

Product datasheet for TL500097V

OriGene Technologies, Inc.

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Anxa11 Mouse shRNA Lentiviral Particle (Locus ID 11744)

Product data:

Product Type: shRNA Lentiviral Particles

Product Name: Anxa11 Mouse shRNA Lentiviral Particle (Locus ID 11744)

Locus ID: 11744

Synonyms: A830099017Rik; Anx11

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: Anxa11 - Mouse shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble

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

RefSeq: <u>BC012875</u>, <u>NM 013469</u>, <u>NM 013469.1</u>, <u>NM 013469.2</u>

UniProt ID: P97384

Summary: Required for midbody formation and completion of the terminal phase of cytokinesis (By

similarity). Binds specifically to calcyclin in a calcium-dependent manner. [UniProtKB/Swiss-

Prot Function]

shRNA Design: These shRNA constructs were designed against multiple splice variants at this gene locus. To

be used in comparison with the target-specific shRNA transfected samples.

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

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).

