

Product datasheet for RC200490L3V

OriGene Technologies, Inc.

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Claudin 4 (CLDN4) (NM 001305) Human Tagged ORF Clone Lentiviral Particle

Product data:

Product Type: Lentiviral Particles

Product Name: Claudin 4 (CLDN4) (NM_001305) Human Tagged ORF Clone Lentiviral Particle

Symbol: CLDN4

Synonyms: CPE-R; CPETR; CPETR1; hCPE-R; WBSCR8

Mammalian Cell

Selection:

Puromycin

Vector: pLenti-C-Myc-DDK-P2A-Puro (PS100092)

Tag: Myc-DDK
ACCN: NM 001305

ORF Size: 627 bp

ORF Nucleotide

The ORF insert of this clone is exactly the same as(RC200490).

Sequence:

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.

RefSeg: NM 001305.3

 RefSeq Size:
 1859 bp

 RefSeq ORF:
 630 bp

 Locus ID:
 1364

 UniProt ID:
 014493

 Cytogenetics:
 7q11.23

Domains: PMP22_Claudin

Protein Families: Druggable Genome, Transmembrane





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Protein Pathways: Cell adhesion molecules (CAMs), Leukocyte transendothelial migration, Tight junction

MW: 22.1 kDa

Gene Summary: The protein encoded by this intronless gene belongs to the claudin family. Claudins are

integral membrane proteins that are components of the epithelial cell tight junctions, which regulate movement of solutes and ions through the paracellular space. This protein is a high-affinity receptor for Clostridium perfringens enterotoxin (CPE) and may play a role in internal organ development and function during pre- and postnatal life. This gene is deleted in Williams-Beuren syndrome, a neurodevelopmental disorder affecting multiple systems.

[provided by RefSeq, Sep 2013]