

Product Data Sheet

Anti-CD247 (Phospho-Y142) Antibody

Catalog #	Source	Reactivity	Applications			
CPA6505	Rabbit	Н, М	WB, IH			
Description	o CD247 (Phospho-Y142)					
Immunogen	KI	LH-conjugated synthetic pl	nosphopeptide corresponding to residues surrounding			
	Y	142 of human CD247 prote	in. The exact sequence is proprietary.			
Purification	Tł	he antibody was purified b	ibody was purified by immunogen affinity chromatography.			
Specificity	Recognizes endogenous levels of CD247 protein only when phosphorylated at Y14					
Clonality	Ро	Polyclonal				
Conjugation						
Form	Li	quid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.				
Dilution	W	WB (1/500 - 1/1000), IH (1/50 - 1/200)				
Gene Symbol	CI	D247				
Alternative N	ames Cl	D3Z; T3Z; TCRZ; T-cell surfa	ce glycoprotein CD3 zeta chain; T-cell receptor T3 zeta			
	cł	nain; CD247				
Entrez Gene	91	19 (Human); 12503 (Mous	e)			
SwissProt	P2	20963 (Human); P24161 (I	/louse)			
Storage/Stabi	lity Sł	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fr	eeze/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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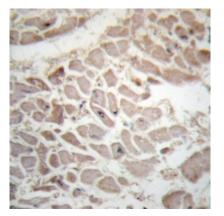
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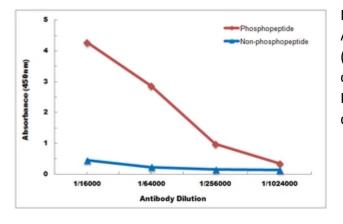
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For research purposes only, not for human use

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Western blot analysis of CD247 (Phospho-Y142) expression in HEK293T (A), LOVO (B), HCT116 (C), mouse liver (D) whole cell lysates. (Predicted band size: 18 kD; Observed band size: 16 kD)





Immunohistochemical analysis of CD247 (Phospho-Y142) staining in human heart 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.

Direct ELISA antibody dose-response curve using Anti-CD247 (Phospho-Y142) Antibody. Antigen (Phosphopeptide and non-phosphopeptide) concentration is 5 ug/ml. Goat Anti-Rabbit IgG (H&L) -HRP was used as the secondary antibody, and signal was developed by TMB substrate.

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