

## **Product Data Sheet**

## **Anti-CD19 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA7303	Rabbit	H, M, R	WB, IH		
Description	R	Rabbit polyclonal antibody t	o CD19		
Immunogen	К	(LH-conjugated synthetic pe	ptide encompassing a sequence within the center		
	r	egion of human CD19. The	exact sequence is proprietary.		
Purification	Т	he antibody was purified b	y immunogen affinity chromatography.		
Specificity	R	Recognizes endogenous leve	ls of CD19 protein.		
Clonality	Р	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	а	and 0.01% sodium azide.			
Dilution	V	NB (1/500 - 1/1000), IH (1/10	0 - 1/200)		
Gene Symbol	C	CD19			
Alternative Na	a <mark>mes</mark> B	3-lymphocyte antigen CD19	B-lymphocyte surface antigen B4; Differentiation		
	а	intigen CD19; T-cell surface	antigen Leu-12; CD19		
Entrez Gene	9	930 (Human); 12478 (Mouse	2)		
SwissProt	Р	215391 (Human); P25918 (N	1ouse)		
Storage/Stabi	lity S	hipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fi	reeze/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

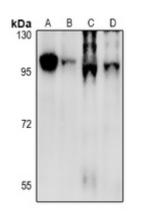
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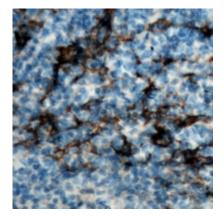
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Western blot analysis of CD19 expression in mouse spleen (A), rat spleen (B), SGC7901 (C), HCT116 (D) whole cell lysates. (Predicted band size: 61 kD; Observed band size: 100 kD)



Immunohistochemical analysis of CD19 staining in human tonsil 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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