

Product Data Sheet

Anti-SHARP1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA3433	Rabbit				
	Nauni		WB, IH		
Description	escription Rabbit polyclonal antibody to SHARP1				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the N-term			
		region of human SHARP1. The exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity Recognizes endogenous levels of SHARP1 protein.			s of SHARP1 protein.		
Clonality		Polyclonal			
Conjugation					
Form		Liquid 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 (1/100) - 1/200)		
Gene Symbol		BHLHE41			
Alternative Na	ames	BHLHB3; DEC2; SHARP1; Clas	s E basic helix-loop-helix protein 41; bHLHe41; Class B		
		basic helix-loop-helix protein	3; bHLHb3; Differentially expressed in chondrocytes		
		protein 2; hDEC2; Enhancer-o	of-split and hairy-related protein 1; SHARP-1		
Entrez Gene		79365 (Human); 79362 (Mou	se)		
SwissProt		Q9C0J9 (Human); Q99PV5 (N	louse); O35779 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
		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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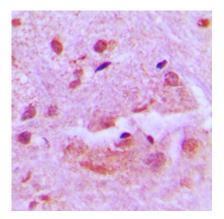
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Western blot analysis of SHARP1 expression in SHSY5Y (A), RAW264.7 (B), PC12 (C) whole cell lysates. (Predicted band size: 50 kD; Observed band size: 50 kD)



Immunohistochemical analysis of SHARP1 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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