

Product Data Sheet

Anti-PKC theta (Phospho-S676) Antibody

Catalog #	Source	Reactivity	Applications	
CPA1939	Rabbit	H, M, R, B, D, Mk, P	WB, IH, IF/IC	
Description	Rab	Rabbit polyclonal antibody to PKC theta (Phospho-S676)		
Immunogen	KLH	I-conjugated synthetic phosphopeptic	de corresponding to residues surrounding	
	S67	6 of human PKC theta protein. The ex	act sequence is proprietary.	
Purification	The	antibody was purified by immunoge	n affinity chromatography.	
Specificity	Rec	Recognizes endogenous levels of PKC theta protein only when phosphorylated at		
	S67	6.		
Clonality	Poly	yclonal		
Conjugation				
Form	Liqu	uid in 0.42% Potassium phosphate, 0.	87% Sodium chloride, pH 7.3, 30% glycerol,	
	and	0.01% sodium azide.		
Dilution	WB	(1/500 - 1/1000), IH (1/100 - 1/200), IF	/IC (1/100 - 1/500)	
Gene Symbol	PRK	CQ		
Alternative Na	ames PRK	CT; Protein kinase C theta type; nPKC	2-theta	
Entrez Gene	558	8 (Human); 18761 (Mouse)		
SwissProt	Q04	1759 (Human); Q02111 (Mouse); Q9V	VTQ0 (Rat)	
Storage/Stabi	lity Ship	oped at 4°C. Upon delivery aliquot an	d store at -20°C for one year. Avoid	
	free	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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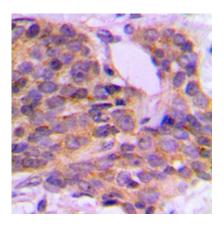
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Western blot analysis of PKC theta (Phospho-S676) expression in HeLa UV-treated (A), SP2/0 UV-treated (B), PC12 UV-treated (C) whole cell lysates. (Predicted band size: 81 kD; Observed band size: 80 kD)



Immunohistochemical analysis of PKC theta (Phospho-S676) 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.

Immunofluorescent analysis of PKC theta (Phospho-S676) staining in HeLa 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 humidified 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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