

Product datasheet for TL312861

OriGene Technologies, Inc.

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GABA A Receptor gamma 2 (GABRG2) Human shRNA Plasmid Kit (Locus ID 2566)

Product data:

Product Type: shRNA Plasmids

Product Name: GABA A Receptor gamma 2 (GABRG2) Human shRNA Plasmid Kit (Locus ID 2566)

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Synonyms: CAE2; DEE74; ECA2; EIEE74; FEB8; GEFSP3

Vector: pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Puromycin

Selection:

Format: Lentiviral plasmids

Components: GABRG2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 2566).

5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pGFP-C-shLenti Vector, TR30021, included for free.

RefSeq: NM 000816, NM 198903, NM 198904, NM 198904.1, NM 198904.2, NM 198903.1,

NM 198903.2, NM 000816.1, NM 000816.2, NM 000816.3, BC059389, BC059389.1, BC069348,

BC069348.1, BC036030, BC052926, BC074795, NM 198904.4

UniProt ID: P18507

Summary: This gene encodes a gamma-aminobutyric acid (GABA) receptor. GABA is the major inhibitory

neurotransmitter in the mammlian brain, where it acts at GABA-A receptors, which are ligand-

gated chloride channels. GABA-A receptors are pentameric, consisting of proteins from

several subunit classes: alpha, beta, gamma, delta and rho. Mutations in this gene have been

associated with epilepsy and febrile seizures. Multiple transcript variants encoding different

shRNA Design: These shRNA constructs were designed against multiple splice variants at this gene locus. To

isoforms have been identified for this gene. [provided by RefSeq, Jul 2008]

be certain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u>.

If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.





Performance Guaranteed:

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).