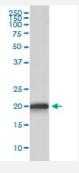


CLDN5 monoclonal antibody, clone AAHA-3

Catalog # MAB19903 Size 100 uL

Applications



Western Blot (Cell lysate)

Western blot analysis of human fetal brain lysate with CLDN5 monoclonal antibody.

Specification	
Product Description	Rabbit monoclonal antibody raised against synthetic peptide of human CLDN5.
Immunogen	A synthetic peptide corresponding to human CLDN5.
Host	Rabbit
Reactivity	Human
Form	Liquid
Purification	Affinity purification
Isotype	lgG
Recommend Usage	Immunohistochemistry (1:50-1:200) Western Blot (1:500-1:2000) The optimal working dilution should be determined by the end user.
Storage Buffer	In PBS, pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Storage Instruction	Store at -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and st ored frozen at -20°C for a longer time. Avoid repeated freezing and thawing.



Product Information

Note

This product contains sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only.

Applications

Western Blot (Cell lysate)

Western blot analysis of human fetal brain lysate with CLDN5 monoclonal antibody.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections)

Gene Info — CLDN5	
Entrez GenelD	<u>7122</u>
Protein Accession#	<u>000501</u>
Gene Name	CLDN5
Gene Alias	AWAL, BEC1, CPETRL1, TMVCF
Gene Description	claudin 5
Omim ID	602101
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a member of the claudin family. Claudins are integral membrane proteins and components of tight junction strands. Tight junction strands serve as a physical barrier to prevent s olutes and water from passing freely through the paracellular space between epithelial or endothel ial cell sheets. Mutations in this gene have been found in patients with velocardiofacial syndrome. Alternatively spliced transcript variants encoding the same protein have been found for this gene. [provided by RefSeq
Other Designations	androgen withdrawal and apoptosis induced protein RVP1-like transmembrane protein deleted in velocardiofacial syndrome

Pathway

- Cell adhesion molecules (CAMs)
- Leukocyte transendothelial migration
- Tight junction



Disease

- Chromosome Deletion
- Schizophrenia