

## **Product Data Sheet**

### **Anti-Neuropilin 2 Antibody**

Catalog #	Source	Reactivity	Applications		
CPA4135	Rabbit	H, M, R, P	WB, IH		
Description	Ra	Rabbit polyclonal antibody to Neuropilin 2			
Immunogen	KL	H-conjugated synthetic pe	ptide encompassing a sequence within the center		
	re	gion of human Neuropilin	2. The exact sequence is proprietary.		
Purification	Th	ie antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous leve	els of Neuropilin 2 protein.		
Clonality	Ро	blyclonal			
Conjugation					
Form	Lic	quid in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	NF	RP2			
Alternative Na	ames VE	GF165R2; Neuropilin-2; V	ascular endothelial cell growth factor 165 receptor 2		
Entrez Gene	88	328 (Human); 18187 (Mou	se); 81527 (Rat)		
SwissProt	06	60462 (Human); O35375 (I	Mouse); O35276 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

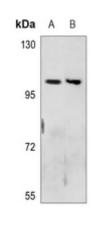
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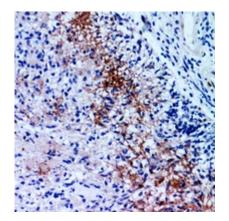
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Western blot analysis of Neuropilin 2 expression in A375 (A), DLD (B) whole cell lysates. (Predicted band size: 104 kD; Observed band size: 105 kD)



Immunohistochemical analysis of Neuropilin 2 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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