bs-1038R

[Primary Antibody]

CHRNA4 Rabbit pAb



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- DATASHEET		400-901-9800
Host: Rabbit	Isotype: IgG	Applications: WB (1:500-2000) IHC-P (1:100-500)
Clonality: Polyclonal GenelD: 1137	SWISS: P43681	IHC-F (1:100-500) IF (1:100-500) ELISA (1:5000-10000)
Target: CHRNA4 Immunogen: KLH conjugated synthetic peptide derived from human CHRNA4: 151-250/627. < Cytoplasmic >		Reactivity: Human, Mouse, Rat
Purification: affinity purified b	y Protein A	
Concentration: 1mg/ml Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.		Predicted MW.: ⁶⁷ kDa Subcellular Location: ^{Cell} membrane
belongs to a supe role in fast signal receptors can bin change in confor conducting chan an integral memb either nAChR beta Mutations in this Polymorphisms in	es a nicotinic acetylcholine receptor, which erfamily of ligand-gated ion channels that play a transmission at synapses. These pentameric d acetylcholine, which causes an extensive mation that leads to the opening of an ion- nel across the plasma membrane. This protein is brane receptor subunit that can interact with a-2 or nAChR beta-4 to form a functional recepto gene cause nocturnal frontal lobe epilepsy type n this gene that provide protection against n have been described.	r.



25 ug total protein per lane of various lysates (see on figure) probed with CHRNA4 polyclonal antibody, unconjugated (bs-1038R) at 1:1000 dilution and 4°C overnight incubation. Followed by conjugated secondary antibody incubation at r.t. for 60 min.



Tissue/cell: rat brain tissue; 4% Paraformaldehyde-fixed and paraffinembedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum,C-0005) at 37°C for 20 min; Incubation: Anti-CHRNA4 Polyclonal Antibody, Unconjugated(bs-1038R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody(SP-0023) and DAB(C-0010) staining

- SELECTED CITATIONS -

• [IF=4.367] Sallam MY et al. Brainstem cholinergic pathways diminish cardiovascular and neuroinflammatory actions of endotoxemia in rats: Role of NFκB/α7/α4β2AChRs signaling. Neuropharmacology. 2019 Jun 25;157:107683. IHC ;Rat. 31247270