

## **Product Data Sheet**

## Anti-CD61 (Phospho-Y773) Antibody

Catalog #	Source	Reactivity	Applications	
CPA4041	Rabbit	H, M, R, B, D, Mk, P, Rb, S	WB, IH	
Description	Ra	bbit polyclonal antibody to CD6:	1 (Phospho-Y773)	
Immunogen	KL	H-conjugated synthetic phospho	peptide corresponding to residue	s surrounding
	Y7	73 of human CD61 protein. The	exact sequence is proprietary.	
Purification	Th	e antibody was purified by imm	unogen affinity chromatography.	
Specificity	Re	cognizes endogenous levels of C	D61 protein only when phosphor	ylated at Y773.
Clonality	Ро	lyclonal		
Conjugation				
Form	Liq	uid in 0.42% Potassium phospha	ate, 0.87% Sodium chloride, pH 7.	3, 30% glycerol,
	an	d 0.01% sodium azide.		
Dilution	WI	B (1/500 - 1/1000), IH (1/100 - 1/20	00)	
Gene Symbol	ITC	GB3		
Alternative N	ames GP	3A; Integrin beta-3; Platelet me	mbrane glycoprotein IIIa; GPIIIa; C	CD61
Entrez Gene	36	90 (Human); 16416 (Mouse)		
SwissProt	PO	5106 (Human); O54890 (Mouse)	)	
Storage/Stabi	lity Sh	ipped at 4°C. Upon delivery aliqu	uot and store at -20°C for one yea	r. Avoid
	fre	eze/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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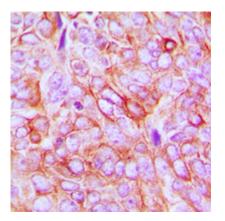
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R

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of CD61 (Phospho-Y773) expression in HUVEC (A), Raw264.7 (B) whole cell lysates. (Predicted band size: 87 kD; Observed band size: 130 kD)



Immunohistochemical analysis of CD61 (Phospho-Y773) 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.

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