

Product datasheet for TR512660

Taar1 Mouse shRNA Plasmid (Locus ID 111174)

Product data:

Product Type: shRNA Plasmids

Product Name: Taar1 Mouse shRNA Plasmid (Locus ID 111174)

Locus ID: 111174

Synonyms: taR-1; Tar1; Trar1

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycir

Selection:

Puromycin

Format: Retroviral plasmids

Components: Taar1 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

111174). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: <u>BC125368</u>, <u>BC125370</u>, <u>NM 053205</u>, <u>NM 053205.1</u>

UniProt ID: Q923Y8

Summary: Receptor for trace amines, including beta-phenylethylamine (b-PEA), p-tyramine (p-TYR),

octopamine and tryptamine, with highest affinity for b-PEA and p-TYR. Unresponsive to classical biogenic amines, such as epinephrine and histamine and only partially activated by dopamine and serotonin. Trace amines are biogenic amines present in very low levels in

mammalian tissues. Although some trace amines have clearly defined roles as neurotransmitters in invertebrates, the extent to which they function as true

neurotransmitters in vertebrates has remained speculative. Trace amines are likely to be involved in a variety of physiological functions that have yet to be fully understood. The signal

transduced by this receptor is mediated by the G(s)-class of G-proteins which activate

adenylate cyclase.[UniProtKB/Swiss-Prot Function]

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.



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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).