

Product datasheet for TL301244

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

TAS1R1 Human shRNA Plasmid Kit (Locus ID 80835)

Product data:

Product Type: shRNA Plasmids

Product Name: TAS1R1 Human shRNA Plasmid Kit (Locus ID 80835)

Chloramphenicol (34 ug/ml)

Locus ID:

Synonyms: GM148; GPR70; T1R1; TR1

Vector: pGFP-C-shLenti (TR30023) E. coli Selection:

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

TAS1R1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 80835). Components:

5µg purified plasmid DNA per construct

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

NM 138697, NM 177539, NM 177540, NM 177541, NM 177540.1, NM 177540.2, RefSeq:

NM 177541.1, NM 138697.1, NM 138697.2, NM 138697.3, NM 177539.1, BC136515,

BC136516, NM 177540.3, NM 138697.4

UniProt ID: Q7RTX1

Summary: The protein encoded by this gene is a G protein-coupled receptor and is a component of the

> heterodimeric amino acid taste receptor T1R1+3. The T1R1+3 receptor responds to L-amino acids but not to D-enantiomers or other compounds. Most amino acids that are perceived as

sweet activate T1R1+3, and this activation is strictly dependent on an intact T1R1+3

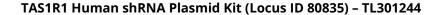
heterodimer. Multiple transcript variants encoding different isoforms have been found for

this gene. [provided by RefSeq, Jun 2010]

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

be certain that your variant of interest is targeted, please contact techsupport@origene.com. 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).