

Product datasheet

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ARG21352 anti-CD42b antibody [MM2/174]

Package: 100 μg Store at: -20°C

Summary

Product Description Mouse Monoclonal antibody [MM2/174] recognizes CD42b

Tested Reactivity Hu

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

Specificity Human CD42b.

Host Mouse

Clonality Monoclonal

Clone MM2/174

Isotype IgG1, kappa

Target Name CD42b
Species Human

Immunogen Human plasma membrane

Conjugation Un-conjugated

Alternate Names CD antigen CD42b; Antigen CD42b-alpha; DBPLT3; VWDP; CD42B; GP-Ib alpha; Glycoprotein Ibalpha;

BDPLT1; BSS; CD42b-alpha; GPIbA; GPIb-alpha; Platelet glycoprotein Ib alpha chain; BDPLT3; GP1B

Application Instructions

Application table	Application	Dilution
	FACS	< 1 µg/10^6 cells
	ICC/IF	Assay-dependent
	IHC-Fr	Assay-dependent
	IHC-P	Assay-dependent
	IP	Assay-dependent
	WB	Assay-dependent
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

Properties

Form	Liquid
Buffer	BBS (pH 8.2)
Concentration	0.1 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. 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

Database links <u>GeneID: 2811 Human</u>

Swiss-port # P07359 Human

Gene Symbol GP1BA

Gene Full Name glycoprotein Ib (platelet), alpha polypeptide

Background Glycoprotein Ib (GP Ib) is a platelet surface membrane glycoprotein composed of a heterodimer, an

alpha chain and a beta chain, that is linked by disulfide bonds. The Gp Ib functions as a receptor for von Willebrand factor (VWF). The complete receptor complex includes noncovalent association of the alpha and beta subunits with platelet glycoprotein IX and platelet glycoprotein V. The binding of the GP Ib-IX-V complex to VWF facilitates initial platelet adhesion to vascular subendothelium after vascular injury, and also initiates signaling events within the platelet that lead to enhanced platelet activation, thrombosis, and hemostasis. This gene encodes the alpha subunit. Mutations in this gene result in Bernard-Soulier syndromes and platelet-type von Willebrand disease. The coding region of this gene is known to contain a polymophic variable number tandem repeat (VNTR) domain that is associated with susceptibility to nonarteritic anterior ischemic optic neuropathy. [provided by RefSeq, Oct 2013]

Function GP-Ib, a surface membrane protein of platelets, participates in the formation of platelet plugs by

binding to the A1 domain of vWF, which is already bound to the subendothelium. [UniProt]

Calculated Mw 72 kDa

PTM Glycocalicin, which is approximately coextensive with the extracellular part of the molecule, is cleaved

off by calpain during platelet lysis.