

## **Product datasheet for TL311276**

## OriGene Technologies, Inc.

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## NAB2 Human shRNA Plasmid Kit (Locus ID 4665)

**Product data:** 

**Product Type:** shRNA Plasmids

Product Name: NAB2 Human shRNA Plasmid Kit (Locus ID 4665)

**Locus ID:** 4665

Synonyms: MADER

**Vector:** pGFP-C-shLenti (TR30023)

E. coli Selection: Chloramphenicol (34 ug/ml)

**Mammalian Cell** 

Selection:

Puromycin

Format: Lentiviral plasmids

Components: NAB2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 4665).

5µg purified plasmid DNA per construct

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

RefSeq: NM 005967, NM 001330305, NM 005967.1, NM 005967.2, NM 005967.3, BC065931,

BC007756, NM 005967.4

UniProt ID: 015742

**Summary:** This gene encodes a member of the family of NGFI-A binding (NAB) proteins, which function

in the nucleus to repress transcription induced by some members of the EGR (early growth response) family of transactivators. NAB proteins can homo- or hetero-multimerize with

other EGR or NAB proteins through a conserved N-terminal domain, and repress

transcription through two partially redundant C-terminal domains. Transcriptional repression by the encoded protein is mediated in part by interactions with the nucleosome remodeling and deactylase (NuRD) complex. Alternatively spliced transcript variants have been described,

but their biological validity has not been determined. [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).