

Product Data Sheet

Anti-ARHGEF1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2358	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to ARHGEF1			
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
	I	region of human ARHGEF1. 1	he exact sequence is proprietary.		
Purification	-	The antibody was purified by	immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	s of ARHGEF1 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	v	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	1	ARHGEF1			
Alternative Na	ames	Rho guanine nucleotide exch	ange factor 1; 115 kDa guanine nucleotide exchange		
	1	factor; p115-RhoGEF; p115R	noGEF; Sub1.5		
Entrez Gene	(9138 (Human); 16801 (Mous	e)		
SwissProt	(Q92888 (Human); Q61210 (N	1ouse); Q9Z1I6 (Rat)		
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y 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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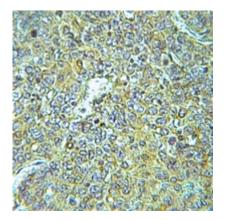
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kDa A B 130 95 Western blot analysis of ARHGEF1 expression in HuT78 (A), A549 (B) whole cell lysates. (Predicted band size: 102 kD; Observed band size: 125; 90 kD)



Immunohistochemical analysis of ARHGEF1 staining in human liver cancer 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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