

Product Data Sheet

Anti-PKC alpha (Phospho-T638) Antibody

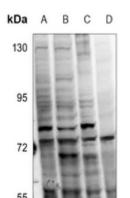
Catalog #	Source	Reactivity	Applications		
CPA4782	Rabbit	H, M, R, B	WB, IH		
Description		Rabbit polyclonal antibody to PKC alpha (Phospho-T638)			
Immunogen		KLH-conjugated synthetic pl	osphopeptide corresponding to residues surrounding		
		T638 of human PKC alpha p	otein. The exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of PKC alpha protein only when phosphorylated at			
		Т638.			
Clonality		Polyclonal			
Conjugation					
Form		Liquid 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), IH (1/10	0 - 1/200)		
Gene Symbol		PRKCA			
Alternative Names		PKCA; PRKACA; Protein kinase C alpha type; PKC-A; PKC-alpha			
Entrez Gene		5578 (Human); 18750 (Mou	se)		
SwissProt		P17252 (Human); P20444 (N	1ouse); P05696 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
		freeze/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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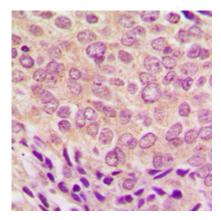




For research purposes only, not for human use

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Western blot analysis of PKC alpha (Phospho-T638) expression in A549 (A), EC9706 (B), C6 (C), BV2 (D) whole cell lysates. (Predicted band size: 76 kD; Observed band size: 80; 76 kD)



Immunohistochemical analysis of PKC alpha (Phospho-T638) 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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