

## **Product Data Sheet**

## **Anti-CHK2** Antibody

Catalog #	Source	Reactivity	Applications		
CPA5711	Rabbit	H, M, R, B, P	WB, IH		
Description		Rabbit polyclonal antibody to	СНК2		
Immunogen		KLH-conjugated synthetic pep	tide encompassing a sequence within the center		
		region of human CHK2. The ex	act sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels	s of CHK2 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 (1/50 -	1/100)		
Gene Symbol		CHEK2			
Alternative Na	ames	CDS1; CHK2; RAD53; Serine/tl	nreonine-protein kinase Chk2; CHK2 checkpoint		
		homolog; Cds1 homolog; Huc	ds1; hCds1; Checkpoint kinase 2		
Entrez Gene		11200 (Human); 50883 (Mous	se)		
SwissProt		O96017 (Human); Q9Z265 (M	ouse)		
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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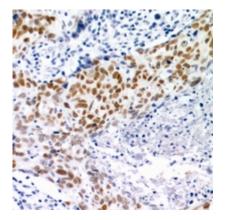
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For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of CHK2 expression in COS7 UV treated 30' (A) whole cell lysates. (Predicted band size: 60 kD; Observed band size: 62 kD)



Immunohistochemical analysis of CHK2 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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