

Product datasheet for TP316887L

OriGene Technologies, Inc.

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CLDN19 (NM_148960) Human Recombinant Protein

Product data:

Product Type: Recombinant Proteins

Description: Recombinant protein of human claudin 19 (CLDN19), transcript variant 1, 1 mg

Species: Human
Expression Host: HEK293T

Expression cDNA Clone >Peptide sequence encoded by RC216887 or AA Sequence: Blue=ORF Red=Cloning site Green=Tag(s)

MANSGLQLLGYFLALGGWVGIIASTALPQWKQSSYAGDAIITAVGLYEGLWMSCASQSTGQVQCKLYDS LLALDGHIQSARALMVVAVLLGFVAMVLSVVGMKCTRVGDSNPIAKGRVAIAGGALFILAGLCTLTAVS WYATLVTQEFFNPSTPVNARYEFGPALFVGWASAGLAVLGGSFLCCTCPEPERPNSSPQPYRPGPSAAA

REPVVKLPASAKGPLGV

TRTRPLEQKLISEEDLAANDILDYKDDDDKV

Recombinant protein using RC216887 also available, TP316887

Tag: C-Myc/DDK

Predicted MW: 23 kDa

Concentration: >0.05 μg/μL as determined by microplate BCA method

Purity: > 80% as determined by SDS-PAGE and Coomassie blue staining

Buffer: 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol

Preparation: Recombinant protein was captured through anti-DDK affinity column followed by

conventional chromatography steps.

Note: For testing in cell culture applications, please filter before use. Note that you may experience

some loss of protein during the filtration process.

Storage: Store at -80°C.

Stability: Stable for 12 months from the date of receipt of the product under proper storage and

handling conditions. Avoid repeated freeze-thaw cycles.

RefSeq: NP 683763

Locus ID: 149461



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UniProt ID: Q8N6F1

RefSeq Size: 2859

Cytogenetics: 1p34.2

RefSeq ORF: 672

Synonyms: HOMG5

Summary: The product of this gene belongs to the claudin family. It plays a major role in tight junction-

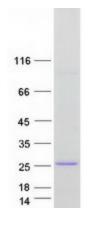
specific obliteration of the intercellular space, through calcium-independent cell-adhesion activity. Defects in this gene are the cause of hypomagnesemia renal with ocular involvement (HOMGO). HOMGO is a progressive renal disease characterized by primary renal magnesium wasting with hypomagnesemia, hypercalciuria and nephrocalcinosis associated with severe ocular abnormalities such as bilateral chorioretinal scars, macular colobomata, significant myopia and nystagmus. Alternatively spliced transcript variants encoding distinct isoforms

have been identified for this gene. [provided by RefSeq, Jun 2010]

Protein Families: Transmembrane

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

Product images:



Coomassie blue staining of purified CLDN19 protein (Cat# [TP316887]). The protein was produced from HEK293T cells transfected with CLDN19 cDNA clone (Cat# [RC216887]) using MegaTran 2.0 (Cat# [TT210002]).

Wegarran 2.0 (Cat# [17210002]).