

Product datasheet for TR311161

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NMBR Human shRNA Plasmid Kit (Locus ID 4829)

Product data:

Product Type: shRNA Plasmids

Product Name: NMBR Human shRNA Plasmid Kit (Locus ID 4829)

Locus ID: 4829

Synonyms: BB1; BB1R; BRS1; NMB-R

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Puromycin

Format: Retroviral plasmids

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

4829). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001324307, NM 001324308, NM 002511, NM 002511.1, NM 002511.2, NM 002511.3,

BC095542, NM 002511.4

UniProt ID: P28336

Summary: This gene encodes a 7-transmembrane G protein-coupled receptor that binds neuromedin B,

which is a growth factor and mitogen for gastrointestinal epithelial tissue and for normal and

neoplastic lung. This receptor may play a role in smooth muscle contraction, neuronal responses, and the regulation of cell growth. Antagonists of this receptor have a potential therapeutic use in inhibiting tumor cell growth. Polymorphisms in this gene may be

associated with a susceptibility for schizophrenia. Alternative splicing of this gene results in

multiple transcript variants. [provided by RefSeq, Apr 2016]

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 <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).