

Product Data Sheet

Anti-Cadherin 7 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4369	Rabbit	H, M, R, C	WB, IH, IF/IC		
Description	Ra	Rabbit polyclonal antibody to Cadherin 7			
Immunogen	KL	-H-conjugated synthetic po	eptide encompassing a sequence within the C-term		
	re	region of human Cadherin 7. The exact sequence is proprietary.			
Purification	Tł	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of Cadherin 7 protein.		
Clonality	Pc	olyclonal			
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 (1/50) - 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	CI	DH7			
Alternative Na	ames CI	DH7L1; Cadherin-7			
Entrez Gene	10	1005 (Human); 241201 (Mouse); 29162 (Rat)			
SwissProt	Q	9ULB5 (Human); Q8BM92	(Mouse); Q5DWV2 (Rat)		
Storage/Stabi	lity Sh	ipped 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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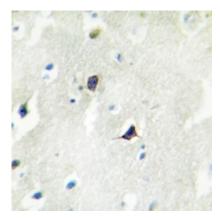
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Western blot analysis of Cadherin 7 expression in K562 (A), NIH3T3 (B) whole cell lysates. (Predicted band size: 87 kD; Observed band size: 87 kD)



Immunohistochemical analysis of Cadherin 7 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 Cadherin 7 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 hidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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