

CLDN1 rabbit monoclonal antibody

Catalog # H00009076-K Size 100 ug x up to 3

Specification	
Product Description	Rabbit monoclonal antibody raised against a human CLDN1 peptide using ARM Technology.
Immunogen	A synthetic peptide of human CLDN1 is used for rabbit immunization. Customer or Abnova will decide on the preferred peptide sequence.
Host	Rabbit
Library Construction	Non-fusion antibody library from rabbit spleen (ARM Technology).
Expression	Overexpression vector and transfection into 293H cell line.
Reactivity	Human
Purification	Protein A
Isotype	lgG
Quality Control Testing	Antibody reactive against human CLDN1 peptide by ELISA and mammalian transfected lysate by W estern Blot.
Storage Buffer	In 1x PBS, pH 7.4
Storage Instruction	Store at -20°C or lower. Aliquot to avoid repeated freezing and thawing.
Deliverable	Up to three rabbit IgG clones of 100 ug each will be delivered to customer.
Note	 Customer may provide cell or tissue lysate for antibody screening. Rabbit monoclonal antibody generated by ARM technology is amenable to antibody engineering in cluding F(ab)₂, lgG, scFv and different Fc and non-Fc conjugates per customer request.

Applications

Western Blot (Transfected lysate)

Protocol Download



ELISA

Gene Info — CLDN1	
Entrez GenelD	9076
GeneBank Accession#	CLDN1
Gene Name	CLDN1
Gene Alias	CLD1, ILVASC, SEMP1
Gene Description	claudin 1
Omim ID	<u>603718</u> <u>607626</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	Tight junctions represent one mode of cell-to-cell adhesion in epithelial or endothelial cell sheets, f orming continuous seals around cells and serving as a physical barrier to prevent solutes and wat er from passing freely through the paracellular space. These junctions are comprised of sets of continuous networking strands in the outwardly facing cytoplasmic leaflet, with complementary groov es in the inwardly facing extracytoplasmic leaflet. The protein encoded by this gene, a member of the claudin family, is an integral membrane protein and a component of tight junction strands. Los s of function mutations result in neonatal ichthyosis-sclerosing cholangitis syndrome. [provided by RefSeq
Other Designations	senescence-associated epithelial membrane protein 1

Pathway

- Cell adhesion molecules (CAMs)
- Leukocyte transendothelial migration
- Pathogenic Escherichia coli infection EHEC
- Tight junction

Disease

- Genetic Predisposition to Disease
- Hepatitis C



- Substance Abuse
- Tobacco Use Disorder