

Product Data Sheet

Anti-Ephrin B3 Antibody

Catalog #	Source	Reactivity	Applications		
CPA1370	Rabbit	H, M, R, B, D, P, Rb	WB, IH		
Description	Rab	bit polyclonal antibody to Ep	bhrin B3		
Immunogen	KLH	-conjugated synthetic peptic	le encompassing a sequence within the C-term		
	regi	on of human Ephrin B3. The	exact sequence is proprietary.		
Purification	The	antibody was purified by im	munogen affinity chromatography.		
Specificity	Rec	ognizes endogenous levels o	f Ephrin B3 protein.		
Clonality	Polyclonal				
Conjugation					
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/50 - 1/	100)		
Gene Symbol	EFN	B3			
Alternative Na	ames EPL	G8; LERK8; Ephrin-B3; EPH-r	elated receptor transmembrane ligand ELK-L3;		
	EPH	I-related receptor tyrosine ki	nase ligand 8; LERK-8		
Entrez Gene	194	9 (Human); 13643 (Mouse)			
SwissProt	Q15	5768 (Human); O35393 (Mou	ise)		
Storage/Stabi	lity Ship	oped at 4°C. Upon delivery a	iquot and store at -20°C for one year. Avoid		
	free	eze/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

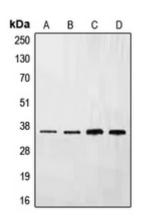
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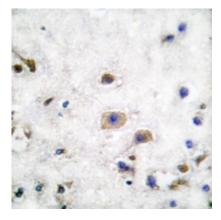


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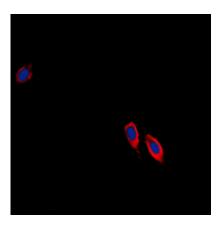
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Western blot analysis of Ephrin B3 expression in HepG2 (A), NIH3T3 (B), mouse liver (C), rat liver (D) whole cell lysates. (Predicted band size: 35 kD; Observed band size: 36 kD)



Immunohistochemical analysis of Ephrin B3 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.



Immunofluorescent analysis of Ephrin B3 staining in NIH3T3 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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