

## **Product Data Sheet**

### Anti-CHK1 (Phospho-S345) Antibody

Catalog #	Source	Reactivity	Applications		
CPA4600	Rabbit	H, M, R	WB, IH, IF/IC		
Description	F	Rabbit polyclonal antibody to CHK1 (Phospho-S345)			
Immunogen	K	(LH-conjugated synthetic pr	osphopeptide corresponding to residues surrounding		
	S	345 of human CHK1 protein	n. The exact sequence is proprietary.		
Purification	Т	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of CHK1 protein only when phosphorylated at \$345.		
Clonality	P	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	C	CHEK1			
Alternative N	ames C	CHK1; Serine/threonine-pro	tein kinase Chk1; CHK1 checkpoint homolog; Cell cycle		
	C	checkpoint kinase; Checkpo	nt kinase-1		
Entrez Gene	1	l111 (Human); 12649 (Mou	se); 140583 (Rat)		
SwissProt	C	014757 (Human); 035280 (I	Mouse); Q91ZN7 (Rat)		
Storage/Stabi	ility S	Shipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	f	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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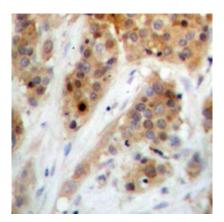
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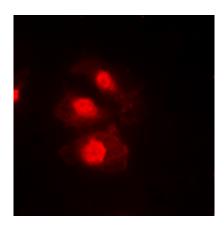
kDa A B C D E

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Western blot analysis of CHK1 (Phospho-S345) expression in Hela (A), HGC27 (B), mouse testis (C), mouse kidney (D), rat testis (E) whole cell lysates. (Predicted band size: 54 kD; Observed band size: 56 kD)





Immunohistochemical analysis of CHK1 (Phospho-S345) 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 CHK1 (Phospho-S345) staining in HeLa 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 °C in a hidified 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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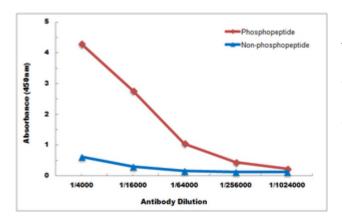
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Direct ELISA antibody dose-response curve using Anti-CHK1 (Phospho-S345) Antibody. Antigen (Phosphopeptide and non-phosphopeptide) concentration is 5 ug/ml. Goat Anti-Rabbit IgG (H&L) -HRP was used as the secondary antibody, and signal was developed by TMB substrate.

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