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## $\beta$ 2-Adrenergic Receptor (D6H2) Rabbit mAb

For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W	<b>Reactivity:</b> H	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 50-100	<b>Source/Isotype:</b> Rabbit IgG	<b>UniProt ID:</b> #P07550	<b>Entrez-Gene Id:</b> 154
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<b>Product Usage Information</b>	<b>Application</b> Western Blotting	<b>Dilution</b> 1:1000
<b>Storage</b>	Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at $-20^{\circ}\text{C}$ . Do not aliquot the antibody.	
<b>Specificity/Sensitivity</b>	$\beta$ 2-Adrenergic Receptor (D6H2) Rabbit mAb recognizes endogenous levels of total $\beta$ 2-adrenergic receptor protein.	
<b>Source / Purification</b>	Monoclonal antibody is produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human $\beta$ 2-adrenergic receptor protein.	
<b>Background</b>	<p>There are four major Adrenergic Receptor (AR) subtypes (<math>\alpha</math>1, <math>\alpha</math>2, <math>\beta</math>1, <math>\beta</math>2). Each of the subtypes has been classified by their unique responses to agonists and antagonists. Adrenergic receptors belong to the family of guanine nucleotide-binding, regulatory protein-coupled receptors (GPCR) which transverse the plasma membrane seven times. The transmembrane regions are hydrophobic and are interconnected by hydrophilic loops (1). <math>\beta</math>2-Adrenergic Receptor (<math>\beta</math>2AR) is the most studied receptor of the catecholamine system. <math>\beta</math>2AR stimulation occurs through the catecholamines epinephrine (adrenaline) and norepinephrine (noradrenaline) acting as neuromodulators in the central nervous system and as hormones in the vascular system. <math>\beta</math>2AR activation results in coupling to heterotrimeric G proteins and activation of the second messengers cAMP and phosphatidylinositol, ultimately leading to changes in cellular physiology. GPCR kinases (GRKs) terminate <math>\beta</math>2AR signaling through phosphorylation of the GPCR and by recruiting <math>\beta</math>-arrestin. <math>\beta</math>-arrestin binding uncouples the receptor from the G protein, thereby terminating G protein-mediated signaling (desensitization), and initiating clathrin-mediated endocytosis (internalization) of <math>\beta</math>2AR (2). <math>\beta</math>-adrenergic blocking agents (beta blockers) are drugs that block catecholamines from binding to <math>\beta</math>AR and are prescribed for cardiac arrhythmias, cardioprotection after myocardial infarction (heart attack), and hypertension (3).</p>	
<b>Background References</b>	<ol style="list-style-type: none"> <li>1. Dohlman, H.G. et al. (1987) <i>Biochemistry</i> 26, 2657-64.</li> <li>2. Nobles, K.N. et al. (2011) <i>Sci Signal</i> 4, ra51.</li> <li>3. Baker, J.G. et al. (2011) <i>Trends Pharmacol Sci</i> 32, 227-34.</li> </ol>	
<b>Species Reactivity</b>	Species reactivity is determined by testing in at least one approved application (e.g., western blot).	
<b>Western Blot Buffer</b>	IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v BSA, 1X TBS, 0.1% Tween@ 20 at $4^{\circ}\text{C}$ with gentle shaking, overnight.	
<b>Applications Key</b>	<b>W:</b> Western Blotting	
<b>Cross-Reactivity Key</b>	<b>H:</b> Human	
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