be/index.php/index.php/analyte/2520>