



VWF Polyclonal Antibody

VWD is associated with a deficiency of VWF; type II by normal to decreased		
Reactivity Human;Rat;Mouse Applications IHC;IF Gene Name VWF F8VWF Protein Name von Willebrand factor (vWF) [Cleaved into: von Willebrand antigen 2 (von Willebrand antigen II)] Immunogen Synthesized peptide derived from part region of human protein AA range: 911-960 Specificity VWF Polyclonal Antibody detects endogenous levels of protein. Formulation Liquid in PBS containing 50% glycerol, and 0.02% sodium azide. Source Polyclonal, Rabbit,IgG Purification The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen. Dilution IHC-p 1:50-300. IF 1:50-200 Concentration 1 mg/ml Purity ≥90% Storage Stability -20°C/1 year Synonyms Observed Band 309kD Cell Pathway Secreted Secreted, extracellular space, extracellular matrix . Localized to storage granules. Tissue Specificity Plasma. Function disease: Defects in VWF are associated with various forms of von Willebrand disease (VWD) [MIM:193400, 277480]. VWD is characterized by frequent bleeding (gingival, minor skin quantitative lacerations, menorrhagia, etc.). Type VWD is associated with various forms of von Willebrand antigen 2 is required for example; type III by a virtual absence of VWF. There are subtypes to H) of type II VWD: for example; type III by a normal to decreased plasma level of VWF; type III by a virtual absence of VWF. There are subtypes to H) of type II VWD: for example; type III by a normal to decreased plasma level of VWF; type III by a virtual absence of VWF. There are subtypes to H) of type II VWD: for example; type III so parand to decreased arbeison of platelets to the sites of vascular injury by forming and polecular bridge adhesion of platelets to the sites of vascular injury by forming and elecular bridge adhesion of platelets to the sites of vascular injury by forming and elecular bridge adhesion of benestasis, it promote to the sites of the si	Catalog No	BYab-06901
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GPIb-IX-V. Also acts as a chaperone for coagu

Background

This gene encodes a glycoprotein involved in hemostasis. The encoded preproprotein is proteolytically processed following assembly into large multimeric complexes. These complexes function in the adhesion of platelets to sites of vascular injury and the transport of various proteins in the blood. Mutations in this gene result in von Willebrand disease, an inherited bleeding disorder. An unprocessed pseudogene has been found on chromosome 22. [provided by RefSeq, Oct 2015],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 30min).

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