

Product datasheet for SC207215

OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

Treacher Collins syndrome protein (TCOF1) (NM_001008656) Human 3' UTR Clone

Product data:

Product Type: 3' UTR Clones

Product Name: Treacher Collins syndrome protein (TCOF1) (NM_001008656) Human 3' UTR Clone

Vector: pMirTarget (PS100062)

Symbol: TCOF1

Synonyms: MFD1; nucleolar trafficking phosphoprotein; Treacher Collins-Franceschetti syndrome 1;

Treacher Collins syndrome protein; treacle

ACCN: NM 001008656

Insert Size: 568 bp

Insert Sequence: >SC207215 3'UTR clone of NM_001008656

The sequence shown below is from the reference sequence of NM_001008656. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

AAAAAAAAAAAA

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).





MW:

Treacher Collins syndrome protein (TCOF1) (NM_001008656) Human 3' UTR Clone - SC207215

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.

21.1

RefSeq: <u>NM 001008656.2</u>

Summary: This gene encodes a nucleolar protein with a LIS1 homology domain. The protein is involved

in ribosomal DNA gene transcription through its interaction with upstream binding factor (UBF). Mutations in this gene have been associated with Treacher Collins syndrome, a disorder which includes abnormal craniofacial development. Multiple transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Sep 2008]

Locus ID: 6949