

## **Product datasheet for TL311749**

## Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com technic original company of the company

**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

EU: info-de@origene.com CN: techsupport@origene.cn

## LGI1 Human shRNA Plasmid Kit (Locus ID 9211)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** LGI1 Human shRNA Plasmid Kit (Locus ID 9211)

**Locus ID:** 9211

Synonyms: ADLTE; ADPAEF; ADPEAF; EPITEMPIN; EPT; ETL1; IB1099

Vector: pGFP-C-shLenti (TR30023)

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

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: LGI1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 9211). 5µg

purified plasmid DNA per construct

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

RefSeq: NM 001308275, NM 001308276, NM 005097, NR 131777, NM 005097.1, NM 005097.2,

NM 005097.3, BC022500, BC022500.1

UniProt ID: 095970

**Summary:** This gene encodes a member of the secreted leucine-rich repeat (LRR) superfamily and

shares homology with members of the SLIT protein family. The encoded protein may regulate

the activity of voltage-gated potassium channels and may be involved in neuronal growth

regulation and cell survival. This gene is rearranged as a result of translocations in glioblastoma cell lines, and it is frequently down-regulated or rearranged in malignant gliomas. Mutations in this gene result in autosomal dominant lateral temporal epilepsy. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Apr 2015]

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

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 custom shRNA service.





## 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).