

Product Data Sheet

Anti-PS1D Antibody

Catalog #	Source	e Reactivity	Applications	
CPA4316	Rabbit		WB, IH	
Description		Rabbit polyclonal antibody to	PS1D	
Immunogen		KLH-conjugated synthetic per	otide encompassing a sequence within the center	
	region of human PS1D. The exact sequence is proprietary.			
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous level	s of PS1D protein.	
Clonality		Polyclonal		
Conjugation				
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glyce			
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IH (1/100	- 1/200)	
Gene Symbol		ZCCHC17		
Alternative Na	ames	PS1D; Nucleolar protein of 40	kDa; pNO40; Pnn-interacting nucleolar protein;	
		Putative S1 RNA-binding dom	ain protein; PS1D protein; Zinc finger CCHC	
		domain-containing protein 17	7	
Entrez Gene		51538 (Human); 619605 (Mo	use)	
SwissProt		Q9NP64 (Human); Q9ESX4 (N	louse)	
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

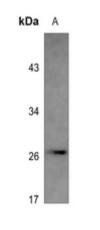
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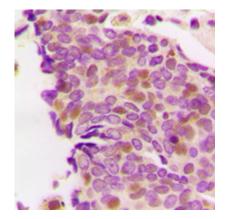
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Western blot analysis of PS1D expression in mouse spleen (A) whole cell lysates. (Predicted band size: 27 kD; Observed band size: 27 kD)



Immunohistochemical analysis of PS1D 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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