

Product Data Sheet

Anti-WIPF1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA5031	Rabbit	H, M, R	WB, IH		
Description		Rabbit polyclonal antibody to	WIPF1		
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the C-term		
		region of human WIPF1. The	exact sequence is proprietary.		
Purification		The antibody was purified by	immunogen affinity chromatography.		
Specificity		Recognizes endogenous leve	s of WIPF1 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 (1/10) - 1/200)		
Gene Symbol	,	WIPF1			
Alternative Na	ames	WASPIP; WIP; WAS/WASL-int	eracting protein family member 1; Protein PRPL-2;		
		Wiskott-Aldrich syndrome pr	otein-interacting protein; WASP-interacting protein		
Entrez Gene		7456 (Human); 215280 (Mou	se); 117538 (Rat)		
SwissProt	1	O43516 (Human); Q8K1I7 (N	ouse); Q6IN36 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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

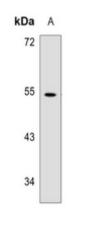
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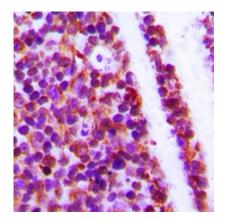
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Western blot analysis of WIPF1 expression in HL60 (A) whole cell lysates. (Predicted band size: 51 kD; Observed band size: 51 kD)



Immunohistochemical analysis of WIPF1 staining in human tonsil 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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