



E92048

ADRB2 Polyclonal Antibody

Catalog Number: E92048

Amount: 100ul

Background: There are four major Adrenergic Receptor (AR) subtypes ($\alpha 1$, $\alpha 2$, $\beta 1$, $\beta 2$). 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). $\beta 2$ -Adrenergic Receptor ($\beta 2AR$) is the most studied receptor of the catecholamine system. $\beta 2AR$ 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. $\beta 2AR$ 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 $\beta 2AR$ signaling through phosphorylation of the GPCR and by recruiting β -arrestin. β -arrestin binding uncouples the receptor from the G protein, thereby terminating G protein-mediated signaling (desensitization), and initiating clathrin-mediated endocytosis (internalization) of $\beta 2AR$ (2). β -adrenergic blocking agents (beta blockers) are drugs that block catecholamines from binding to βAR and are prescribed for cardiac arrhythmias, cardioprotection after myocardial infarction (heart attack), and hypertension (3).

Species: Rabbit

Isotype: IgG

Storage/Stability: Store at -20oC or -80oC. Avoid freeze / thaw cycles. Buffer: PBS with 0.02% sodium azide, 50% glycerol, pH7.3.

Synonyms: ADRB2R; ADRBR; B2AR; BAR; BETA2AR;

Immunogen: A synthetic peptide of human ADRB2

Purification: Affinity purification

Reactivity: H M R

Applications: WB IHC

Molecular Weight: 47kDa

Swiss-Prot No. : P07550

Gene ID: 154

References: 1. Dohlman, H.G. et al. (1987) Biochemistry 26, 2657-64. 2. Nobles, K.N. et al. (2011) Sci Signal 4, ra51. 3. Baker, J.G. et al. (2011) Trends Pharmacol Sci 32, 227-34.

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