

Product datasheet for RC204771L4

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OriGene Technologies, Inc.

TIMM8A (NM_004085) Human Tagged Lenti ORF Clone

Product data:

Product Type: Expression Plasmids

Product Name: TIMM8A (NM_004085) Human Tagged Lenti ORF Clone

Tag: mGFP

Symbol: TIMM8A

Synonyms: DDP; DDP1; DFN1; MTS; TIM8

Mammalian Cell

Selection:

Vector: pLenti-C-mGFP-P2A-Puro (PS100093)

Puromycin

E. coli Selection: Chloramphenicol (34 ug/mL)

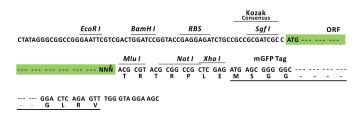
ORF Nucleotide The ORF insert of this clone is exactly the same as(RC204771).

Sequence:

Restriction Sites: Sgfl-Mlul

Cloning Scheme:





^{*} The last codon before the Stop codon of the ORF

ACCN: NM_004085

ORF Size: 291 bp





TIMM8A (NM_004085) Human Tagged Lenti ORF Clone - RC204771L4

OTI Disclaimer: The molecular sequence of this clone aligns with the gene accession number as a point of

reference only. However, individual transcript sequences of the same gene can differ through naturally occurring variations (e.g. polymorphisms), each with its own valid existence. This clone is substantially in agreement with the reference, but a complete review of all prevailing

variants is recommended prior to use. More info

OTI Annotation: This clone was engineered to express the complete ORF with an expression tag. Expression

varies depending on the nature of the gene.

Components: The ORF clone is ion-exchange column purified and shipped in a 2D barcoded Matrix tube

containing 10ug of transfection-ready, dried plasmid DNA (reconstitute with 100 ul of water).

Reconstitution Method: 1. Centrifuge at 5,000xg for 5min.

2. Carefully open the tube and add 100ul of sterile water to dissolve the DNA.

3. Close the tube and incubate for 10 minutes at room temperature.

4. Briefly vortex the tube and then do a quick spin (less than 5000xg) to concentrate the liquid

at the bottom.

5. Store the suspended plasmid at -20°C. The DNA is stable for at least one year from date of

shipping when stored at -20°C.

RefSeq: <u>NM 004085.2</u>

 RefSeq Size:
 1459 bp

 RefSeq ORF:
 294 bp

 Locus ID:
 1678

 UniProt ID:
 060220

Cytogenetics: Xq22.1

Protein Families: Druggable Genome

MW: 11 kDa

Gene Summary: This translocase is involved in the import and insertion of hydrophobic membrane proteins

from the cytoplasm into the mitochondrial inner membrane. The gene is mutated in Mohr-Tranebjaerg syndrome/Deafness Dystonia Syndrome (MTS/DDS) and it is postulated that MTS/DDS is a mitochondrial disease caused by a defective mitochondrial protein import

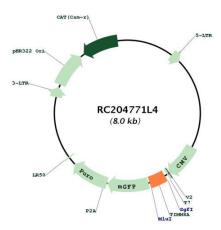
system. Defects in this gene also cause Jensen syndrome; an X-linked disease with

opticoacoustic nerve atrophy and muscle weakness. This protein, along with TIMM13, forms a 70 kDa heterohexamer. Alternative splicing results in multiple transcript variants encoding

distinct isoforms.[provided by RefSeq, Mar 2009]



Product images:



Circular map for RC204771L4