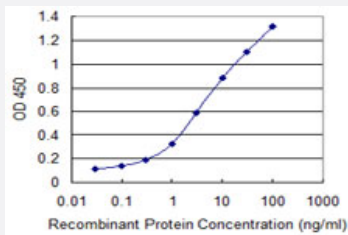


CLDN19 monoclonal antibody (M02), clone 2F2

Catalog # H00149461-M02

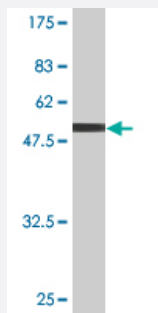
Size 100 ug

Applications



Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged CLDN19 is 0.1 ng/ml as a capture antibody.



Western Blot detection against Immunogen (48.95 KDa) .

Specification

Product Description

Mouse monoclonal antibody raised against a full-length recombinant CLDN19.

Immunogen

CLDN19 (AAH30524, 1 a.a. ~ 211 a.a) full-length recombinant protein with GST tag. MW of the GST tag alone is 26 KDa.

Sequence

MANSGLQLLG YFLALGGWVGIIASTALPQWKQSSYAGDAITAVGLYEGLWMSCASQSTGQVQCK
LYDSLLALDGHISARALMVAVLLGFVAMVLSVVGKCTRVGDSNPIAKGRVAIAGGALFILAGL
CTLTAVSWYATLVTQEFPNPSTPVNARYEFGPALFVGWASAGLAVLGGSFLLCCTCPEPERPNSS
PQPYRPGPSAAAREYV

Host

Mouse

Reactivity

Human

Isotype	IgG2a Kappa
Quality Control Testing	Antibody Reactive Against Recombinant Protein. Western Blot detection against Immunogen (48.95 KDa) .
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

Applications

- Western Blot (Recombinant protein)

[Protocol Download](#)

- Sandwich ELISA (Recombinant protein)

Detection limit for recombinant GST tagged CLDN19 is 0.1 ng/ml as a capture antibody.

[Protocol Download](#)

- ELISA

Gene Info — CLDN19

Entrez GeneID	149461
GeneBank Accession#	BC030524
Protein Accession#	AAH30524
Gene Name	CLDN19
Gene Alias	-
Gene Description	claudin 19
Omim ID	248190 610036
Gene Ontology	Hyperlink

Gene 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. Two transcript variants encoding distinct isoforms have been identified for this gene. [provided by RefSeq]

Other Designations

OTTHUMP00000008733

Pathway

- [Cell adhesion molecules \(CAMs\)](#)
- [Leukocyte transendothelial migration](#)
- [Tight junction](#)