

## Product datasheet for **SC205656**

### Arginase 1 (ARG1) (NM\_000045) Human 3' UTR Clone

#### Product data:

**Product Type:** 3' UTR Clones  
**Product Name:** Arginase 1 (ARG1) (NM\_000045) Human 3' UTR Clone  
**Vector:** pMirTarget (PS100062)  
**Symbol:** ARG1  
**ACCN:** NM\_000045  
**Insert Size:** 451 bp  
**Insert Sequence:** >SC205656 3'UTR clone of NM\_000045

The sequence shown below is from the reference sequence of NM\_000045. The complete sequence of this clone may contain minor differences, such as SNPs.  
**Blue**=Stop Codon **Red**=Cloning site

```
GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAAGCCAAGAAGGGCGGAAAGATCGCCGTG
TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC
CCTATTGACTACCTTAACCCACCTAAGTAATGTGGAAACATCCGATATAAATCTCATAGTTAATGGCA
TAATTAGAAAGCTAATCATTTTTCTAAGCATAGAGTTATCCTTCTAAAGACTTGTCTTTTCAGAAAAAT
GTTTTTCCAATTAGTATAAACTCTACAAATCCCTCTTGGTGTAATAAATTCAAGATGTGGAAATTCTAAC
TTTTTTGAAATTTAAAAGCTTATTTTTCTAACTTGGCAAAAGACTTATCCTTAGAAAGAGAAGTGAC
ATTGATTTCCAATTAATAATTTGCTGGCATTAAAAATAAGCACACTTACATAAGCCCCCATAACATAGAG
TGGGACTCTTGGAAATCAGGAGACAAAGCTACCACATGTGGAAAGGTAATGTGTCCATGTCATTCAAA
AAATGTGATTTTTTATAATAAACTCTTTATAACAAGA
ACGCGTAAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA
CGAGATTTGATTCCACCGCCCTTCTATGAAAGG
```

**Restriction Sites:** SgfI-MluI

**OTI Disclaimer:** Our molecular clone sequence data has been matched to the sequence identifier above as a point of reference. Note that the complete sequence of this clone is largely the same as the reference sequence but may contain minor differences, e.g., single nucleotide polymorphisms (SNPs).

**Components:** The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The package also includes 100 pmols of both the corresponding 5' and 3' vector primers in separate vials.

**RefSeq:** [NM\\_000045.4](#)



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**Summary:** Arginase catalyzes the hydrolysis of arginine to ornithine and urea. At least two isoforms of mammalian arginase exist (types I and II) which differ in their tissue distribution, subcellular localization, immunologic crossreactivity and physiologic function. The type I isoform encoded by this gene, is a cytosolic enzyme and expressed predominantly in the liver as a component of the urea cycle. Inherited deficiency of this enzyme results in argininemia, an autosomal recessive disorder characterized by hyperammonemia. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2011]

**Locus ID:** 383

**MW:** 17.8