

Product Data Sheet

Anti-DNA-PKcs (Phospho-S2612) Antibody

Catalog #	Source	e Reactivity	Applications		
CPA6230	Rabbit	н, М	WB, IH		
Description		Rabbit polyclonal antibody to DNA-PKcs (Phospho-S2612)			
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
		S2612 of human DNA-PKcs protein. The exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of DNA-PKcs protein only when phosphorylated at			
		S2612.			
Clonality		Polyclonal			
Conjugation					
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	/50 - 1/200)		
Gene Symbol		PRKDC			
Alternative Names		HYRC; HYRC1; DNA-dependent protein kinase catalytic subunit; DNA-PK catalytic			
		subunit; DNA-PKcs; DNPk	(1; p460		
Entrez Gene	5591 (Human); 19090 (Mouse)				
SwissProt		P78527 (Human); P97313	B (Mouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery 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

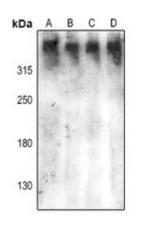
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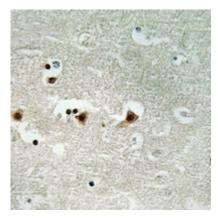
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Western blot analysis of DNA-PKcs (Phospho-S2612) expression in mouse heart (A), NIH3T3L1 (B), HEK293T (C), SGC7901 (D) whole cell lysates. (Predicted band size: 469 kD; Observed band size: 470 kD)



Immunohistochemical analysis of DNA-PKcs (Phospho-S2612) staining in human 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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