

## **Product Data Sheet**

## **Anti-CLEC-6 Antibody**

Catalog #	Source	Reactivity	Applications
CQA4111	Rabbit	H, M, R	WB, IF/IC
Description	Rab	bit polyclonal antibody	to CLEC-6
Immunogen	Reco	ombinant fusion protei	n of human CLEC-6. The exact sequence is proprietary.
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	vels of CLEC-6 protein
Clonality	Poly	rclonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/2000), IF/IC (	1/50 - 1/200)
Gene Symbol	CLE	C4D	
Alternative N	ames CLEO	CSF8; MCL; C-type lecti	n domain family 4 member D; C-type lectin superfamily
	mer	nber 8; C-type lectin-lil	ke receptor 6; CLEC-6
Entrez Gene	338	339 (Human); 17474 (N	1ouse); 362432 (Rat)
SwissProt	Q8V	VXI8 (Human); Q9Z2H6	(Mouse); Q69FH1 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon deliv	ery aliquot and store at -20°C for one year. Avoid
	free	ze/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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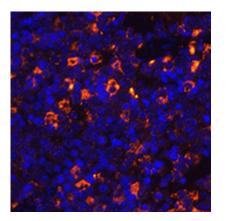
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17

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of CLEC-6 expression in HEK293T (A) whole cell lysates. (Predicted band size: 24 kD; Observed band size: 27 kD)



Immunofluorescent analysis of CLEC-6 staining in mouse spleen. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a humidified chamber. Cells were washed with PBST and incubated with a AF594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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