

Product Data Sheet

Anti-Cyclin A1/2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2335	Rabbit	H, M, R, B, C, Rb	WB, IH, IF/IC		
Description	Rabb	it polyclonal antibody to	Cyclin A1/2		
Immunogen	KLH-	conjugated synthetic pep	tide encompassing a sequence within the center		
	regio	region of human Cyclin A1/2. The exact sequence is proprietary.			
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous levels	s of Cyclin A1/2 protein.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqu	d in 0.42% Potassium pho	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB	1/500 - 1/1000), IH (1/50 -	1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	CCN	A1; CCNA2			
Alternative Na	ames CCN	CCNA1; Cyclin-A1; CCNA2; CCN1; CCNA; Cyclin-A2; Cyclin-A			
Entrez Gene 8900), 890 (Human); 12427, 12428 (Mouse); 295052 (Rat)			
SwissProt	P783	96, P20248 (Human); Q6	1456, P51943 (Mouse); Q6AY13 (Rat)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery	aliquot and store at -20°C for one year. Avoid		
	free	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

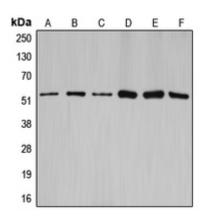
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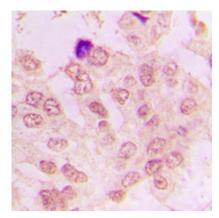


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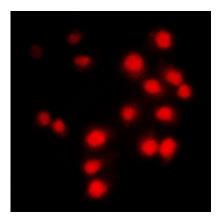
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Western blot analysis of Cyclin A1/2 expression in HEK293T (A), NIH3T3 (B), rat brain (C), SW626 (D), SKOV3 (E), A2780 (F) whole cell lysates. (Predicted band size: 52; 48 kD; Observed band size: 53 kD)



Immunohistochemical analysis of Cyclin A1/2 staining in human lung 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 Cyclin A1/2 staining in NIH3T3 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 humidified 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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