

Product Data Sheet

Anti-CD179b Antibody

Catales #	Source	Doostivity	Applications		
Catalog #	Source		Applications		
CPA4018	Rabbit	н, М, R	WB, IH		
Description		Rabbit polyclonal antibody	to CD179b		
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the C-term			
		region of human CD179b. T	he exact sequence is proprietary.		
Purification		The antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	els of CD179b protein.		
Clonality Polyclonal					
Conjugation					
Form Lie		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) - 1/100)		
Gene Symbol		IGLL1			
Alternative Na	ames	IGL1; Immunoglobulin lamb	da-like polypeptide 1; CD179 antigen-like family		
		member B; Ig lambda-5; Im	nunoglobulin omega polypeptide;		
		Immunoglobulin-related pro	otein 14.1; CD179b		
Entrez Gene		3543 (Human); 16136 (Mou	se)		
SwissProt		P15814 (Human); P20764 (N	Aouse)		
Storage/Stabi	lity	Shipped at 4°C. Upon delivery 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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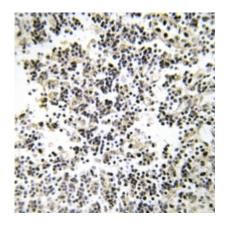
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kDa A B C D E

For research purposes only, not for human use

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Western blot analysis of CD179b expression in HEK293T (A), mouse liver (B), mouse spleen (C), rat liver (D), rat spleen (E) whole cell lysates. (Predicted band size: 22 kD; Observed band size: 23 kD)



Immunohistochemical analysis of CD179b 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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