

## Product Information

### MemDX™ Membrane Protein Human CLDN19 (Claudin 19) Expressed *in vitro* *E.coli* expression system, Full Length

Cat. No.: **MPX2232K**

This product is for research use only and is not intended for diagnostic use.

This product is a Human CLDN19 membrane protein expressed *in vitro* *E.coli* expression system. The protein is for research use only and is not approved for use in humans or in clinical diagnosis.

#### Product Specifications

##### Host Species

Human

##### Target Protein

CLDN19

##### Protein Length

Full Length

##### Protein Class

Receptor

##### TMD

4

##### Sequence

MANSGLQLLGYFLALGGWVGIIASTALPQWKQSSYAGDAIITAVGLYEGLWMSCASQSTGQVQCKLYDSLLALDGHIQSARALMVV

#### Product Description

##### Expression Systems

*in vitro* *E.coli* expression system

##### Tag

10xHis tag at the N-terminus

##### Protein Format

Soluble

##### Form

Liquid or Lyophilized powder

##### Buffer

Tris/PBS-based buffer, 6% Trehalose, pH 8.0

### **Storage**

Aliquot and store at -20°C or lower. For long term storage, we recommend to store at -70°C or lower. Avoid freeze/thaw cycles.

### **Target**

#### **Target Protein**

CLDN19

#### **Full Name**

Claudin 19

### **Introduction**

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.

#### **Alternative Names**

CLDN19; HOMG5; claudin-19; Claudin 19

#### **Gene ID**

[149461](#)

#### **UniProt ID**

[Q8N6F1](#)