

**DISEASE:**  
**Von Willebrand disease type 3**

<b>NAME:</b>	Von Willebrand disease type 3
<b>DESCRIPTION:</b>	A form of von Willebrand disease (VWD) characterized by a bleeding disorder associated with a total or near-total absence of Willebrand factor (VWF) in the plasma and cellular compartments, also leading to a profound deficiency of plasmatic factor VIII (FVIII). It is the most severe form of VWD.
<b>ORPHACODE:</b>	166096
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">MeSH</a> <a href="#">ICD-10</a> <a href="#">OMIM</a>
<b>ANALYTE(S):</b>	<a href="#">VWF</a>
<b>CREATED:</b>	13 May 2019 - 01:02
<b>CHANGED:</b>	22 Jun 2023 - 16:14

---

Source URL: <http://gentest.healthdata.be/disease/58>

## RELATED CONTENT

---

### Related Genetic Tests

- Trombosis - Hemostasis (gene panel)
- Von Willebrand disease

### Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Menselijke Erfelijheid - KUL

### Related Analytes

- von Willebrand factor

### Related Gene Panels

- Trombosis - Hemostasis (107 genes) - KUL

---

Source URL: <http://gentest.healthdata.be/disease/58>