

# Product datasheet for TP316887M

## CLDN19 (NM\_148960) Human Recombinant Protein

### **Product data:**

#### **Product Type: Recombinant Proteins** Recombinant protein of human claudin 19 (CLDN19), transcript variant 1, 100 µg **Description:** Species: Human HEK293T **Expression Host:** Expression cDNA Clone >RC216887 representing NM\_148960 or AA Sequence: Red=Cloning site Green=Tags(s) MANSGLQLLGYFLALGGWVGIIASTALPQWKQSSYAGDAIITAVGPYEGLWMSCASQSTGQVQCKLYDSL LALDGHIQSARALMVVAVLLGFVAMVLSVVGMKCTRVGDSNPIAKGRVAIAGGALFILAGLCTLTAVSWY ATLVTQEFFNPSTPVNARYEFGPALFVGWASAGLAVLGGSFLCCTCPEPERPNSSPQPYRPGPSAAAREP VVKLPASAKGPLGV **TRTRPLEQKLISEEDLAANDILDYKDDDDKV** Tag: C-Myc/DDK Predicted MW: 23 kDa **Concentration:** >0.05 µg/µL as determined by microplate BCA method > 80% as determined by SDS-PAGE and Coomassie blue staining Purity: **Buffer:** 25 mM Tris-HCl, 100 mM glycine, pH 7.3, 10% glycerol Recombinant protein was captured through anti-DDK affinity column followed by **Preparation:** 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. Store at -80°C. Storage: Stable for 12 months from the date of receipt of the product under proper storage and Stability: handling conditions. Avoid repeated freeze-thaw cycles. **RefSeq:** NP 683763 Locus ID: 149461 **UniProt ID:** Q8N6F1



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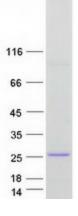
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### OriGene Technologies, Inc.

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	CLDN19 (NM_148960) Human Recombinant Protein – TP316887M
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 image	es:

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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]).

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