# DISEASE:
# Von Willebrand disease type 1

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| **NAME:** | Von Willebrand disease type 1 |
| **DESCRIPTION:** | A form of von Willebrand disease (VWD) characterized by a bleeding disorder associated with a partial, quantitative plasmatic deficiency of an otherwise structurally and functionally normal von Willebrand factor (VWF). |
| **ORPHACODE:** | 166078 |
| **XREF(S):** | Orphanet<br>MeSH<br>OMIM<br>ICD-10 |
| **ANALYTE(S):** | VWF |
| **CREATED:** | 13 May 2019 - 01:02 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

# RELATED CONTENT

## Related Genetic Tests

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

## Related Laboratories

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

## Related Analytes

- von Willebrand factor

## Related Gene Panels

- Trombosis - Hemostasis (107 genes) - KUL