

CHRNA7 Antibody (N-term)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP13898a**Specification**

CHRNA7 Antibody (N-term) - Product Information

Application	WB,E
Primary Accession	P36544
Other Accession	NP_000737.1
Reactivity	Human
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit IgG
Calculated MW	56449
Antigen Region	8-37

CHRNA7 Antibody (N-term) - Additional Information**Gene ID** 1139;89832**Other Names**

Neuronal acetylcholine receptor subunit alpha-7, CHRNA7, NACHRA7

Target/Specificity

This CHRNA7 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 8-37 amino acids from the N-terminal region of human CHRNA7.

Dilution

WB~~1:1000

E~~Use at an assay dependent concentration.

Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

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

Precautions

CHRNA7 Antibody (N-term) is for research use only and not for use in diagnostic or therapeutic procedures.

CHRNA7 Antibody (N-term) - Protein Information**Name** CHRNA7 ([HGNC:1960](#))**Synonyms** NACHRA7

Function Component of neuronal acetylcholine receptors (nAChRs) that function as pentameric, ligand-gated cation channels with high calcium permeability among other activities. nAChRs are excitatory neurotransmitter receptors formed by a collection of nAChR subunits known to mediate synaptic transmission in the nervous system and the neuromuscular junction. Each nAChR subunit confers differential attributes to channel properties, including activation, deactivation and desensitization kinetics, pH sensitivity, cation permeability, and binding to allosteric modulators (PubMed:[15609996](#), PubMed:[33735609](#), PubMed:[8145738](#)). CHRNA7 forms homopentameric neuronal acetylcholine receptors abundantly expressed in the central nervous system, characterized by fast desensitization and high calcium permeability (PubMed:[31560909](#), PubMed:[33735609](#), PubMed:[38382524](#), PubMed:[8145738](#)). Also forms heteropentamers with CHRNB2, mainly expressed in basal forebrain cholinergic neurons. Involved in the modulation of calcium-dependent signaling pathways and influences the release of neurotransmitters, including dopamine, glutamate and GABA (PubMed:[33239400](#)). Also expressed in non-neuronal cells such as immune cells like lymphocytes, monocytes and macrophages (PubMed:[12508119](#), PubMed:[16968406](#), PubMed:[25259522](#)). In T cells, activation induces metabotropic signaling that results in an increase of intracellular Ca²⁺ concentrations, independent of ionotropic receptor functions (PubMed:[17709503](#)). In macrophages, required for acetylcholine-mediated inhibition of TNF and other inflammatory cytokine release (PubMed:[12508119](#)). Once activated by acetylcholine, nicotine or other agonists, selectively inhibits production of pro-inflammatory cytokines while leaving anti-inflammatory cytokines undisturbed (PubMed:[12508119](#), PubMed:[25259522](#)). Stimulates the cholinergic anti-inflammatory pathway, controlling inflammation by inhibiting NFκB nuclear translocation and activating the JAK2-STAT3 pathway, independently of ion channel activity (PubMed:[16968406](#), PubMed:[25259522](#)). Also expressed in the urothelium where it modulates reflex bladder activity by increasing intracellular calcium through internal stores and decreasing basal ATP release (By similarity).

Cellular Location

Postsynaptic cell membrane {ECO:0000250|UniProtKB:Q05941}; Multi-pass membrane protein. Cell membrane; Multi-pass membrane protein. Note=TMEM35A/NACHO promotes its trafficking to the cell membrane (PubMed:27789755). RIC3 promotes its trafficking to the cell membrane (By similarity) {ECO:0000250|UniProtKB:Q05941, ECO:0000269|PubMed:27789755}

Tissue Location

Expressed in neuronal cells (PubMed:8145738). Expressed in macrophages (at protein level) (PubMed:12508119)

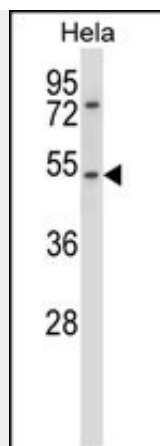
CHRNA7 Antibody (N-term) - Protocols

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

CHRNA7 Antibody (N-term) - Images





CHRNA7 Antibody (N-term) (Cat. #AP13898a) western blot analysis in HeLa cell line lysates (35ug/lane). This demonstrates the CHRNA7 antibody detected the CHRNA7 protein (arrow).

CHRNA7 Antibody (N-term) - Background

The nicotinic acetylcholine receptors (nAChRs) are members of a superfamily of ligand-gated ion channels that mediate fast signal transmission at synapses. The nAChRs are thought to be hetero-pentamers composed of homologous subunits. The proposed structure for each subunit is a conserved N-terminal extracellular domain followed by three conserved transmembrane domains, a variable cytoplasmic loop, a fourth conserved transmembrane domain, and a short C-terminal extracellular region. The protein encoded by this gene forms a homo-oligomeric channel, displays marked permeability to calcium ions and is a major component of brain nicotinic receptors that are blocked by, and highly sensitive to, alpha-bungarotoxin. Once this receptor binds acetylcholine, it undergoes an extensive change in conformation that affects all subunits and leads to opening of an ion-conducting channel across the plasma membrane. This gene is located in a region identified as a major susceptibility locus for juvenile myoclonic epilepsy and a chromosomal location involved in the genetic transmission of schizophrenia. An evolutionarily recent partial duplication event in this region results in a hybrid containing sequence from this gene and a novel FAM7A gene. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

CHRNA7 Antibody (N-term) - References

Chernyavsky, A.I., et al. Am. J. Physiol., Cell Physiol. 299 (5), C903-C911 (2010) :
Saccone, N.L., et al. Genes Brain Behav. 9(7):741-750(2010)
Ruano, G., et al. Pharmacogenomics 11(7):959-971(2010)
Jin, Y., et al. Int. J. Immunogenet. (2010) In press :
Schraufstatter, I.U., et al. J Stem Cells 4(4):203-215(2009)