

## **Product Data Sheet**

## Anti-CHK1 (Phospho-S317) Antibody

Catalog #	Source	Reactivity	Applications		
CPA1224	Rabbit	H, M, R	WB, IH		
Description	Rabbit polyclonal antibody to CHK1 (Phospho-S317)				
Immunogen	К	KLH-conjugated synthetic pl	nosphopeptide corresponding to residues surrounding		
	S	317 of human CHK1 protei	n. The exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	els of CHK1 protein only when phosphorylated at S317.		
Clonality Polyclonal					
Conjugation					
Form	L	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/10	00 - 1/200)		
Gene Symbol	C	CHEK1			
Alternative Na	ames C	CHK1; Serine/threonine-pro	tein kinase Chk1; CHK1 checkpoint homolog; Cell cycle		
	С	heckpoint kinase; Checkpo	int kinase-1		
Entrez Gene	1	.111 (Human); 12649 (Mou	se); 140583 (Rat)		
SwissProt	C	014757 (Human); 035280 (	Mouse); Q91ZN7 (Rat)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fi	reeze/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

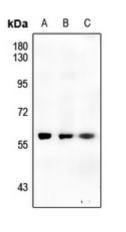
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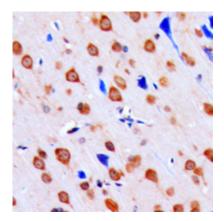
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Western blot analysis of CHK1 (Phospho-S317) expression in HEK293T-UV 6h (A), HEK293T-UV 4h (B), HEK293T (C) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 56 kD)



Immunohistochemical analysis of CHK1 (Phospho-S317) 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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