

Product datasheet for TL504467

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

Ube2v2 Mouse shRNA Plasmid (Locus ID 70620)

Product data:

Product Type: shRNA Plasmids

Product Name: Ube2v2 Mouse shRNA Plasmid (Locus ID 70620)

Locus ID:

1110021H13Rik; 4632410D19Rik; 5730524P06Rik; 6820402M05; Al848315; C81524; MMS2 Synonyms:

Vector: pGFP-C-shLenti (TR30023) E. coli Selection: Chloramphenicol (34 ug/ml)

Mammalian Cell

Selection:

Puromycin

Format:

Lentiviral plasmids

Components: Ube2v2 - Mouse, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 70620).

5µg purified plasmid DNA per construct

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

BC058374, BC083098, BC094903, NM 001159351, NM 023585, NM 023585.1, NM 023585.2, RefSeq:

NM 023585.3, NM 023585.4, NM 001159351.1, BC029742, BC044722

UniProt ID: Q9D2M8

Summary: Has no ubiquitin ligase activity on its own. The UBE2V2/UBE2N heterodimer catalyzes the

> synthesis of non-canonical poly-ubiquitin chains that are linked through 'Lys-63'. This type of poly-ubiquitination does not lead to protein degradation by the proteasome. Mediates transcriptional activation of target genes. Plays a role in the control of progress through the cell cycle and differentiation. Plays a role in the error-free DNA repair pathway and

contributes to the survival of cells after DNA damage.[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.



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).