

Product Data Sheet

Anti-BMPR1B Antibody

Catalog #	Source	Reactivity	Applications
CQA1854	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rabb	it polyclonal antibody t	O BMPR1B
Immunogen	Reco	mbinant full length pro	tein of human BMPR1B
Purification	The	antibody was purified b	y immunogen affinity chromatography.
Specificity	Reco	gnizes endogenous leve	els of BMPR1B protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqui	d in 0.42% Potassium p	hosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB (1/500 - 1/2000), IH (1/50	- 1/200), IF/IC (1/50 - 1/200)
Gene Symbol	BMP	R1B	
Alternative Na	ames Bone	e morphogenetic protei	n receptor type-1B; BMP type-1B receptor; BMPR-1B;
	CDw	293	
Entrez Gene	658	(Human); 12167 (Mous	e)
SwissProt	0002	238 (Human); P36898 (I	Mouse)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid
	freez	e/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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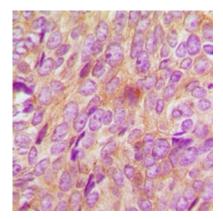
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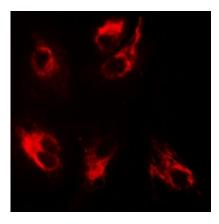
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Western blot analysis of BMPR1B expression in mouse brain (A) whole cell lysates. (Predicted band size: 56; 60 kD; Observed band size: 51 kD)



Immunohistochemical analysis of BMPR1B staining in human breast 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 BMPR1B 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 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.

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