

Anti-GABBR2 (pS893) Antibody

Rabbit polyclonal antibody to GABBR2 (pS893) Catalog # AP61281

Product Information

Application WB
Primary Accession O75899
Other Accession O80T41

Reactivity Human, Mouse, Rat

Host Rabbit
Clonality Polyclonal
Calculated MW 105821

Additional Information

Gene ID 9568

Other Names GPR51; GPRC3B; Gamma-aminobutyric acid type B receptor subunit 2;

GABA-B receptor 2; GABA-B-R2; GABA-BR2; GABABR2; Gb2; G-protein coupled

receptor 51; HG20

Target/Specificity KLH-conjugated synthetic peptide encompassing a sequence within the

C-term region of human GABBR2 (pS893). The exact sequence is proprietary.

Dilution WB~~WB (1/500 - 1/1000)

Format Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30%

glycerol, and 0.09% (W/V) sodium azide.

Storage Store at -20 °C.Stable for 12 months from date of receipt

Protein Information

Name GABBR2

Synonyms GPR51, GPRC3B

Function Component of a heterodimeric G-protein coupled receptor for GABA,

formed by GABBR1 and GABBR2 (PubMed: 15617512, PubMed: 18165688, PubMed: 22660477, PubMed: 24305054, PubMed: 9872316, PubMed: 9872744). Within the heterodimeric GABA receptor, only GABBR1 seems to bind

agonists, while GABBR2 mediates coupling to G proteins (PubMed:18165688). Ligand binding causes a conformation change that triggers signaling via guanine nucleotide-binding proteins (G proteins) and modulates the activity of down-stream effectors, such as adenylate cyclase (PubMed:10075644, PubMed:10773016, PubMed:24305054). Signaling inhibits adenylate cyclase, stimulates phospholipase A2, activates potassium channels, inactivates

voltage-dependent calcium-channels and modulates inositol phospholipid hydrolysis (PubMed:10075644, PubMed:10773016, PubMed:10906333, PubMed:9872744). Plays a critical role in the fine-tuning of inhibitory synaptic transmission (PubMed:22660477, PubMed:9872744). Pre-synaptic GABA receptor inhibits neurotransmitter release by down-regulating high-voltage activated calcium channels, whereas postsynaptic GABA receptor decreases neuronal excitability by activating a prominent inwardly rectifying potassium (Kir) conductance that underlies the late inhibitory postsynaptic potentials (PubMed:10075644, PubMed:22660477, PubMed:9872316, PubMed:9872744). Not only implicated in synaptic inhibition but also in hippocampal long-term potentiation, slow wave sleep, muscle relaxation and antinociception (Probable).

Cellular Location

Cell membrane; Multi-pass membrane protein. Postsynaptic cell membrane {ECO:0000250 | UniProtKB:O88871}; Multi-pass membrane protein. Note=Coexpression of GABBR1 and GABBR2 is required for GABBR1 maturation and transport to the plasma membrane. In contrast, GABBR2 does not depend on GABBR1 for transport to the cell membrane

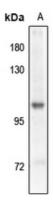
Tissue Location

Highly expressed in brain, especially in cerebral cortex, thalamus, hippocampus, frontal, occipital and temporal lobe, occipital pole and cerebellum, followed by corpus callosum, caudate nucleus, spinal cord, amygdala and medulla (PubMed:10087195, PubMed:10328880, PubMed:10727622, PubMed:9872744). Weakly expressed in heart, testis and skeletal muscle (PubMed:10087195, PubMed:10727622)

Background

KLH-conjugated synthetic peptide encompassing a sequence within the C-term region of human GABBR2 (pS893). The exact sequence is proprietary.

Images



Western blot analysis of GABBR2 (pS893) expression in U87MG (A) whole cell lysates.

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