

Product Data Sheet

Anti-PKR Antibody

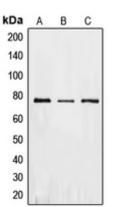
Catalog #	Source	e Reactivity	Applications		
CPA1952	Rabbit	-	WB, IH, IF/IC		
Description		Rabbit polyclonal antibody t			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
			xact sequence is proprietary.		
Purification		C	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	els of PKR protein.		
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 (1/10	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol		EIF2AK2			
Alternative Names		PKR; PRKR; Interferon-induced, double-stranded RNA-activated protein kinase;			
		Eukaryotic translation initiat	ion factor 2-alpha kinase 2; eIF-2A protein kinase 2;		
		Interferon-inducible RNA-de	pendent protein kinase; P1/eIF-2A protein kinase;		
		Protein kinase RNA-activate	d; PKR; Tyrosine-protein kinase EIF2AK2; p68 kinase		
Entrez Gene		5610 (Human); 19106 (Mou	se)		
SwissProt		P19525 (Human); Q03963 (I	Mouse)		
Storage/Stabi	lity	Shipped at 4 $^\circ$ C. Upon deliv	ery aliquot and store at -20 $^\circ$ 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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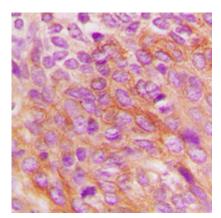
Coherion



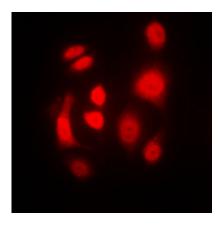
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Western blot analysis of PKR expression in HeLa (A), MCF7 (B), Caco2 (C) whole cell lysates. (Predicted band size: 62 kD; Observed band size: 74 kD)



Immunohistochemical analysis of PKR 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 PKR staining in MCF7 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 $^{\circ}$ 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.

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