

Product Data Sheet

Anti-Beta-2 Adrenergic Receptor (Phospho-S355/S356) Antibody

Catalog #	Source	Reactivity	Applications
CPA5865	Rabbit	H, M, R	WB, IH
Description	Rabbit polyclonal antibody to Beta-2 Adrenergic Receptor (Phospho-S355/S356)		
Immunogen	KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding		
	S355/S356 of human Beta-2 Adrenergic Receptor protein. The exact sequence is		

proprietary.

Purification The antibody was purified by immunogen affinity chromatography.

Specificity Recognizes endogenous levels of Beta-2 Adrenergic Receptor protein only when

phosphorylated at S355/S356.

Clonality Polyclonal

Conjugation

Form 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 - 1/200)

Gene Symbol ADRB2

Alternative Names ADRB2R; B2AR; Beta-2 adrenergic receptor; Beta-2 adrenoreceptor; Beta-2

adrenoceptor

Entrez Gene 154 (Human); 11555 (Mouse)

SwissProt P07550 (Human); P18762 (Mouse); P10608 (Rat)

Storage/Stability Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid

freeze/thaw cycles.

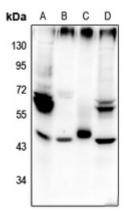
Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC- Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, 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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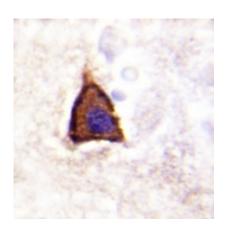
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Western blot analysis of Beta-2 Adrenergic Receptor (Phospho-S355/S356) expression in Hela (A), rat prostate (B), mouse lung (C), EC9706 (D) whole cell lysates. (Predicted band size: 46 kD; Observed band size: 47; 68 kD)



Immunohistochemical analysis of Beta-2 Adrenergic Receptor (Phospho-S355/S356) 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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