

## **Product datasheet for TP325251**

## OriGene Technologies, Inc.

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## CLDN19 (NM\_001123395) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Recombinant protein of human claudin 19 (CLDN19), transcript variant 2, 20 μg

Species: Human
Expression Host: HEK293T

**Expression cDNA Clone** >RC225251 representing NM\_001123395

or AA Sequence: Red=Cloning site Green=Tags(s)

MANSGLQLLGYFLALGGWVGIIASTALPQWKQSSYAGDAIITAVGLYEGLWMSCASQSTGQVQCKLYDSL LALDGHIQSARALMVVAVLLGFVAMVLSVVGMKCTRVGDSNPIAKGRVAIAGGALFILAGLCTLTAVSWY ATLVTQEFFNPSTPVNARYEFGPALFVGWASAGLAVLGGSFLCCTCPEPERPNSSPQPYRPGPSAAAREY

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**TRTRPLEQKLISEEDLAANDILDYKDDDDKV** 

Tag: C-Myc/DDK
Predicted MW: 21.9 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 001116867

 Locus ID:
 149461

 UniProt ID:
 Q8N6F1





Cytogenetics: 1p34.2

RefSeq ORF: 633

Synonyms: HOMG5

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

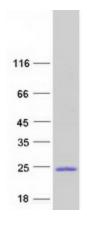
specific obliteration of the intercellular space, through calcium-independent cell-adhesion activity. Defects in this gene are the cause of hypomagnesemia renal with ocular involvement (HOMGO). HOMGO is a progressive renal disease characterized by primary renal magnesium wasting with hypomagnesemia, 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# TP325251). The protein was produced from HEK293T cells transfected with CLDN19 cDNA clone (Cat# [RC225251]) using MegaTran 2.0 (Cat# [TT210002]).