

Product Data Sheet

Anti-CD213a1 (Phospho-Y405) Antibody

Catalog #	Source	Reactivity	Α	pplications		
CPA4686	Rabbit	H, M, Mk	V	VB, IF/IC		
Description	Rabb	Rabbit polyclonal antibody to CD213a1 (Phospho-Y405)				
Immunogen	KLH-	onjugated synthetic phosphopeptide corresponding to residues surrounding				
	Y405	of human CD213a1 pr	otein. The exact s	equence is proprietary.		
Purification	The a	antibody was purified b	y immunogen aff	inity chromatography.		
Specificity	ecificity Recognizes endogenous levels of CD213a1 protein only when phosphorylated a					
	Y405					
Clonality	Polyc	clonal				
Conjugation						
Form	Liqui	d in 0.42% Potassium p	hosphate, 0.87%	Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.				
Dilution	WB (1/500 - 1/1000), IF/IC (1	/100 - 1/500)			
Gene Symbol	IL13F	RA1				
Alternative Na	ames IL13F	R; IL13RA; Interleukin-1	3 receptor subun	it alpha-1; IL-13 receptor subunit		
	alpha	a-1; IL-13R subunit alpł	a-1; IL-13R-alpha	-1; IL-13RA1; Cancer/testis antigen 19;		
	CT19	; CD213a1				
Entrez Gene	3597	(Human); 16164 (Mou	se)			
SwissProt	P785	52 (Human); O09030 (Mouse)			
Storage/Stabi	lity Ship	oed at 4°C. Upon delive	ry aliquot and sto	ore at -20°C for one year. Avoid		
	freez	e/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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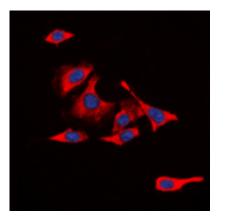
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For research purposes only, not for human use

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Western blot analysis of CD213a1 (Phospho-Y405) expression in HCT116 (A), HuvEc (B) whole cell lysates. (Predicted band size: 48 kD; Observed band size: 49 kD)



Immunofluorescent analysis of CD213a1 (Phospho-Y405) staining in HCT116 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 hidified 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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