

Product Data Sheet

Anti-Cyclin E1 Antibody

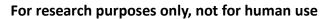
Catalog #	Source	Reactivity	Applications		
CPA5791	Rabbit	H, M, R, Mk	WB, IH, IF/IC		
Description	Rabb	it polyclonal antibody to	OCyclin E1		
Immunogen	KLH-c	conjugated synthetic pe	otide encompassing a sequence within the center		
	regio	n of human Cyclin E1. Tl	ne exact sequence is proprietary.		
Purification	The a	ntibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous leve	s of Cyclin E1 protein.		
Clonality	Polyc	lonal			
Conjugation					
Form	Liquid	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and O	.01% sodium azide.			
Dilution	WB (2	1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/100 - 1/500)		
Gene Symbol	CCNE	1			
Alternative Na	ames CCNE	; G1/S-specific cyclin-E1			
Entrez Gene	898 (Human); 12447 (Mouse)		
SwissProt	P248	64 (Human); Q61457 (N	louse); P39949 (Rat)		
Storage/Stabi	lity Shipp	ed at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	freeze	e/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

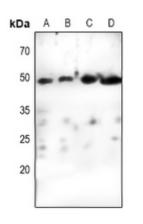
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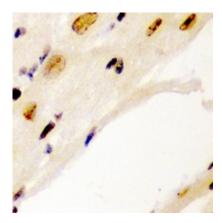
Coherion



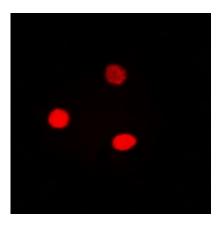
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Western blot analysis of Cyclin E1 expression in HepG2 (A), MCF7 (B), K562 (C), mouse lung (D) whole cell lysates. (Predicted band size: 47 kD; Observed band size: 48 kD)



Immunohistochemical analysis of Cyclin E1 staining in human heart 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 Cyclin E1 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 Alexa Fluor 647-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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