

## Datasheet

### CLDN19 monoclonal antibody (M02), clone 2F2

**Catalog Number:** H00149461-M02

**Regulation Status:** For research use only (RUO)

**Product Description:** Mouse monoclonal antibody raised against a full-length recombinant CLDN19.

**Clone Name:** 2F2

**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:**

MANSGQLQLGYFLALGGWVGIIASTALPQWKQSSYAG  
DAITAVGLYEGLWMSCASQSTGQVQCKLYDSLALD  
GHIQSARALMVVAVLLGFVAMVLSVVGMKCTRVGDSN  
PIAKGRVAIAGGALFILAGLCTLTAVSWYATLVTQEFFN  
PSTPVNARYEFGPALFVGWASAGLAVLGGSFCLCTCP  
EPERPNSSPQPYPYRPGPSAAAREYV

**Host:** Mouse

**Reactivity:** Human

**Applications:** ELISA, S-ELISA, WB-Re

(See our web site product page for detailed applications information)

**Protocols:** See our web site at

<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

**Isotype:** IgG2a Kappa

**Storage Buffer:** In 1x PBS, pH 7.4

**Storage Instruction:** Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.

**Entrez GeneID:** 149461

**Gene Symbol:** CLDN19

**Gene Alias:** -

**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]