

Product Data Sheet

Anti-VEGFR1 Antibody

Catalog #	Source	Reactivity	Applications
CPA4411	Rabbit	H, M, R, C, Mk, P, S	WB, IH
Description	F	Rabbit polyclonal antibody to \	/EGFR1
Immunogen	K	KLH-conjugated synthetic pept	de encompassing a sequence within the C-term
	r	region of human VEGFR1. The	exact sequence is proprietary.
Purification	Т	The antibody was purified by ir	nmunogen affinity chromatography.
Specificity	F	Recognizes endogenous levels	of VEGFR1 protein.
Clonality	P	Polyclonal	
Conjugation			
Form	L	iquid in 0.42% Potassium pho	sphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	а	and 0.01% sodium azide.	
Dilution	V	WB (1/500 - 1/1000), IH (1/50 - 1	/100)
Gene Symbol	F	ELT1	
Alternative Na	ames F	ELT; FRT; VEGFR1; Vascular end	othelial growth factor receptor 1; VEGFR-1; Fms-like
	t	zyrosine kinase 1; FLT-1; Tyrosir	ne-protein kinase FRT; Tyrosine-protein kinase
	r	receptor FLT; FLT; Vascular perr	neability factor receptor
Entrez Gene	2	2321 (Human); 14254 (Mouse)	; 54251 (Rat)
SwissProt	P	P17948 (Human); P35969 (Mo	use); P53767 (Rat)
Storage/Stabi	l <mark>ity</mark> S	Shipped at 4°C. Upon delivery a	aliquot and store at -20°C for one year. Avoid
	f	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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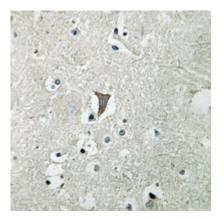
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For research purposes only, not for human use

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Western blot analysis of VEGFR1 expression in HEK293T (A) whole cell lysates. (Predicted band size: 150 kD; Observed band size: 150 kD)



Immunohistochemical analysis of VEGFR1 staining in human brain 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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