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Von Willebrand disease type 3

INSERM

Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. [Von Willebrand disease type 3](#). ORPHA:166096

Type 3 von Willebrand disease (type 3 VWD) is the most severe form of VWD (see this term) characterized by a bleeding disorder associated with a total or near-total absence of Willebrand factor (von Willebrand factor; VWF) in the plasma and cellular compartments, also leading to a profound deficiency of plasmatic factor VIII (FVIII).