

## Product datasheet for SC202837

## MCCC1 (NM 020166) Human 3' UTR Clone

**Product data:** 

**Product Type:** 3' UTR Clones

**Product Name:** MCCC1 (NM\_020166) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

MCCC1 Symbol:

Synonyms: MCC-B; MCCA ACCN: NM 020166

Insert Size: 250 bp

>SC202837 3'UTR clone of NM\_020166 **Insert Sequence:** 

The sequence shown below is from the reference sequence of NM\_020166. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

GAAGAATCAGACAAAAGGGAATCGGAATAAACTCCAGCAAGGAAATGGCCAGTTAAGTAGTGTCTTCTC TCTCCACCAAAAAGAGGAAGTGCCTCCAGCTTTTCTGGGGGTCTCATAAAGAGCAGTTTTACTAAATGA TTGTATGCTTATGCTGAACACCTTTCATATTGGAGAATCATGCATTTGGGTCACTAATTATCTCAAAAT

ATTTCATACTAATAAAGTTGAATTATTTTTTATTGGAAGCCAA

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).

The cDNA clone is shipped in a 2-D bar-coded Matrix tube as 10 ug dried plasmid DNA. The Components:

package also includes 100 pmols of both the corresponding 5' and 3' vector primers in

separate vials.

NM 020166.5 RefSeq:



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



## MCCC1 (NM\_020166) Human 3' UTR Clone - SC202837

Summary: This gene encodes the large subunit of 3-methylcrotonyl-CoA carboxylase. This enzyme

functions as a heterodimer and catalyzes the carboxylation of 3-methylcrotonyl-CoA to form

3-methylglutaconyl-CoA. Mutations in this gene are associated with 3-

Methylcrotonylglycinuria, an autosomal recessive disorder of leucine catabolism. [provided

by RefSeq, Jul 2008]

**Locus ID:** 56922

**MW:** 9.6