

## Recombinant Anti-VEGFR1 Rabbit mAb

Catalog #	Source	Reactivity	Applications
CMA4478	Rabbit	H, M, R	WB, IF/IC
<b>Description</b>	Recombinant rabbit monoclonal antibody to VEGFR1		
<b>Immunogen</b>	KLH-conjugated synthetic peptide encompassing a sequence within human VEGFR1 protein. The exact sequence is proprietary.		
<b>Purification</b>	The antibody was purified by immunogen affinity chromatography.		
<b>Specificity</b>	Recognizes endogenous levels of VEGFR1 protein		
<b>Clonality</b>	Monoclonal		
<b>Conjugation</b>			
<b>Form</b>	Liquid in PBS containing 50% glycerol, 0.2% BSA and 0.01% sodium azide.		
<b>Dilution</b>	WB (1/500 - 1/1000), IF/IC (1/50 - 1/200)		
<b>Gene Symbol</b>	FLT1		
<b>Alternative Names</b>	FLT; FRT; VEGFR1; Vascular endothelial growth factor receptor 1; VEGFR-1; Fms-like tyrosine kinase 1; FLT-1; Tyrosine-protein kinase FRT; Tyrosine-protein kinase receptor FLT; FLT; Vascular permeability factor receptor		
<b>Entrez Gene</b>	2321 (Human); 14254 (Mouse); 54251 (Rat)		
<b>SwissProt</b>	P17948 (Human); P35969 (Mouse); P53767 (Rat)		
<b>Storage/Stability</b>	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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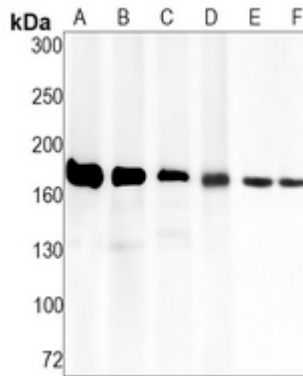
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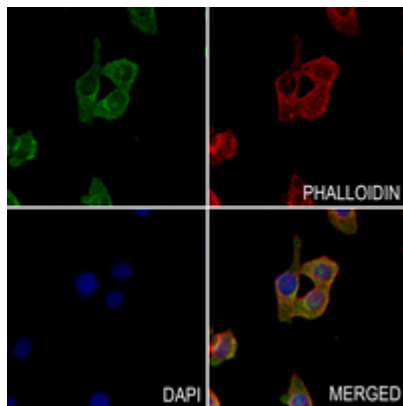
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# Product Data Sheet



Western blot analysis of VEGFR1 expression in HEK293T (A), Jurkat (B), K562 (C), mouse kidney (D), mouse muscle (E), rat muscle (F) whole cell lysates. (Predicted band size: 150 kD; Observed band size: 180 kD)



Immunofluorescent analysis of VEGFR1 staining in MCF7 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 humidified 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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