



# Anti-VWF monoclonal antibody, clone 4F3 (CABT-B184)

This product is for research use only and is not intended for diagnostic use.

## PRODUCT INFORMATION

<b>Specificity</b>	Recognizes human VWF.
<b>Target</b>	VWF
<b>Immunogen</b>	von Willebrand factor, D3 domain
<b>Isotype</b>	IgG1, κ
<b>Source/Host</b>	Mouse
<b>Species Reactivity</b>	Human
<b>Clone</b>	4F3
<b>Purification</b>	Protein G affinity purified
<b>Conjugate</b>	Unconjugated
<b>Applications</b>	This clone is suitable for ELISA, western blot Recommended working dilution: Approximately 2 µg/ml. Each laboratory should determine an optimum working titer for use in its particular application. Other applications have not been tested but use in such assays should not necessarily be excluded
<b>Molecular Weight</b>	>300 kDa
<b>Concentration</b>	1 mg/mL
<b>Size</b>	100ug, 500ug
<b>Buffer</b>	0.1M Sodium Phosphate, pH 7.4, 0.15M NaCl, 0.05% (w/v) Sodium Azide

<b>Preservative</b>	0.05% Sodium Azide
<b>Storage</b>	-20°C
<b>Ship</b>	Cold packs

## BACKGROUND

<b>Introduction</b>	Von Willebrand factor (vWF) is a blood glycoprotein involved in 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. vWF 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. It is deficient or defective in von Willebrand disease and is involved in a large number of other diseases, including thrombotic thrombocytopenic purpura, Heydes syndrome, and hemolytic-uremic syndrome.
---------------------	--

<b>Keywords</b>	VWF;von Willebrand factor;VWD;F8VWF;coagulation factor VIII VWF
-----------------	---

## GENE INFORMATION

<b>Entrez Gene ID</b>	<a href="#">7450</a>
<b>UniProt ID</b>	<a href="#">P04275</a>