

Product Data Sheet

Anti-Beta-2 Adrenergic Receptor Antibody

Catalog #	Source	Reactivity	Applications		
CPA3117	Rabbit	H, M, R, B, P	WB, IH, IF/IC		
Description	R	abbit polyclonal antibody to	t polyclonal antibody to Beta-2 Adrenergic Receptor		
Immunogen	K	LH-conjugated synthetic per	otide encompassing a sequence within the center		
	re	egion of human Beta-2 Adre	nergic Receptor. The exact sequence is proprietary.		
Purification	TI	he antibody was purified by	immunogen affinity chromatography.		
Specificity	R	ecognizes endogenous level	s of Beta-2 Adrenergic Receptor protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	aı	nd 0.01% sodium azide.			
Dilution	W	VB (1/500 - 1/1000), IH (1/50 ·	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	А	DRB2			
Alternative Na	ames A	DRB2R; B2AR; Beta-2 adren	ergic receptor; Beta-2 adrenoreceptor; Beta-2		
	а	drenoceptor			
Entrez Gene	1	54 (Human); 11555 (Mouse			
SwissProt	P	07550 (Human); P18762 (M	ouse); P10608 (Rat)		
Storage/Stabi	lity Sł	hipped at 4°C. Upon deliver	v aliquot and store at -20°C for one year. Avoid		
	fr	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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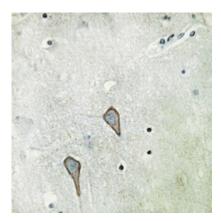
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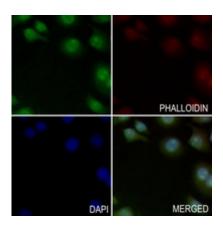
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Western blot analysis of Beta-2 Adrenergic Receptor expression in HeLa (A), Raw264.7 (B) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 68 kD)



Immunohistochemical analysis of Beta-2 Adrenergic Receptor 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 Beta-2 Adrenergic Receptor staining in A549 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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