

Product Data Sheet

Anti-CD61 Antibody

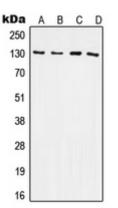
Catalog #	Source	Reactivity		Applications		
CPA1617	Rabbit	H, M, R, B, D, Mk, P, I	Rb, S	WB, IH, IF/IC, IP		
Description	R	abbit polyclonal antibody to	CD61			
Immunogen	К	LH-conjugated synthetic per	otide encomp	assing a sequence within the C-term		
	re	region of human CD61. The exact sequence is proprietary.				
Purification	Т	he antibody was purified by	immunogen	affinity chromatography.		
Specificity	R	Recognizes endogenous levels of CD61 protein.				
Clonality	P	Polyclonal				
Conjugation						
Form	Li	iquid in 0.42% Potassium ph	osphate, 0.87	% Sodium chloride, pH 7.3, 30% glycerol,		
	a	nd 0.01% sodium azide.				
Dilution	W	VB (1/500 - 1/1000), IH (1/100	- 1/200), IF/I	C (1/100 - 1/500), IP (1/10 - 1/100)		
Gene Symbol	TI	TGB3				
Alternative N	ames G	SP3A; Integrin beta-3; Platele	t membrane	glycoprotein IIIa; GPIIIa; CD61		
Entrez Gene	3	690 (Human); 16416 (Mouse	e)			
SwissProt	P	205106 (Human); O54890 (M	ouse)			
Storage/Stabi	i lity Sl	hipped at 4°C. Upon delivery	aliquot and	store at -20°C for one year. Avoid		
	fr	reeze/thaw cycles.				

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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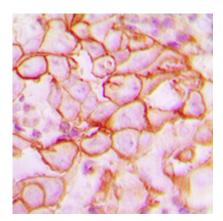
Cohesion



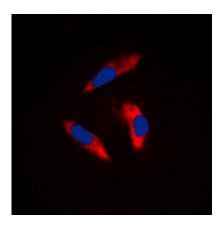
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Western blot analysis of CD61 expression in HeLa (A), Raw264.7 (B), NIH3T3 (C), H9C2 (D) whole cell lysates. (Predicted band size: 87 kD; Observed band size: 130 kD)



Immunohistochemical analysis of CD61 staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of CD61 staining in Raw264.7 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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