

# **Product Data Sheet**

### Anti-CD179a Antibody

Catalog #	Source	e Reactivity	Applications		
CPA5841	Rabbit	t H	WB, IH		
Description		Rabbit polyclonal antibody	to CD179a		
Immunogen		KLH-conjugated synthetic p	eptide encompassing a sequence within the C-term		
		region of human CD179a.	he exact sequence is proprietary.		
Purification		The antibody was purified	by immunogen affinity chromatography.		
Specificity		Recognizes endogenous lev	vels of CD179a 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 (1/5	0 - 1/100)		
Gene Symbol		VPREB1			
Alternative Na	ames	VPREB; Immunoglobulin io	ta chain; CD179 antigen-like family member A; Protein		
		VPreB1; V(pre)B protein; V	preB protein; CD179a		
Entrez Gene		7441 (Human)			
SwissProt		P12018 (Human)			
Storage/Stabi	ility	Shipped at 4°C. Upon deliv	ery 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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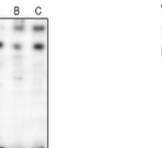
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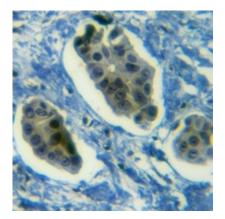
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For research purposes only, not for human use

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Western blot analysis of CD179a expression in Myla2059 (A), HuT78 (B), K562 (C) whole cell lysates. (Predicted band size: 16 kD; Observed band size: 16 kD)



Immunohistochemical analysis of CD179a 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.

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