

Product datasheet for TL311707

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LMO2 Human shRNA Plasmid Kit (Locus ID 4005)

Product data:

Product Type: shRNA Plasmids

Product Name: LMO2 Human shRNA Plasmid Kit (Locus ID 4005)

Locus ID: 4005

Synonyms: LMO-2; RBTN2; RBTNL1; RHOM2; TTG2

Vector: pGFP-C-shLenti (TR30023)

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

Mammalian Cell

Selection:

Puromycin

Format: Lentiviral plasmids

Components: LMO2 - Human, 4 unique 29mer shRNA constructs in lentiviral GFP vector(Gene ID = 4005).

5µg purified plasmid DNA per construct

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

RefSeq: NM 001142315, NM 001142316, NM 005574, NM 005574.1, NM 005574.2, NM 005574.3,

NM 001142315.1, BC034041, BC034041.1, BC035607, BC042426, BC073973, BM690599,

BM699014, NM 005574.4

UniProt ID: P25791

Summary: LMO2 encodes a cysteine-rich, two LIM-domain protein that is required for yolk sac

erythropoiesis. The LMO2 protein has a central and crucial role in hematopoietic development and is highly conserved. The LMO2 transcription start site is located approximately 25 kb downstream from the 11p13 T-cell translocation cluster (11p13 ttc), where a number T-cell acute lymphoblastic leukemia-specific translocations occur. Alternative splicing results in multiple transcript variants encoding different isoforms.

[provided by RefSeq, Nov 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).