

## **Product Data Sheet**

## Anti-VEGFD Antibody

Catalog #	Source	Reactivity	Applications			
CPA1428	Rabbit	H, R, B	WB, IH, IF/IC			
Description	F	Rabbit polyclonal antibody	to VEGFD			
Immunogen	ŀ	KLH-conjugated synthetic pe	eptide encompassing a sequence within the center			
	r	region of human VEGFD. Th	e exact sequence is proprietary.			
Purification		The antibody was purified by immunogen affinity chromatography.				
Specificity		Recognizes endogenous levels of VEGFD protein.				
Clonality	F	Polyclonal				
Conjugation						
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,				
	ā	and 0.01% sodium azide.				
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200)				
Gene Symbol	F	FIGF				
Alternative Na	ames \	VEGFD; Vascular endothelia	l growth factor D; VEGF-D; c-Fos-induced growth factor;			
	F	FIGF				
Entrez Gene		2277 (Human)				
SwissProt O43915 (Human)		O43915 (Human)				
Storage/Stabi	ility S	Shipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid			
	f	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

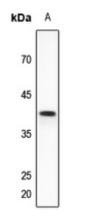
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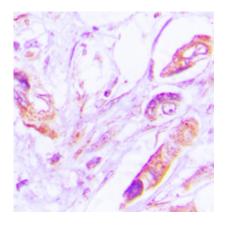
# Cohesion

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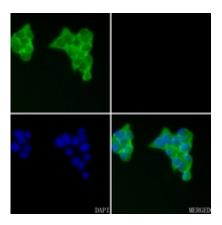
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Western blot analysis of VEGFD expression in rat lung (A) whole cell lysates. (Predicted band size: 40 kD; Observed band size: 40 kD)



Immunohistochemical analysis of VEGFD staining in human lung cancer 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.



Immunofluorescent analysis of VEGFD 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 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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