

## **Product datasheet for SC210930**

## TMPRSS3 (NM 024022) Human 3' UTR Clone

## **Product data:**

**Product Type:** 3' UTR Clones

Product Name: TMPRSS3 (NM\_024022) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: TMPRSS3

Synonyms: DFNB8; DFNB10; ECHOS1; TADG12

**ACCN:** NM\_024022

**Insert Size:** 902 bp

Insert Sequence: >SC210930 3'UTR clone of NM\_024022

The sequence shown below is from the reference sequence of NM\_024022. The complete

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

Blue=Stop Codon Red=Cloning site

GGCAAGTTGGACGCCCGCAAGATCCGCGAGATTCTCATTAAGGCCAAGAAGGGCGGAAAGATCGCCGTG

TAACAATTGGCAGAGCTCAGAATTCAAGCGATCGCC

CCTCA

**ACGCGT**AAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCGACCAAGCGACGCCCAACCTGCCATCA

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).



**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



## TMPRSS3 (NM\_024022) Human 3' UTR Clone - SC210930

**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 024022.4</u>

Summary: This gene encodes a protein that belongs to the serine protease family. The encoded protein

contains a serine protease domain, a transmembrane domain, an LDL receptor-like domain, and a scavenger receptor cysteine-rich domain. Serine proteases are known to be involved in a variety of biological processes, whose malfunction often leads to human diseases and disorders. This gene was identified by its association with both congenital and childhood onset autosomal recessive deafness. This gene is expressed in fetal cochlea and many other tissues, and is thought to be involved in the development and maintenance of the inner ear or the contents of the perilymph and endolymph. This gene was also identified as a tumor-associated gene that is overexpressed in ovarian tumors. Alternatively spliced transcript

variants have been described. [provided by RefSeq, Jan 2012]

**Locus ID:** 64699

MW: 33.5