

# **Product Data Sheet**

# **Anti-PKR Antibody**

Catalog #	Source	e Reactivity	Applications	
CPA4395	Rabbit	t Н, М	WB, IH	
Description		Rabbit polyclonal antibody t	o PKR	
Immunogen		KLH-conjugated synthetic pe	ptide encompassing a sequence within the C-term	
		region of human PKR. The e	kact sequence is proprietary.	
Purification		The antibody was purified b	y immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	els of PKR protein.	
Clonality		Polyclonal		
Conjugation				
Form		Liquid in 0.42% Potassium p	hosphate, 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)	
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	Aouse)	
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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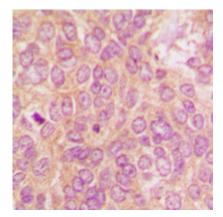
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**kDa** A B C D 180 130 95 72 55 Western blot analysis of PKR expression in HCC827 (A), SGC7901 (B), A375 (C), MCF7 (D) whole cell lysates. (Predicted band size: 62 kD; Observed band size: 74 kD)



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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