

Product Data Sheet

Anti-CD70 Antibody

Catalog #	Source	Reactivity	Applications		
CPA6022	Rabbit	Н, М	WB, IH		
Description	F	Rabbit polyclonal antibody	to CD70		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
	r	region of human CD70. The	exact sequence is proprietary.		
Purification	I	Γhe antibody was purified t	by immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous lev	els of CD70 protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	ā	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	0 - 1/100)		
Gene Symbol	(CD70			
Alternative Names		CD27L; CD27LG; TNFSF7; CD70 antigen; CD27 ligand; CD27-L; Tumor necrosis factor			
	I	ligand superfamily member	7; CD70		
Entrez Gene	ç	970 (Human); 21948 (Mous	e)		
SwissProt	F	P32970 (Human); O55237 (Mouse)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	f	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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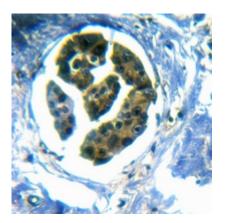
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For research purposes only, not for human use

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Western blot analysis of CD70 expression in Myla2059 (A), HCT116 (B), CT26 (C) whole cell lysates. (Predicted band size: 21 kD; Observed band size: 75; 95 kD)



Immunohistochemical analysis of CD70 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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