

# **Product Data Sheet**

### Anti-PKC gamma Antibody

Catalog #	Source	Reactivity	Applications		
CPA1936	Rabbit	H, M, R, B, Mk, Rb	WB, IH		
Description	Rab	Rabbit polyclonal antibody to PKC gamma			
Immunogen	KLH	-conjugated synthetic pep	tide encompassing a sequence within the center		
	regi	on of human PKC gamma.	The exact sequence is proprietary.		
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous levels	s of PKC gamma protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	iid in 0.42% Potassium pho	osphate, 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	PRK	CG			
Alternative Na	ames PKC	G; Protein kinase C gamma	a type; PKC-gamma		
Entrez Gene	558	2 (Human); 18752 (Mouse	e); 24681 (Rat)		
SwissProt	P05	129 (Human); P63318 (Mc	ouse); P63319 (Rat)		
Storage/Stabi	<b>lity</b> Ship	ped at 4°C. Upon delivery	v 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

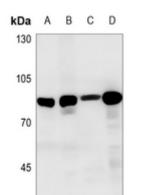
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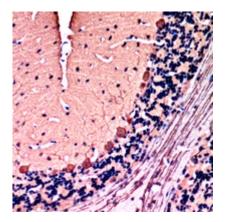


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Western blot analysis of PKC gamma expression in A549 (A), H1792 (B), mouse kidney (C), rat brain (D) whole cell lysates. (Predicted band size: 78 kD; Observed band size: 83 kD)



Immunohistochemical analysis of PKC gamma staining in rat brain 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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