

# **Product Data Sheet**

## **Anti-EMR2** Antibody

Catalog #	Source	e Reactivity	Applications	
CPA3918	Rabbit	H, M, R, D	WB, IF/IC	
Description		Rabbit polyclonal antibody to	DEMR2	
Immunogen		KLH-conjugated synthetic pe	otide encompassing a sequence within the center	
		region of human EMR2. The	exact sequence is proprietary.	
Purification		The antibody was purified by	immunogen affinity chromatography.	
Specificity		Recognizes endogenous leve	ls of EMR2 protein.	
Clonality		Polyclonal		
Conjugation				
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycero			
		and 0.01% sodium azide.		
Dilution		WB (1/500 - 1/1000), IF/IC (1/5	50 - 1/200)	
Gene Symbol		EMR2		
Alternative Na	ames	EGF-like module-containing	nucin-like hormone receptor-like 2; EGF-like module	
		receptor 2; CD312		
Entrez Gene		30817 (Human)		
SwissProt		Q9UHX3 (Human)		
Storage/Stabi	lity	Shipped at 4°C. Upon deliver	y 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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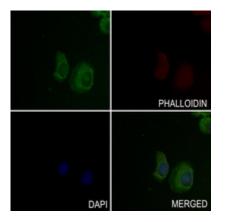
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**kDa** A B C D E 130 105 70 45 For research purposes only, not for human use

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Western blot analysis of EMR2 expression in 3T3L1 (A), PMVEC (B), A549 (C), K562 (D), Jurkat (E) whole cell lysates. (Predicted band size: 90 kD; Observed band size: 100 kD)



Immunofluorescent analysis of EMR2 staining in A549 cells. 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 hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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