

GluR-2 (Phospho-Tyr876) rabbit pAb

Cat No.: ES16165

For research use only

Overview

Product Name GluR-2 (Phospho-Tyr876) rabbit pAb

Host species Rabbit
Applications IHC;IF;WB

Species Cross-Reactivity Human; Mouse; Rat

Recommended dilutions IHC-p 1:50-200, WB 1:500-2000

Immunogen Synthesized peptide derived from human GluR-2

(Phospho-Tyr876)

Specificity This antibody detects endogenous phospho levels of

GluR-2 (Phospho-Tyr876) at Human:Y876,

Mouse:Y876, Rat:Y876

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Storage Store at -20°C. Avoid repeated freeze-thaw cycles.

Protein Name GluR-2 (Phospho-Tyr876)

Gene Name GRIA2 GLUR2

Cellular localization Cell membrane; Multi-pass membrane protein.

Endoplasmic reticulum membrane ; Multi-pass membrane protein . Cell junction, synapse,

postsynaptic cell membrane; Multi-pass membrane protein. Cell junction, synapse, postsynaptic density

membrane; Multi-p

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

Clonality Polyclonal
Concentration 1 mg/ml
Observed band 99kD
Human Gene ID 2891
Human Swiss-Prot Number P42262

Alternative Names Glutamate receptor 2 (GluR-2;AMPA-selective

glutamate receptor 2; GluR-B; GluR-K2; Glutamate

receptor ionotropic, AMPA 2;GluA2)

Background Glutamate receptors are the predominant excitatory

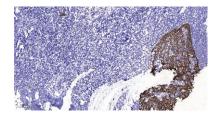


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neurotransmitter receptors in the mammalian brain and are activated in a variety of normal neurophysiologic processes. This gene product belongs to a family of glutamate receptors that are sensitive to

alpha-amino-3-hydroxy-5-methyl-4-isoxazole propionate (AMPA), and function as ligand-activated cation channels. These channels are assembled from 4 related subunits, GRIA1-4. The subunit encoded by this gene (GRIA2) is subject to RNA editing (CAG->CGG; Q->R) within the second transmembrane domain, which is thought to render the channel impermeable to Ca(2+). Human and animal studies suggest that pre-mRNA editing is essential for brain function, and defective GRIA2 RNA editing at the Q/R site may be relevant to amyotrophic lateral sclerosis (ALS) etiology. Alternative splicing, resulting in transcript variants enco



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

