**Delivery** 

**Background** 



## **PRODUCT INFORMATION**

Tag C-Flag Tag

GRIA2 **Target** 

GLUR2, GLURB, GluA2, GluR-K2, HBGR2, NEDLIB, **Synonyms** 

gluR-2, gluR-B

Human GRIA2 full length protein-synthetic Description

nanodisc 6~8weeks P42262

**Uniprot ID HFK293 Expression Host** 

**Protein Families** Ion Channels: Glutamate Receptors

**Protein Pathways** 

The human full length GRIA2 protein has a MW of **Molecular Weight** 

98.8kDa

Lyophilized from nanodisc solubilization buffer (20 mM Tris-HCl, 150 mM NaCl, pH 8.0). Normally 5% Formulation & - 8% trehalose is added as protectants before Reconstitution lyophilization. Please see Certificate of Analysis

for specific instructions of reconstitution. Store at -20°C to -80°C for 12 months in lyophilized form. After reconstitution, if not

intended for use within a month, aliquot and store Storage & Shipping at -80°C (Avoid repeated freezing and thawing). Lyophilized proteins are shipped at ambient

temperature.

Glutamate receptors are the predominant excitatory 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 premRNA 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 encoding different isoforms, (including the flip and flop isoforms that vary in their signal transduction properties), has been noted for this gene. [provided by RefSeq, Jul

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