# DISEASE:
# Von Willebrand disease type 2N

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| **NAME:** | Von Willebrand disease type 2N |
| **DESCRIPTION:** | A subtype of type 2 von Willebrand disease characterized by a bleeding disorder associated with a marked decrease in the affinity of the Willebrand factor (VWF) for factor VIII (FVIII). Abnormal bleeding manifestations are less frequent in this VWD subtype than in other forms of the disease. The disease manifests mainly as soft tissue bleeding (haematoma, post-operative bleeding, etc.). |
| **ORPHACODE:** | 166093 |
| **XREF(S):** | Orphanet<br>ICD-10<br>OMIM |
| **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