

Product Data Sheet

Anti-PKR Antibody

Catalog #	Source	Reactivity	Applications		
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CPA4395	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to	PKR		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
		region of human PKR. The ex	act sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous level	s 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 (1/100	- 1/200)		
Gene Symbol		EIF2AK2			
Alternative Na	ames	PKR; PRKR; Interferon-induce	d, double-stranded RNA-activated protein kinase;		
		Eukaryotic translation initiati	on factor 2-alpha kinase 2; eIF-2A protein kinase 2;		
		Interferon-inducible RNA-dep	endent protein kinase; P1/eIF-2A protein kinase;		
		Protein kinase RNA-activated	; PKR; Tyrosine-protein kinase EIF2AK2; p68 kinase		
Entrez Gene		5610 (Human); 19106 (Mous	e); 54287 (Rat)		
SwissProt		P19525 (Human); Q03963 (N	ouse); Q63184 (Rat)		
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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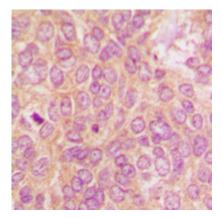
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kDa A B C D 180 130 95 72 55 Western blot analysis of PKR expression in HCC827 (A), SGC7901 (B), A375 (C), MCF7 (D) whole cell lysates. (Predicted band size: 62 kD; Observed band size: 74 kD)



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.

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