

Product Data Sheet

Anti-CD40 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1185	Rabbit	H, M, D, P	WB, IH, IF/IC		
Description	F	Rabbit polyclonal antibody t	o CD40		
Immunogen	K	(LH-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	r	region of human CD40. The exact sequence is proprietary.			
Purification	Т	The antibody was purified b	y immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	els of CD40 protein.		
Clonality	P	Polyclonal			
Conjugation					
Form	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)		
Gene Symbol	C	CD40			
Alternative Na	ames T	NFRSF5; Tumor necrosis far	ctor receptor superfamily member 5; B-cell surface		
	а	antigen CD40; Bp50; CD40L	receptor; CDw40; CD40		
Entrez Gene	9	958 (Human); 21939 (Mouse	e)		
SwissProt	P	25942 (Human); P27512 (N	1ouse)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	f	reeze/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

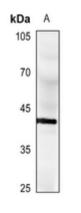
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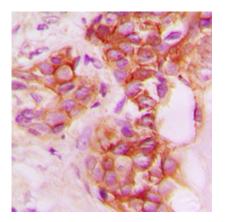
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For research purposes only, not for human use

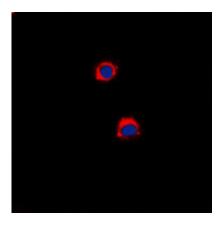
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Western blot analysis of CD40 expression in mouse pancreas (A) whole cell lysates. (Predicted band size: 30 kD; Observed band size: 42 kD)



Immunohistochemical analysis of CD40 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.



Immunofluorescent analysis of CD40 staining in A431 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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