

Product Data Sheet

Anti-CDC42EP5 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4367	Rabbit	H, M, R	WB, IH		
Description	Ra	bbit polyclonal antibody to CDC42EP5			
Immunogen	KL	H-conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	re	gion of human CDC42EP5	The exact sequence is proprietary.		
Purification	Th	ne antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous leve	els of CDC42EP5 protein.		
Clonality	Pc	blyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	'B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	CE	DC42EP5			
Alternative N	ames BC	DRG3; CEP5; Cdc42 effecto	r protein 5; Binder of Rho GTPases 3		
Entrez Gene	14	18170 (Human)			
SwissProt	Q	6NZY7 (Human)			
Storage/Stabi	lity Sh	nipped 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

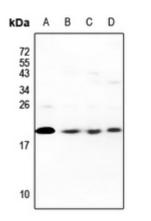
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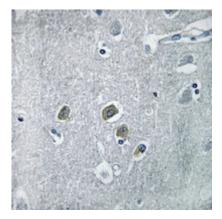


For research purposes only, not for human use

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Western blot analysis of CDC42EP5 expression in CT26 (A), PC12 (B), HepG2 (C), A549 (D) whole cell lysates. (Predicted band size: 15 kD; Observed band size: 19 kD)



Immunohistochemical analysis of CDC42EP5 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.

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