

## **Product datasheet for TL311472**

## OriGene Technologies, Inc.

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## Aquaporin 0 (MIP) Human shRNA Plasmid Kit (Locus ID 4284)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** Aquaporin 0 (MIP) Human shRNA Plasmid Kit (Locus ID 4284)

**Locus ID:** 4284

Synonyms: AQP0; CTRCT15; LIM1; MIP26; MP26

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

**Mammalian Cell** 

Puromycin

Selection:

Format: Lentiviral plasmids

Components: MIP - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 4284). 5µg

purified plasmid DNA per construct

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

RefSeq: NM 012064, NM 012064.1, NM 012064.2, NM 012064.3, BC117474, BC074913, BM696300,

BM705396, BM716579, BM724221, NM 012064.4

UniProt ID: P30301

Summary: Major intrinsic protein is a member of the water-transporting aquaporins as well as the

original member of the MIP family of channel proteins. The function of the fiber cell

membrane protein encoded by this gene is undetermined, yet this protein is speculated to play a role in intracellular communication. The MIP protein is expressed in the ocular lens and is required for correct lens function. This gene has been mapped among aquaporins AQP2, AQP5, and AQP6, in a potential gene cluster at 12q13. [provided by RefSeq, Jul 2008]

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).