

Product Data Sheet

Anti-CD275 Antibody

Catalog #	Source	Reactivity	Applications		
CQA3269	Rabbit	H <i>,</i> M, R	WB, IH		
Description	Ra	bbit polyclonal antibody	to CD275		
Immunogen	Re	combinant full length pro	tein of human CD275		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	ecognizes endogenous lev	els of CD275 protein.		
Clonality	Рс	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/2000), IH (1/50) - 1/200)		
Gene Symbol	IC	OSLG			
Alternative Na	ames B7	7H2; B7RP1; ICOSL; KIAA0	653; ICOS ligand; B7 homolog 2; B7-H2; B7-like protein		
	Gl	50; B7-related protein 1;	B7RP-1; CD275		
Entrez Gene	23	308 (Human); 50723 (Mc	use)		
SwissProt	07	75144 (Human); Q9JHJ8 (Mouse)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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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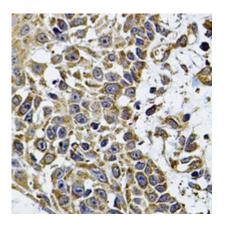
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A B C

For research purposes only, not for human use

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Western blot analysis of CD275 expression in HL60 (A), mouse brain (B), rat spleen (C) whole cell lysates. (Predicted band size: 20; 33; 34 kD; Observed band size: 60 kD)



Immunohistochemical analysis of CD275 staining in human skin 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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