

## **Product Data Sheet**

## **Anti-CDCA7** Antibody

Catalog #	Source	Reactivity	Applications			
CPA3837	Rabbit	H, R, Mk	WB, IF/IC			
Description	Rabl	Rabbit polyclonal antibody to CDCA7				
Immunogen	KLH-	conjugated synthetic	peptide encompassing a sequence within the center			
	regio	on of human CDCA7. T	he exact sequence is proprietary.			
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous le	vels of CDCA7 protein.			
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), IF/IC (	1/100 - 1/500)			
Gene Symbol	CDC	A7				
Alternative Na	ames JPO2	L; Cell division cycle-as	sociated protein 7; Protein JPO1			
Entrez Gene	8387	83879 (Human); 66953 (Mouse); 311742 (Rat)				
SwissProt	Q9B	WT1 (Human); Q9D0N	12 (Mouse); Q4KM91 (Rat)			
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot 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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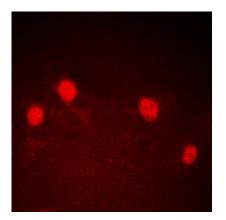
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Western blot analysis of CDCA7 expression in A431 (A), U2OS (B) whole cell lysates. (Predicted band size: 42 kD; Observed band size: 43 kD)



Immunofluorescent analysis of CDCA7 staining in A431 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. DAPI was used to stain the cell nuclei (blue).

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