

Product Data Sheet

Anti-CCT2 Antibody

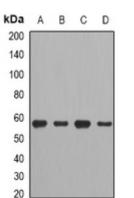
Catalog #	Source	Reactivity	Applications	
CQA1871	Rabbit	H <i>,</i> M, R	WB, IH, IF/IC	
Description	I	Rabbit polyclonal antibody	to CCT2	
Immunogen	I	Recombinant full length pro	tein of human CCT2	
Purification	-	The antibody was purified b	y immunogen affinity chromatography.	
Specificity	I	Recognizes endogenous lev	els of CCT2 protein.	
Clonality	I	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	,	WB (1/500 - 1/2000), IH (1/50) - 1/200), IF/IC (1/50 - 1/200)	
Gene Symbol	(CCT2		
Alternative Na	ames 🤅	99D8.1; CCTB; T-complex pı	otein 1 subunit beta; TCP-1-beta; CCT-beta	
Entrez Gene		10576 (Human); 12461 (Mc	use); 299809 (Rat)	
SwissProt	I	P78371 (Human); P80314 (I	Mouse); Q5XIM9 (Rat)	
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid	
	1	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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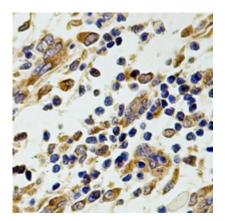




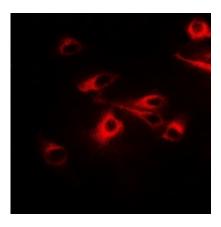
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Western blot analysis of CCT2 expression in A549 (A), HT29 (B), mouse spleen (C), mouse heart (D) whole cell lysates. (Predicted band size: 52; 57 kD; Observed band size: 57 kD)



Immunohistochemical analysis of CCT2 staining in rat stomach 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 CCT2 staining in A549 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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