

Product Data Sheet

Anti-CDC16 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6142	Rabbit	H, M, R, Z	WB, IH		
Description	F	Rabbit polyclonal antibody to	0 CDC16		
Immunogen	ł	KLH-conjugated synthetic pe	otide encompassing a sequence within the center		
	r	region of human CDC16. The	exact sequence is proprietary.		
Purification	7	The antibody was purified by	immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	ls of CDC16 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ä	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/200)		
Gene Symbol	(CDC16			
Alternative Na	ames /	ANAPC6; Cell division cycle p	rotein 16 homolog; Anaphase-promoting complex		
	S	subunit 6; APC6; CDC16 hom	olog; CDC16Hs; Cyclosome subunit 6		
Entrez Gene	8	8881 (Human); 69957 (Mous	e)		
SwissProt	(Q13042 (Human); Q8R349 (N	/louse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	freeze/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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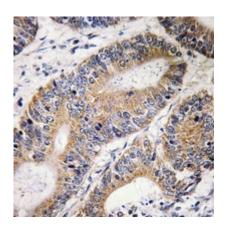
95

72

For research purposes only, not for human use

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Western blot analysis of CDC16 expression in MCF7 (A), A549 (B) whole cell lysates. (Predicted band size: 71 kD; Observed band size: 72 kD)



Immunohistochemical analysis of CDC16 staining in human colon 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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