

Product Data Sheet

Anti-Cyclin D3 (Phospho-T283) Antibody

Catalog #	Source	Reactivity	Applications			
CPA7006	Rabbit	H, M, R, B	WB, IH			
Description	Ra	Rabbit polyclonal antibody to Cyclin D3 (Phospho-T283)				
Immunogen	KL	-H-conjugated synthetic pl	nosphopeptide corresponding to residues surrounding			
	Т2	283 of human Cyclin D3 pr	otein. The exact sequence is proprietary.			
Purification	Th	ne antibody was purified b	y immunogen affinity chromatography.			
Specificity	Re	ecognizes endogenous lev	els of Cyclin D3 protein only when phosphorylated at			
	Т2	283.				
Clonality	Pc	olyclonal				
Conjugation						
Form	Lic	quid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ar	nd 0.01% sodium azide.				
Dilution	W	/B (1/500 - 1/1000), IH (1/10	00 - 1/200)			
Gene Symbol	СС	CND3				
Alternative Na	ames G	1/S-specific cyclin-D3				
Entrez Gene	89	96 (Human); 12445 (Mous	e); 25193 (Rat)			
SwissProt	PB	30281 (Human); P30282 (N	Nouse); P48961 (Rat)			
Storage/Stabi	lity Sh	nipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	fre	eeze/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

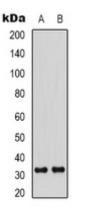
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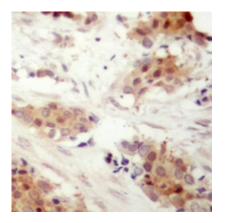
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Western blot analysis of Cyclin D3 (Phospho-T283) expression in HepG2 UV-treated (A), K562 UV-treated (B) whole cell lysates. (Predicted band size: 32 kD; Observed band size: 32 kD)



Immunohistochemical analysis of Cyclin D3 (Phospho-T283) 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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