EMPOWER YOUR RESEARCH

## Product datasheet for TR509434

## Rpn2 Mouse shRNA Plasmid (Locus ID 20014)

## Product data:

Product Type:
Product Name:
Locus ID:
Synonyms:
Vector:
E. coli Selection:

Mammalian Cell
Selection:
Format:
Components:

RefSeq:

UniProt ID:
Summary:
shRNA Design:

shRNA Plasmids

Rpn2 Mouse shRNA Plasmid (Locus ID 20014)
20014
1300012C06Rik; AV261018; Rpn-2
pRS (TR20003)
Ampicillin
Puromycin

Retroviral plasmids
Rpn2 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID = 20014). $5 \mu \mathrm{~g}$ purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.
BC046806, NM 019642, NM 001355163 , NM 001355164 NM 001355165 NM 019642.1 NM 019642.2, NM 019642.3 NM 019642.4 BC010509

## Q9DBG6

Subunit of the oligosaccharyl transferase (OST) complex that catalyzes the initial transfer of a defined glycan ( $\operatorname{Glc}(3) \mathrm{Man}(9) \mathrm{GlcNAc}(2)$ in eukaryotes) from the lipid carrier dolicholpyrophosphate to an asparagine residue within an Asn-X-Ser/Thr consensus motif in nascent polypeptide chains, the first step in protein N-glycosylation. N -glycosylation occurs cotranslationally and the complex associates with the Sec61 complex at the channel-forming translocon complex that mediates protein translocation across the endoplasmic reticulum (ER). All subunits are required for a maximal enzyme activity.[UniProtKB/Swiss-Prot Function]
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 techsupport@origene.com. If you need a special design or shRNA sequence, please utilize our custom shRNA service.

## Performance <br> 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).

