

## **Product datasheet for TL306768**

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# **ALAS1 Human shRNA Plasmid Kit (Locus ID 211)**

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** ALAS1 Human shRNA Plasmid Kit (Locus ID 211)

Locus ID: 21

Synonyms: ALAS; ALAS-H; ALAS3; ALASH; MIG4

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

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

**Mammalian Cell** 

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Selection: Format:

Lentiviral plasmids

Puromycin

**Components:** ALAS1 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 211).

5µg purified plasmid DNA per construct

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

RefSeq: NM 000688, NM 001304443, NM 001304444, NM 199166, NM 199166.1, NM 199166.2,

NM 000688.1, NM 000688.2, NM 000688.3, NM 000688.4, NM 000688.5, BC011798,

BC011798.2, BM546974, NM 000688.6

UniProt ID: P13196

**Summary:** This gene encodes the mitochondrial enzyme which is catalyzes the rate-limiting step in heme

(iron-protoporphyrin) biosynthesis. The enzyme encoded by this gene is the housekeeping enzyme; a separate gene encodes a form of the enzyme that is specific for erythroid tissue. The level of the mature encoded protein is regulated by heme: high levels of heme down-

regulate the mature enzyme in mitochondria while low heme levels up-regulate. A

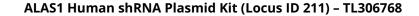
pseudogene of this gene is located on chromosome 12. Alternative splicing results in multiple

transcript variants encoding different isoforms. [provided by RefSeq, Jan 2015]

**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).