

Anti-VWF Rabbit pAb



WL00949

For Research Use Only. Not For Use In Diagnostic Procedures

Product Information

Product name	Anti-VWF Rabbit pAb
Source	Rabbit
Species reactivity	Human
Tested applications	Western blot 1:1000-1:2000

**Suggested working dilutions are given as a guide only. It is recommended that the user titrates the product for use in their own experiment using appropriate negative and positive controls.*

Molecular Wt.	predicted size : 250kDa observed size : 150-200kDa
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Pack size	50/100/200/500/1000 μ l
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Storage	Store at -20°C. Avoid freeze/thaw cycles.
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Storage buffer	Supplied in 20 mM phosphate (pH 7.5), 150 mM NaCl, 100 μ g/ml BSA, 50% glycerol and less than 0.02% sodium azide
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General Information

Background	Von Willebrand disease is a congenital bleeding disorder caused by defects in the von Willebrand factor protein (VWF). VWF is thought to undergo a variety of posttranslational modifications that influence the affinity and availability for Factor VII, including cleavage of the propeptide and formation of N-terminal intersubunit disulfide bonds. VWF is a multimeric glycoprotein that is found in endothelial cells, plasma and platelets, and it is involved in the coagulation of blood at injury sites. VWF acts as a carrier protein for Factor VIII, a cofactor required for coagulation, and it promotes platelet adhesion and aggregation.
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Immunogen	Polyclonal antibody is produced by immunizing animals with recombinant protein of VWF.
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Purification	Polyclonal antibody was purified by Protein A affinity chromatography.
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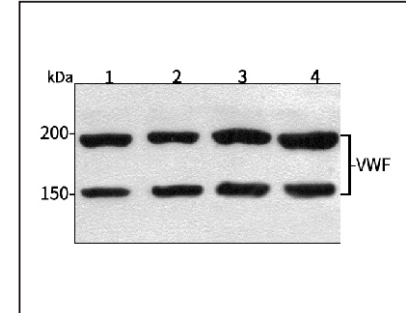
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Product Images



Western blot-Anti-VWF pAb

Lane 1: Human HepG2 cell lysate 24 μ gLane 2: Human Hela cell lysate 24 μ gLane 3: Human BGC-823 cell lysate 24 μ gLane 4: Human MGC-803 cell lysate 24 μ g

Separation gel: 6% polyacrylamide

Electrophoresis: 100V, 4°C, 3h

Transmembrane: 100V, 4°C, 1h

Blocking: 5% w/v nonfat dry milk, 1 \times TBST, at RT with gentle shaking

Primary antibody: 1:1000 in blocking buffer, 4°C, overnight

Visualization: ECL, 30s-2min