

Product Data Sheet

Anti-ARHGAP22 Antibody

Catalog #	Source	Reactivity	Applications		
CPA2742	Rabbit	H, M, R	WB, IH		
Description	Rab	Rabbit polyclonal antibody to ARHGAP22			
Immunogen	KLH	-conjugated synthetic p	eptide encompassing a sequence within the C-term		
	regi	on of human ARHGAP2	2. The exact sequence is proprietary.		
Purification	The	antibody was purified	by immunogen affinity chromatography.		
Specificity	Reco	ognizes endogenous lev	els of ARHGAP22 protein.		
Clonality	Poly	rclonal			
Conjugation					
Form	Liqu	id 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), IH (1/1	00 - 1/200)		
Gene Symbol	ARH	IGAP22			
Alternative Na	ames RHC	GAP2; Rho GTPase-act	vating protein 22; Rho-type GTPase-activating protein 22		
Entrez Gene	585	04 (Human); 239027 (N	louse)		
SwissProt	Q7Z	5H3 (Human); Q8BL80	(Mouse)		
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid		
	free	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

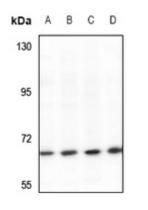
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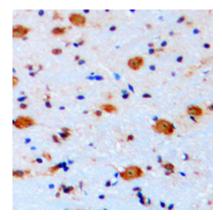


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Western blot analysis of ARHGAP22 expression in SP20 (A), PC12 (B), HEK293T (C), Hela (D) whole cell lysates. (Predicted band size: 76 kD; Observed band size: 68 kD)



Immunohistochemical analysis of ARHGAP22 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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