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Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody

Our Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 primary antibody from PhosphoSolutions is rabbi

Catalog # AN1472

Specification

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody - Product Information

Primary Accession	Q9ERK7
Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Calculated MW	57113

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody - Additional Information

Gene ID	11444

Other Names

Acetylcholine receptor beta 2 neural antibody, ACHB2_HUMAN antibody, ACHN antibody, AChR antibody, Acrb 2 antibody, Acrb2 antibody, b2 nAchR antibody, Cholinergic receptor nicotinic beta 2 antibody, Cholinergic receptor nicotinic beta polypeptide 2 antibody, Cholinergic receptor nicotinic beta polypeptide 2 neuronal antibody, cholinergic receptor nicotinic beta 2 (neuronal) antibody, Chrnb2 antibody, EFNL 3 antibody, EFNL3 antibody, nAChRB2 antibody, Neuronal acetylcholine receptor protein beta 2 chain precursor antibody, Neuronal acetylcholine receptor protein subunit beta 2 antibody, Neuronal acetylcholine receptor subunit beta-2 antibody, Neuronal nicotinic acetylcholine receptor beta 2 antibody

Target/Specificity

Nicotinic acetylcholine receptors (nAChRs) are ionotropic, cholinergic receptors that are divided into 2 types; muscle type and neuronal type. Neuronal nAChRs are pentameric ion channels consisting of 5 identical (homopentamers) or different (heteropentamers) subunits. Heteropentameric neuronal nAChRs mediate fast synaptic transmission in the autonomic nervous system. The predominant hetero-oligomeric nAChR in the CNS contain the subunits α4β2, whereas α3β4 prevail in the PNS. However, the expression of these subunits varies not only by region but also during development (Scholze et al 2011). In the brain, β2-containing receptors greatly outnumber receptors that contain β4 (McGehee & Role, 1995; Albuquerque, et al., 2009), and in most brain regions, targeted deletion of the β2 subunit virtually abolishes [3H]-epibatidine binding and receptor autoradiography (Zoli, et al., 1998) due to the absence of a β subunit required to form functional nAChRs (Champtiaux & Changeux, 2004).

Format

Antigen Affinity Purified from Pooled Serum

Storage

Maintain refrigerated at 2-8°C for up to 6 months. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

Precautions

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

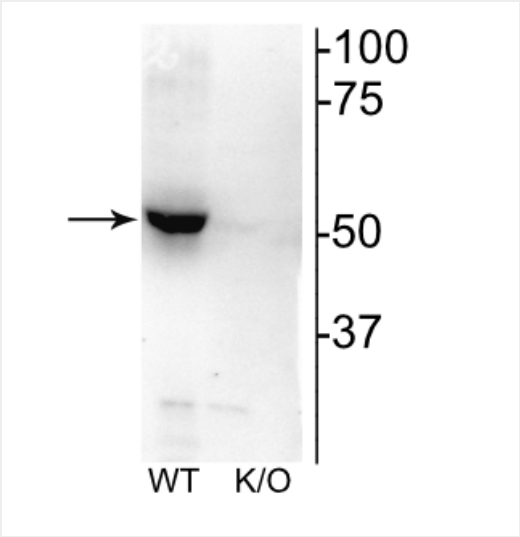
Shipping
Blue Ice

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- Immunoprecipitation
- Flow Cytomety
- Cell Culture

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody - Images



Western blot of mouse habenula lysate showing specific immunolabeling of the ~52 kDa nAChRβ2 protein.

Anti-Nicotinic Acetylcholine Receptor (nAChR) β2 Antibody - Background

Nicotinic acetylcholine receptors (nAChRs) are ionotropic, cholinergic receptors that are divided into 2 types; muscle type and neuronal type. Neuronal nAChRs are pentameric ion channels consisting of 5 identical (homopentamers) or different (heteropentamers) subunits. Heteropentameric neuronal nAChRs mediate fast synaptic transmission in the autonomic nervous system. The predominant hetero-oligomeric nAChR in the CNS contain the subunits α4β2, whereas α3β4 prevail in the PNS. However, the expression of these subunits varies not only by region but also during development (Scholze et al 2011). In the brain, β2-containing receptors greatly outnumber receptors that contain β4 (McGehee & Role, 1995; Albuquerque, et al., 2009), and in most brain regions, targeted deletion of the β2 subunit virtually abolishes [3H]-epibatidine binding and receptor autoradiography (Zoli, et al., 1998) due to the absence of a β subunit required to form functional nAChRs (Champtiaux & Changeux, 2004).