

## **Product datasheet for TR314692**

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## ARG2 Human shRNA Plasmid Kit (Locus ID 384)

**Product data:** 

**Product Type:** shRNA Plasmids

**Product Name:** ARG2 Human shRNA Plasmid Kit (Locus ID 384)

Locus ID: 384

**Vector:** pRS (TR20003)

E. coli Selection: Ampicillin

Mammalian Cell Puromycin

Selection:

Format: Retroviral plasmids

Components: ARG2 - Human, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =

384). 5µg purified plasmid DNA per construct

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

**RefSeq:** NM 001172, NM 001172.1, NM 001172.2, NM 001172.3, BC029050, BC029050.1, BC001350,

BC008464, NM 001172.4

UniProt ID: <u>P78540</u>

**Summary:** Arginase catalyzes the hydrolysis of arginine to ornithine and urea. At least two isoforms of

mammalian arginase exists (types I and II) which differ in their tissue distribution, subcellular

localization, immunologic crossreactivity and physiologic function. The type II isoform

encoded by this gene, is located in the mitochondria and expressed in extra-hepatic tissues, especially kidney. The physiologic role of this isoform is poorly understood; it is thought to play a role in nitric oxide and polyamine metabolism. Transcript variants of the type II gene resulting from the use of alternative polyadenylation sites have been described. [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>.



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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).