

Product datasheet for SC311650

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TJP2 (AK025185) Human Untagged Clone

Product data:

Symbol:

Product Type: Expression Plasmids

Product Name: TJP2 (AK025185) Human Untagged Clone

Tag: Tag Free TJP2

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

Vector: pCMV6 series

>NCBI ORF sequence for AK025185, the custom clone sequence may differ by one or more **Fully Sequenced ORF:**

nucleotides

Restriction Sites: Please inquire

ACCN: AK025185

OTI Disclaimer: Our molecular clone sequence data has been matched to the reference identifier above as a

> point of reference. Note that the complete sequence of our molecular clones may differ from the sequence published for this corresponding reference, e.g., by representing an alternative

RNA splicing form or single nucleotide polymorphism (SNP).

OTI Annotation: This TrueClone is provided through our Custom Cloning Process that includes sub-cloning

> into OriGene's pCMV6 vector and full sequencing to provide a non-variant match to the expected reference without frameshifts, and is delivered as lyophilized plasmid DNA.

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

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: AK025185.1

RefSeq Size: 1811 bp





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RefSeq ORF: 1811 bp Locus ID: 9414 Cytogenetics: 9q21.11

Protein Pathways: Tight junction, Vibrio cholerae infection

Gene 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]