

Product Data Sheet

Anti-CDC2 (Phospho-T161) Antibody

Catalog #	Source	Reactivity	Applications			
CPA1189	Rabbit	H, M, R, B, C, Mk, Z	WB, IH			
Description	Rab	Rabbit polyclonal antibody to CDC2 (Phospho-T161)				
Immunogen	KLH	-conjugated synthetic phosphc	opeptide corresponding to residues surrounding			
	T16	1 of human CDC2 protein. The	exact sequence is proprietary.			
Purification	The	antibody was purified by imm	unogen affinity chromatography.			
Specificity	Rec	Recognizes endogenous levels of CDC2 protein only when phosphorylated at T161.				
Clonality	Poly	Polyclonal				
Conjugation						
Form	Liqu	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 - 1/10	00)			
Gene Symbol	CD	<1; CDK2				
Alternative Na	ames CDC	2; CDC28A; CDKN1; P34CDC2;	Cyclin-dependent kinase 1; CDK1; Cell division			
	con	trol protein 2 homolog; Cell div	vision protein kinase 1; p34 protein kinase			
Entrez Gene	983	(Human)				
SwissProt	P06	493 (Human)				
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery aliqu	uot and store at -20°C for one year. Avoid			
	free	ze/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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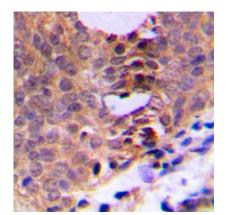
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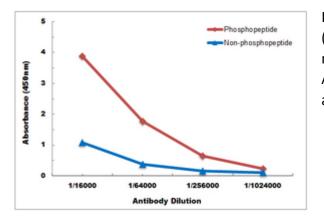
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Western blot analysis of CDC2 (Phospho-T161) expression in HepG2 (A) whole cell lysates. (Predicted band size: 34 kD; Observed band size: 34 kD)



Immunohistochemical analysis of CDC2 (Phospho-T161) 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.



Direct ELISA antibody dose-response curve using Anti-CDC2 (Phospho-T161) 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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