

Product datasheet for TL302433

OriGene Technologies, Inc.

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FAPP2 (PLEKHA8) Human shRNA Plasmid Kit (Locus ID 84725)

Product data:

Product Type: shRNA Plasmids

Product Name: FAPP2 (PLEKHA8) Human shRNA Plasmid Kit (Locus ID 84725)

Locus ID: 84725 Synonyms: FAPP2

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: PLEKHA8 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID =

84725). 5µg purified plasmid DNA per construct

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

RefSeq: NM 001197026, NM 001197027, NM 032639, NM 001350973, NM 001350974,

NM 001350975, NM 032639.1, NM 032639.2, NM 032639.3, NM 001197027.1,

NM 001197026.1, BC002838, BC002838.2, BC053990, NM 001363473, NM 001363474,

NM 001197026.2, NM 032639.4

UniProt ID: Q96|A3

Summary: Cargo transport protein that is required for apical transport from the Golgi complex.

Transports AQP2 from the trans-Golgi network (TGN) to sites of AQP2 phosphorylation. Mediates the non-vesicular transport of glucosylceramide (GlcCer) from the trans-Golgi network (TGN) to the plasma membrane and plays a pivotal role in the synthesis of complex glycosphingolipids. Binding of both phosphatidylinositol 4-phosphate (PIP) and ARF1 are essential for the GlcCer transfer ability. Also required for primary cilium formation, possibly by being involved in the transport of raft lipids to the apical membrane, and for membrane

tubulation.[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 <u>custom shRNA service</u>.







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).