

LGI3 Antibody

Catalog # ASC10653

Product Information

Application WB, E
Primary Accession Q8N145

Other AccessionAAQ88483, 37181338ReactivityHuman, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Isotype IgG
Calculated MW 61704
Concentration (mg/ml) 1 mg/mL
Conjugate Unconjugated

Application NotesLGI3 antibody can be used for the detection of LGI3 by Western blot at 1

□g/mL.

Additional Information

Gene ID 203190

Other Names Leucine-rich repeat LGI family member 3, LGI1-like protein 4, Leucine-rich

glioma-inactivated protein 3, LGI3, LGIL4

Target/Specificity LGI3; Two isoforms of LGI3 are known to exist; this LGI3 antibody will

recognize both. This LGI3 antibody is predicted to be specific to LGI3 and not recognize other LGI proteins. The observed higher molecular weight band

may represent a post-translationally modified form of LGI3.

Reconstitution & Storage LGI3 antibody can be stored at 4°C for three months and -20°C, stable for up

to one year. As with all antibodies care should be taken to avoid repeated freeze thaw cycles. Antibodies should not be exposed to prolonged high

temperatures.

Precautions LGI3 Antibody is for research use only and not for use in diagnostic or

therapeutic procedures.

Protein Information

Name LGI3

Synonyms LGIL4

Function May participate in the regulation of neuronal exocytosis.

Cellular Location Secreted. Cytoplasmic vesicle, secretory vesicle, synaptic vesicle

{ECO:0000250|UniProtKB:Q8K406}. Synapse, synaptosome {ECO:0000250|UniProtKB:Q8K406}. Cell projection, axon

{ECO:0000250|UniProtKB:Q8K406}. Note=Found in the synaptosomal membrane fraction. Within peripheral myelinated axons, LGI3 is highly expressed at the juxtaparanodal membrane and colocalizes with the voltage-gated potassium channels Kv1.1 (KCNA1) and Kv1.2 (KCNA2), and with CNTNAP2, DLG4, ADAM22 and ADAM23 (By similarity) {ECO:0000250|UniProtKB:Q8K406}

Tissue Location

Widely expressed, with highest levels in brain and lung.

Background

LGI3 Antibody: The leucine-rich, glioma inactivated gene 3 (LGI3) is a member of the LGI family in which LGI1 is the exemplar. The LGI family consists of four of highly related proteins containing leucine-rich repeats (LRRs) which are highly similar to other transmembrane signaling molecules and receptors. LGI1 has been identified as a candidate tumor suppressor gene for glioma and plays a role in autodominant lateral temporal epilepsy (ADTLE), an epileptic syndrome characterized by focal seizures with predominant auditory symptoms. Despite its high homology with LGI1 and similar pattern of expression, mutations in LGI3 have not been found to be associated with ADTLE. LGI3 expression is induced in rat astrocyte cultures by the amyloid beta (Abeta) peptide and accumulated on neuronal plasma membranes of aged monkey brains and co-localized with Abeta.

References

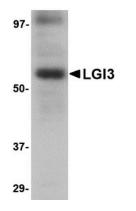
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Berkovic SF, Izzillo P, McMahon JM, et al. LGI1 mutations in temporal lobe epilepsies. Neurology2004; 62:1115-9.

Kimura N, Ishii Y, Suzaki S, et al. Abeta upregulates and colocalizes with LGI3 in cultured rat astrocytes. Cell Mol. Neurobiol.2007; 27:335-50.

Images



Western blot analysis of LGI3 in human brain tissue lysate with LGI3 antibody at 1 µg/mL.

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.