

## **Product Data Sheet**

### **Anti-CDC46 Antibody**

Catalog #	Source	Reactivity	Applications	
CPA1730	Rabbit	H, M, R, B, C, Mk, Rb	WB, IH, IF/IC	
Description	F	Rabbit polyclonal antibody to (	CDC46	
Immunogen	ł	KLH-conjugated synthetic pept	ide encompassing a sequence within the N-term	
	r	region of human CDC46. The exact sequence is proprietary.		
Purification	٦	The antibody was purified by i	nmunogen affinity chromatography.	
Specificity	F	Recognizes endogenous levels	of CDC46 protein.	
Clonality	F	Polyclonal		
Conjugation				
Form	l	Liquid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,	
	ā	and 0.01% sodium azide.		
Dilution	١	WB (1/500 - 1/1000), IH (1/100 -	1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	ſ	MCM5		
Alternative Na	ames (	CDC46; DNA replication licensi	ng factor MCM5; CDC46 homolog; P1-CDC46	
Entrez Gene	Z	4174 (Human)		
SwissProt	F	P33992 (Human); P49718 (Mo	use)	
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery	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

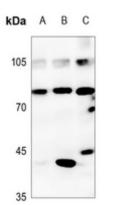
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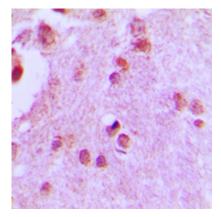


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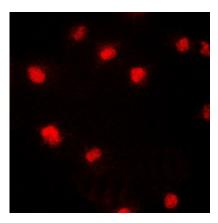
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Western blot analysis of CDC46 expression in mouse spleen (A), mouse kidney (B), rat spleen (C) whole cell lysates. (Predicted band size: 82 kD; Observed band size: 95 kD)



Immunohistochemical analysis of CDC46 staining in human brain 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 CDC46 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 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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