

Product Data Sheet

Anti-SH2D4A Antibody

Catalog #	Source	Reactivity	Applications	
-		-		
CPA4232	Rabbit	H, M, R	WB, IH	
Description		Rabbit polyclonal antibody to	SH2D4A	
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the center	
		region of human SH2D4A. Th	e exact sequence is proprietary.	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	s of SH2D4A 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		SH2D4A		
Alternative Na	ames	PPP1R38; SH2A; SH2 domain	-containing protein 4A; Protein SH(2)A; Protein	
		phosphatase 1 regulatory sul	ounit 38	
Entrez Gene		63898 (Human); 72281 (Mouse); 306376 (Rat)		
SwissProt		Q9H788 (Human); Q9D7V1 (Mouse); Q6AYC8 (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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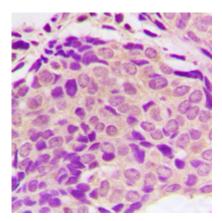
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Western blot analysis of SH2D4A expression in mouse kidney (A), mouse liver (B), rat kidney (C), rat liver (D) whole cell lysates. (Predicted band size: 52 kD; Observed band size: 52 kD)



Immunohistochemical analysis of SH2D4A staining in human breast 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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