

# **Product datasheet for SC210302**

## TJP2 (NM 004817) Human 3' UTR Clone

### **Product data:**

**Product Type:** 3' UTR Clones

Product Name: TJP2 (NM\_004817) Human 3' UTR Clone

**Vector:** pMirTarget (PS100062)

Symbol: TJP2

Synonyms: C9DUPq21.11; DFNA51; DUP9q21.11; FHCA1; PFIC4; X104; ZO2

**ACCN:** NM\_004817

**Insert Size:** 865 bp

Insert Sequence: >SC210302 3' UTR clone of NM\_004817

The sequence shown below is from the reference sequence of NM\_004817. The complete sequence of this clone may contain minor differences, such as SNPs. Red=Cloning site

Blue=Stop Codon

#### CAATTGGCAGAGCTCAGAATTCAAGCGATCGC

ACGCGTAAGCGGCCGCGCATCTAGATTCGAAGAAAATGACCG

**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).



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## TJP2 (NM\_004817) Human 3' UTR Clone - SC210302

**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 004817.3</u>

Summary: This gene encodes a zonula occluden that is a member of the membrane-associated

guanylate kinase homolog family. The encoded protein functions as a component of the tight junction barrier in epithelial and endothelial cells and is necessary for proper assembly of tight junctions. Mutations in this gene have been identified in patients with hypercholanemia, and genomic duplication of a 270 kb region including this gene causes autosomal dominant deafness-51. Alternatively spliced transcripts encoding multiple isoforms have been observed

for this gene. [provided by RefSeq, Nov 2011]

**Locus ID:** 9414