

Claudin 4 (ABT188R) rabbit mAb

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| Catalog No : | YM7088 |
| Reactivity : | Human; |
| Applications : | WB;IHC; ELISA |
| Target : | Claudin-4 |
| Fields : | >>Cell adhesion molecules;>>Tight junction;>>Leukocyte transendothelial migration;>>Pathogenic Escherichia coli infection;>>Hepatitis C |
| Gene Name : | CLDN4 |
| Protein Name : | Claudin-4 (Clostridium perfringens enterotoxin receptor) (CPE-R) (CPE-receptor) (Williams-Beuren syndrome chromosomal region 8 protein) |
| Human Gene Id : | 1364 |
| Human Swiss Prot No : | O14493 |
| Immunogen : | Synthesized peptide derived from human Claudin 4 AA range:150-209 |
| Specificity : | This antibody detects endogenous levels of Claudin-4 |
| Formulation : | PBS, 50% glycerol, 0.05% Proclin 300, 0.05%BSA |
| Source : | Monoclonal, Rabbit IgG1, Kappa |
| Dilution : | IHC 1:100-500, WB 1:500-1000, ELISA 1:5000-20000 |
| Purification : | Recombinant Expression and Affinity purified |
| Storage Stability : | -15°C to -25°C/1 year(Do not lower than -25°C) |
| Molecularweight : | 22kD |
| Background : | The protein encoded by this intronless gene belongs to the claudin family. Claudins are integral membrane proteins that are components of the epithelial cell |

tight junctions, which regulate movement of solutes and ions through the paracellular space. This protein is a high-affinity receptor for Clostridium perfringens enterotoxin (CPE) and may play a role in internal organ development and function during pre- and postnatal life. This gene is deleted in Williams-Beuren syndrome, a neurodevelopmental disorder affecting multiple systems. [provided by RefSeq, Sep 2013],

Function :

disease:Haploinsufficiency of CLDN4 may be the cause of certain cardiovascular and musculo-skeletal abnormalities observed in Williams-Beuren syndrome (WBS), a rare developmental disorder. It is a contiguous gene deletion syndrome involving genes from chromosome band 7q11.23.,function:Plays a major role in tight junction-specific obliteration of the intercellular space.,similarity:Belongs to the claudin family.,subunit:Directly interacts with TJP1/ZO-1, TJP2/ZO-2 and TJP3/ZO-3.,

Subcellular Location :

Cell junction, tight junction . Cell membrane ; Multi-pass membrane protein . CLDN4 is required for tight junction localization in the kidney. .

Expression :

Colon,Fetal brain,Trachea,

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