DISEASE:
Von Willebrand disease type 3

NAME:	Von Willebrand disease type 3
DESCRIPTION:	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.
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XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	VWF
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