

Product Data Sheet

Anti-Ephrin B1 (Phospho-Y317) Antibody

Catalog #	Source	Reactivity	Applications		
CPA1369	Rabbit	H, M, R, C, Z	WB, IH		
Description	F	Ephrin B1 (Phospho-Y317)			
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding			
	Y	Y317 of human Ephrin B1 protein. The exact sequence is proprietary.			
Purification	Т	The antibody was purified by	immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous levels of Ephrin B1 protein only when phosphorylated at			
	Y	Y317.			
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	E	EFNB1			
Alternative Names		EFL3; EPLG2; LERK2; Ephrin-B1; EFL-3; ELK ligand; ELK-L; EPH-related receptor			
	t	tyrosine kinase ligand 2; LER	K-2		
Entrez Gene	1	1947 (Human); 13641 (Mous	e)		
SwissProt	F	P98172 (Human); P52795 (M	ouse); P52796 (Rat)		
Storage/Stabi	lity S	Shipped at 4°C. Upon delivery 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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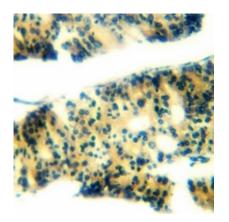
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Western blot analysis of Ephrin B1 (Phospho-Y317) expression in zebrafish (A) whole cell lysates. (Predicted band size: 38 kD; Observed band size: 45 kD)



Immunohistochemical analysis of Ephrin B1 (Phospho-Y317) staining in human colorectal 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.

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