

Product datasheet for TL711618V

OriGene Technologies, Inc.

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Flot2 Rat shRNA Lentiviral Particle (Locus ID 83764)

Product data:

Product Type: shRNA Lentiviral Particles

Product Name: Flot2 Rat shRNA Lentiviral Particle (Locus ID 83764)

Locus ID: 83764

Vector: pGFP-C-shLenti (TR30023)

Format: Lentiviral particles

Components: Flot2 - Rat shRNA lentiviral particles (4 unique 29mer target-specific shRNA, 1 scramble

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

RefSeq: NM 001270800, NM 001270801, NM 031830, NR 073076, NR 073077, NM 031830.1,

NM 031830.2, NM 001270801.1, NM 001270800.1, BC083550

UniProt ID: Q9Z2S9

Summary: colocalized with activated GPI-linked cell adhesion molecules at the plasma membrane where

transmembrane signaling may occur [RGD, Feb 2006]

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.

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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).