

Product Data Sheet

Anti-CD294 Antibody

Catalog #	Source	e Reactivity	Applications		
CPA6480	Rabbit	H <i>,</i> R	WB, IH		
Description		Rabbit polyclonal antibody t	o CD294		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center			
		region of human CD294. The exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.			
Specificity		Recognizes endogenous levels of CD294 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 (1/50	- 1/200)		
Gene Symbol		PTGDR2			
Alternative Na	ames	CRTH2; DL1R; GPR44; Prosta	glandin D2 receptor 2; Chemoattractant		
		receptor-homologous moleo	cule expressed on Th2 cells; G-protein coupled receptor		
		44; CD294			
Entrez Gene		11251 (Human); 309212 (Ra	t)		
SwissProt		Q9Y5Y4 (Human); Q6XKD3 (Rat)		
Storage/Stabi	lity	Shipped at 4°C. Upon delive	ry 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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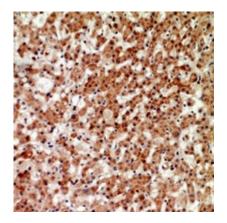


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kDa A B C 55 43 34 For research purposes only, not for human use

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Western blot analysis of CD294 expression in HEK293T (A), HuT78 (B), rat spleen (C) whole cell lysates. (Predicted band size: 43 kD; Observed band size: 43 kD)



Immunohistochemical analysis of CD294 staining in human liver 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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