

Product Data Sheet

Anti-Beta-adducin Antibody

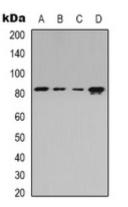
Catalog #	Source	Reactivity	Applications			
CPA7119	Rabbit	H, M, R	WB, IF/IC			
Description	Rabb	Rabbit polyclonal antibody to Beta-adducin				
Immunogen	KLH-	conjugated synthetic p	peptide encompassing a sequence within the center			
	regio	on of human Beta-add	ucin. The exact sequence is proprietary.			
Purification	The	antibody was purified	by immunogen affinity chromatography.			
Specificity	Reco	gnizes endogenous lev	vels of Beta-adducin protein.			
Clonality	Poly	clonal				
Conjugation						
Form	Liqui	d 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	ADD	2				
Alternative Na	ames ADD	B; Beta-adducin; Eryth	rocyte adducin subunit beta			
Entrez Gene	119	(Human); 11519 (Mou	se); 24171 (Rat)			
SwissProt	P356	512 (Human); Q9QYB8	(Mouse); Q05764 (Rat)			
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid			
	freez	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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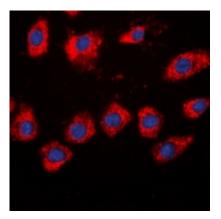




For research purposes only, not for human use

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Western blot analysis of Beta-adducin expression in Jurkat (A), HEK293T (B), Raw264.7 (C), rat spleen (D) whole cell lysates. (Predicted band size: 80 kD; Observed band size: 81 kD)



Immunofluorescent analysis of Beta-adducin 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. DAPI was used to stain the cell nuclei (blue).

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