

Product Data Sheet

Anti-CDC16 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3832	Rabbit	H, Mk, P	WB, IH, IF/IC		
Description	Ra	abbit polyclonal antibody t	o CDC16		
Immunogen	KL	H-conjugated synthetic pe	eptide encompassing a sequence within the C-term		
	re	gion of human CDC16. Th	e exact sequence is proprietary.		
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of CDC16 protein.		
Clonality	Рс	blyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 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	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	C	DC16			
Alternative Na	ames Al	NAPC6; Cell division cycle	protein 16 homolog; Anaphase-promoting complex		
	su	bunit 6; APC6; CDC16 hor	nolog; CDC16Hs; Cyclosome subunit 6		
Entrez Gene	88	381 (Human); 69957 (Mou	se)		
SwissProt	Q	13042 (Human); Q8R349 (Mouse)		
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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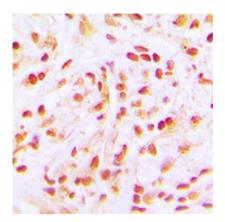
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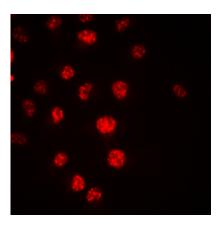
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Western blot analysis of CDC16 expression in HeLa (A), HT1080 (B) whole cell lysates. (Predicted band size: 71 kD; Observed band size: 72 kD)



Immunohistochemical analysis of CDC16 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 CDC16 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 a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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