

Product Data Sheet

Anti-PI3K p85 alpha Antibody

Catalog #	Source	Reactivity	Applications		
CPA3286	Rabbit	H, M, R, B	WB, IH		
Description		Rabbit polyclonal antibody to PI3K p85 alpha			
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
		region of human PI3K p85 al	bha. The exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	s of PI3K p85 alpha protein.		
Clonality		Polyclonal			
Form		Liquid 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)		
Gene Symbol		PIK3R1			
Alternative Names		GRB1; Phosphatidylinositol 3-kinase regulatory subunit alpha; PI3-kinase regulatory			
		subunit alpha; PI3K regulator	y subunit alpha; PtdIns-3-kinase regulatory subunit		
		alpha; Phosphatidylinositol 3	-kinase 85 kDa regulatory subunit alpha; PI3-kinase		
		subunit p			
Entrez Gene		5295 (Human); 18708 (Mous	e); 25513 (Rat)		
SwissProt		P27986 (Human); P26450 (M	ouse); Q63787 (Rat)		
Storage/Stabi	ility	Shipped at 4°C. Upon deliver	y 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

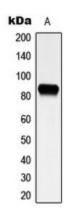
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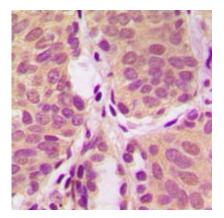


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Western blot analysis of PI3K p85 alpha expression in HeLa (A) whole cell lysates.



Immunohistochemical analysis of PI3K p85 alpha staining in human prostate 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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