

Product datasheet for TR512098

Arntl2 Mouse shRNA Plasmid (Locus ID 272322)

Product data:

Product Type: shRNA Plasmids

Product Name: Arntl2 Mouse shRNA Plasmid (Locus ID 272322)

Locus ID: 272322

Synonyms: 4632430A05Rik; bHLHe6; BMAL2; CLIF; MOP9

Vector: pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: Arntl2 - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

272322). 5µg purified plasmid DNA per construct

29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.

RefSeq: BC108965, NM 001289679, NM 001289680, NM 001289681, NM 172309, NM 172309.1,

NM 172309.2, NM 001289681.1, NM 001289679.1, NM 001289680.1, BC108966

UniProt ID: Q2VPD4

Summary: This gene encodes a basic helix-loop-helix transcription factor belonging to the PAS (Per, Arnt,

Sim) superfamily. The PAS proteins play important roles in adaptation to low atmospheric and cellular oxygen levels, exposure to certain environmental pollutants, and diurnal oscillations in light and temperature. This protein forms a transcriptionally active

heterodimer with the circadian Clock protein, the structurally related Mop4, and hypoxia-inducible factors, such as Hif1alpha. Consistent with its role as a biologically relevant partner of circadian and hypoxia factors, this protein is coexpressed in regions of the brain such as the thalamus, hypothalamus, and amygdala. Alternatively spliced transcript variants encoding

different isoforms have been described for this gene. [provided by RefSeq, Feb 2014]

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 custom shRNA service.



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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).