

Product Data Sheet

Anti-CD253 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4282	Rabbit	H, M, R	WB, IH		
Description	ļ	Rabbit polyclonal antibody t	o CD253		
Immunogen	I	KLH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	I	region of human CD253. The	exact sequence is proprietary.		
Purification	-	The antibody was purified by	v immunogen affinity chromatography.		
Specificity	I	Recognizes endogenous leve	ls of CD253 protein.		
Clonality	I	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	i	and 0.01% sodium azide.			
Dilution	,	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	-	TNFSF10			
Alternative Na	ames	APO2L; TRAIL; Tumor necros	is factor ligand superfamily member 10; Apo-2 ligand;		
		Apo-2L; TNF-related apoptos	is-inducing ligand; Protein TRAIL; CD253		
Entrez Gene	:	8743 (Human); 22035 (Mous	e)		
SwissProt	I	P50591 (Human); P50592 (N	louse)		
Storage/Stabi	lity :	Shipped at 4°C. Upon deliver	y 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

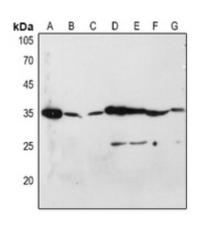
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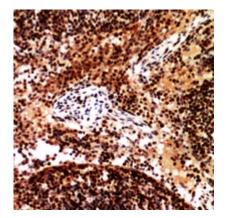
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Western blot analysis of CD253 expression in HEK293T (A), Hela (B), H460 (C), mouse kidney (D), mouse testis (E), rat kidney (F), rat testis (G) whole cell lysates. (Predicted band size: 32 kD; Observed band size: 33 kD)



Immunohistochemical analysis of CD253 staining in human lymph node 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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