

# **VWF** Antibody

Purified Mouse Monoclonal Antibody (Mab) Catalog # AM8430b

#### **Product Information**

**Application** WB, IHC-P, FC, E

Primary Accession
Reactivity
Human
Host
Mouse
Clonality
Monoclonal
Isotype
IgG1

Clone Names 907CT12.1.9 Calculated MW 309265

#### **Additional Information**

**Gene ID** 7450

Other Names von Willebrand factor, vWF, von Willebrand antigen 2, von Willebrand antigen

II, VWF, F8VWF

Target/Specificity This antibody is generated from a mouse immunized with .VWF recombinant

protein.

**Dilution** WB~~1:1000 IHC-P~~1:100~500 FC~~1:100 E~~Use at an assay dependent

concentration.

**Format** Purified monoclonal antibody supplied in PBS with 0.09% (W/V) sodium azide.

This antibody is purified through a protein G column, followed by dialysis

against PBS.

**Storage** Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store

at -20°C in small aliquots to prevent freeze-thaw cycles.

**Precautions** VWF Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

#### **Protein Information**

Name VWF

Synonyms F8VWF

**Function** Important in the maintenance of hemostasis, it promotes adhesion of

platelets to the sites of vascular injury by forming a molecular bridge between

sub-endothelial collagen matrix and platelet- surface receptor complex

GPIb-IX-V. Also acts as a chaperone for coagulation factor VIII, delivering it to the site of injury, stabilizing its heterodimeric structure and protecting it from premature clearance from plasma.

**Cellular Location** Secreted. Secreted, extracellular space, extracellular matrix. Note=Localized

to storage granules

Tissue Location Plasma.

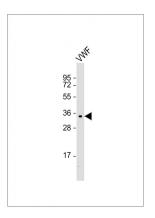
## **Background**

Important in the maintenance of hemostasis, it promotes adhesion of platelets to the sites of vascular injury by forming a molecular bridge between sub-endothelial collagen matrix and platelet-surface receptor complex GPIb-IX-V. Also acts as a chaperone for coagulation factor VIII, delivering it to the site of injury, stabilizing its heterodimeric structure and protecting it from premature clearance from plasma.

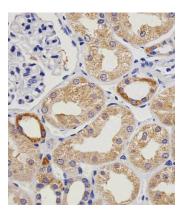
#### References

Bonthron D.,et al.Nucleic Acids Res. 14:7125-7128(1986). Mancuso D.J.,et al.J. Biol. Chem. 264:19514-19527(1989). Scherer S.E.,et al.Nature 440:346-351(2006). Verweij C.L.,et al.EMBO J. 5:1839-1847(1986). Verweij C.L.,et al.EMBO J. 5:3074-3074(1986).

### **Images**

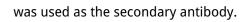


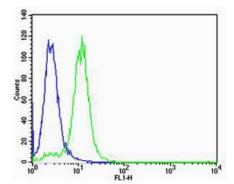
Anti-VWF at dilution + VWF whole cell lysate Lysates/proteins at 20 µg per lane. Secondary Goat Anti-mouse IgG, (H+L), Peroxidase conjugated at 1/10000 dilution. Predicted band size : 309 kDa Blocking/Dilution buffer: 5% NFDM/TBST.



Immunohistochemical analysis of paraffin-embedded H. kidney section using VWF(Cat#AM8430b). AM8430b was diluted at 1:25 dilution. A peroxidase-conjugated goat anti-mouse IgG at 1:400 dilution was used as the secondary antibody, followed by DAB staining.

Flow cytometric analysis of K562 cells using VWF(green, Cat#AM8430b) compared to an isotype control of mouse IgG1(blue). AP20600c was diluted at 1:100 dilution. An Alexa Fluor® 488 goat anti-mouse IgG at 1:400 dilution





Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.