

Product datasheet for TL513689V

OriGene Technologies, Inc.

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Rbm45 Mouse shRNA Lentiviral Particle (Locus ID 241490)

Product data:

Product Type: shRNA Lentiviral Particles

Product Name: Rbm45 Mouse shRNA Lentiviral Particle (Locus ID 241490)

Locus ID: 241490

Synonyms: Drb1; Drbp1; G430095G15Rik

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: Rbm45 - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble

control), 0.5 ml each, >10^7 TU/ml.

RefSeq: <u>BC057890</u>, <u>NM 153405</u>, <u>NM 153405.1</u>, <u>NM 153405.2</u>, <u>NM 178090</u>

UniProt ID: Q8BHN5

Summary: RNA-binding protein with binding specificity for poly(C). May play an important role in neural

development (By similarity).[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 techsupport@origene.com.

OriGene guarantees that the sequences in the shRNA expression cassettes are verified to

If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u>.

Performance

Guaranteed: 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).

