For Research Use Only

CD36 Monoclonal antibody

Catalog Number:66395-1-lg 13 Publications

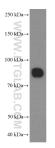


Basic Information	Catalog Number:	GenBank Accession Number:		Purification Method:	
	66395-1-lg	BC008406		Protein A purification	
	Size:	GeneID (NCBI):		CloneNo.: 1A8C5	
	Nanodron and 1000 ug/ml by Bradford				
	method using BSA as the standard; CD36 molecule (thrombospondin		Recommended Dilutions: WB 1:1000-1:6000		
	Source:	receptor)	cate (thrombosponam	IHC 1:500-1:2000	
	Mouse	Calculated MW:		IF 1:200-1:800	
	Isotype:	472 aa, 53 k			
	lgG1	Observed M	1W:		
	Immunogen Catalog Number: AG13541				
Applications	Tested Applications:		Positive Con	trols:	
	FC, IF, IHC, WB, ELISA Cited Applications:		WB : human heart tissue, human milk, human spleen tissue, human placenta tissue		
	IF, IHC, WB		IHC : human	spleen tissue, human heart tissue	
	Species Specificity: IF : human		IF : human li	liver cancer tissue,	
	Cited Species: human				
	Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0				
	CD36, also named as GP3B and GP4, is an 88-kDa membrane glycoprotein present on platelets, monocytes, erythroid precursors, endothelial cells, and several tumor cell lines. CD36 binds to collagen, thrombospondin, anionic phospholipids, long chain fatty acids and may function in the transport and/or as a regulator of fatty acid transport and oxidized LDL. CD36 may function as a cell adhesion molecule. It mediates cytoadherence of Plasmodium falciparum parasitized erythrocytes. Mutation of CD36 will cause platelet glycoprotein IV deficiency which known as CD36 deficiency. Genetic variations in CD36 are associated with susceptibility to coronary heart disease type 7 (CHDS7).				
Background Information	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia	ction as a cell adh tes. Mutation of C	nesion molecule. It medi D36 will cause platelet	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which known	
	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia (CHDS7).	ction as a cell adh tes. Mutation of C	nesion molecule. It medi D36 will cause platelet	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which known	
	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia (CHDS7).	ction as a cell adh tes. Mutation of C ations in CD36 ar	nesion molecule. It medi D36 will cause platelet e associated with suscep	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which known tibility to coronary heart disease type 7	
	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia (CHDS7). Author Yan Cao	ction as a cell adh tes. Mutation of C ations in CD36 ar Pubmed ID	nesion molecule. It medi D36 will cause platelet e associated with suscep Journal	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which known tibility to coronary heart disease type 7 Application WB	
	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia (CHDS7). Author Yan Cao Simon Riis	ction as a cell adh tes. Mutation of C ations in CD36 ar Pubmed ID 34599913	esion molecule. It medi D36 will cause platelet e associated with suscep Journal Eur J Pharmacol	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which known tibility to coronary heart disease type 7 Application WB ts IHC	
Background Information Notable Publications	and oxidized LDL CD36 may func falciparum parasitized erythrocy as CD36 deficiency. Genetic varia (CHDS7). Author Yan Cao Simon Riis	ttion as a cell adh tes. Mutation of C ations in CD36 ar Pubmed ID 34599913 31430404 35656026 r after shipment. d 50% glycerol p	nesion molecule. It medi D36 will cause platelet e associated with suscep Journal Eur J Pharmacol Scand J Med Sci Spo Oxid Med Cell Longe	d/or as a regulator of fatty acid transport ates cytoadherence of Plasmodium glycoprotein IV deficiency which knowr tibility to coronary heart disease type 7 Application WB ts IHC	

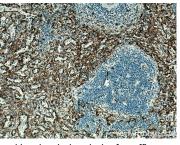
For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free
in USA), or 1(312) 455-8498 (outside USA)E: proteintech@ptglab.comW: ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

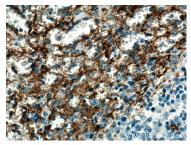
Selected Validation Data



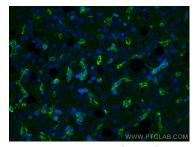
human heart tissue were subjected to SDS PAGE followed by western blot with 66395-1-1g (CD36 Antibody) at dilution of 1:3000 incubated at room temperature for 1.5 hours.



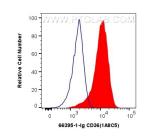
Immunohistochemical analysis of paraffinembedded human spleen tissue slide using 66395-1-Ig (CD36 antibody) at dilution of 1:1000 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunohistochemical analysis of paraffinembedded human spleen tissue slide using 66395-1-Ig (CD36 antibody) at dilution of 1:1000 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



Immunofluorescent analysis of (4% PFA) fixed human liver cancer tissue using CD36 antibody (66395-1-1g, Clone: 1A8C5) at dilution of 1:400 and CoraLite®488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L).



1X10^6 THP-1 cells were surface stained with 0.4 ug Anti-Human CD36 (66395-1-1g, Clone:1A8C5) and Coralite@488-Conjugated AffiniPure Goat Anti-Mouse IgG(H+L) at dilution 1:1000 (red), or 0.4 ug Mouse IgG1 Isotype Control (MOPC-21) (65124-1-1g, Clone: MOPC-21) (blue). Cells were not fixed.