

## **Product datasheet for SC201594**

## COMP (NM 000095) Human 3' UTR Clone

## Product data:

**Product Type:** 3' UTR Clones

**Product Name:** COMP (NM\_000095) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: COMP

Synonyms: CTS2; EDM1; EPD1; MED; PSACH; THBS5; TSP5

**ACCN:** NM\_000095

**Insert Size:** 172 bp

Insert Sequence: >SC201594 3'UTR clone of NM\_000095

The sequence shown below is from the reference sequence of NM\_000095. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

TATGAGACCCATCAGCTGCGGCAAGCCTAGGGACCAGGGTGAGGACCCGCCGGATGACAGCCACCCTCACCCGGGCTGGATGGGGGGCTCTGCACCCCAGCCCCAAGGGGTGGCCGTCCTGAGGGGGAAGTGAGAAGGGC

TCAGAGAGGACAAAATAAAGTGTGTGCAGGGA

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.

**RefSeg:** NM 000095.3



**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



## COMP (NM\_000095) Human 3' UTR Clone - SC201594

Summary: The protein encoded by this gene is a noncollagenous extracellular matrix (ECM) protein. It

consists of five identical glycoprotein subunits, each with EGF-like and calcium-binding (thrombospondin-like) domains. Oligomerization results from formation of a five-stranded coiled coil and disulfides. Binding to other ECM proteins such as collagen appears to depend on divalent cations. Contraction or expansion of a 5 aa aspartate repeat and other mutations can cause pseudochondroplasia (PSACH) and multiple epiphyseal dysplasia (MED). [provided

by RefSeq, Jul 2016]

**Locus ID:** 1311

**MW:** 5.8