

## **Product datasheet for SC203379**

## ASS1 (NM 054012) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

**Product Name:** ASS1 (NM\_054012) Human 3' UTR Clone

Symbol: ASS1

Synonyms: ASS; CTLN1

Mammalian Cell N

Selection:

Neomycin

**Vector:** pMirTarget (PS100062)

**ACCN:** NM\_054012

**Insert Size:** 283 bp

Insert Sequence: >SC203379 3'UTR clone of NM\_054012

The sequence shown below is from the reference sequence of NM\_054012. The complete

sequence of this clone may contain minor differences, such as SNPs.

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CGTCTCCAGAGCAAGGTCACTGCCAAATAGACCCGTGTACAATGAGGAGCTGGGGCCTCCTCAATTTGCAGATCCCCCAAGTACAGGCGCTAATTGTTGTAATTTGTAATTGTGACTTGTTCTCCCCGGCTGGCAGCGTAGTGGGGCTGCCAGCCTGCAAGCCTGCAAACGTTGTCATCGAAGGGAAGGGTGGGGGGCAGCTGCGGTGGGGAGCTATAAAAATGACAATTAAAAGAGACACTAGTCTTTT

ATTTCTA

CGAGATTTCGATTCCACCGCCGCCTTCTATGAAAGG

Restriction Sites: Sgfl-Mlul

**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: <u>NM 054012.4</u>



**OriGene Technologies, Inc.** 9620 Medical Center Drive, Ste 200

CN: techsupport@origene.cn

Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com



## ASS1 (NM\_054012) Human 3' UTR Clone - SC203379

**Summary:** The protein encoded by this gene catalyzes the penultimate step of the arginine biosynthetic

pathway. There are approximately 10 to 14 copies of this gene including the pseudogenes scattered across the human genome, among which the one located on chromosome 9 appears to be the only functional gene for argininosuccinate synthetase. Mutations in the chromosome 9 copy of this gene cause citrullinemia. Two transcript variants encoding the

same protein have been found for this gene. [provided by RefSeq, Aug 2012]

Locus ID: 445

MW: 10.4