

Product datasheet

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ARG42214 anti-vWF antibody [3E2D10]

Package: 50 μg Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [3E2D10] recognizes vWF

Tested Reactivity Hu

Tested Application FACS, ICC/IF, IHC-P, IP, WB

Host Mouse

Clonality Monoclonal
Clone 3E2D10

Isotype IgG1, kappa

Target Name vWF

Species Human

Immunogen Synthetic peptide corresponding to aa. 845-949 of Human vWF.

Conjugation Un-conjugated

Alternate Names VWD; von Willebrand factor; vWF; von Willebrand antigen II; F8VWF

Application Instructions

Application table	Application	Dilution
	FACS	0.5 - 2 μg/10^6 cells
	ICC/IF	0.5 - 2 μg/ml
	IHC-P	0.5 - 2 μg/ml
	IP	1 - 4 μg/mg protein lysate
	WB	0.5 - 2 μg/ml
Application Note	Antigen Retrieval: Boil tissue section in 10 mM Tris with 1 mM EDTA (pH 9.0) for 10-20 min, followed by cooling at RT for 20 min. * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid	
Purification	Purification with Protein G.	
Buffer	PBS, 0.05% Sodium azide and 0.1 mg/ml BSA.	
Preservative	0.05% Sodium azide	
Stabilizer	0.1 mg/ml BSA	

Concentration 0.2 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

before use.

Note For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol VWF

Gene Full Name von Willebrand factor

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]

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. [UniProt]

Calculated Mw 309 kDa

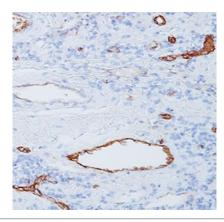
PTM All cysteine residues are involved in intrachain or interchain disulfide bonds.

N- and O-glycosylated. [UniProt]

Cellular Localization Secreted. Secreted, extracellular space, extracellular matrix. Note=Localized to storage granules.

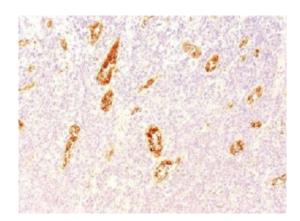
[UniProt]

Images



ARG42214 anti-vWF antibody [3E2D10] IHC-P image

Immunohistochemistry: Formalin-fixed and paraffin-embedded Human tonsil tissue. Antigen Retrieval: Boil tissue section in 10 mM Tris with 1 mM EDTA (pH 9.0) for 10-20 min, followed by cooling at RT for 20 min. The tissue section was stained with ARG42214 anti-vWF antibody [3E2D10].



ARG42214 anti-vWF antibody [3E2D10] IHC-P image

Immunohistochemistry: Formalin-fixed and paraffin-embedded Human pancreas tissue. Antigen Retrieval: Boil tissue section in 10 mM Tris with 1 mM EDTA (pH 9.0) for 10-20 min, followed by cooling at RT for 20 min. The tissue section was stained with ARG42214 antivWF antibody [3E2D10].