

Product Data Sheet

Anti-SYK (Phospho-Y348) Antibody

Catalog #	Source	Reactivity	Applications		
CPA2128	Rabbit	H, M, R, C, D, P, Rb	WB, IH		
Description	Rab	Rabbit polyclonal antibody to SYK (Phospho-Y348)			
Immunogen	KLH	-conjugated synthetic phos	phopeptide corresponding to residues surrounding		
	Y34	3 of human SYK protein. Th	e exact sequence is proprietary.		
Purification	The	antibody was purified by ir	nmunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous levels	of SYK protein only when phosphorylated at Y348.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	id in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/50 - 1	/100)		
Gene Symbol	SYK				
Alternative Na	ames Tyrc	sine-protein kinase SYK; Sp	oleen tyrosine kinase; p72-Syk		
Entrez Gene	685	0 (Human); 20963 (Mouse)	; 25155 (Rat)		
SwissProt	P43	405 (Human); P48025 (Mo	use); Q64725 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery a	aliquot and store at -20°C for one year. Avoid		
	free	ze/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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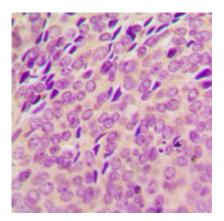
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For research purposes only, not for human use

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Western blot analysis of SYK (Phospho-Y348) expression in HEK293T (A) whole cell lysates. (Predicted band size: 72 kD; Observed band size: 72 kD)



Immunohistochemical analysis of SYK (Phospho-Y348) 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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