

# Product Name: Claudin-3 (phospho Tyr219) Rabbit Polyclonal Antibody Catalog #: APRab04471

For research use only.

#### **Summary**

**Description** Rabbit polyclonal Antibody

**Host** Rabbit

ApplicationWB,IHC,ICC/IF,ELISAReactivityHuman,Rat,MouseConjugationUnconjugatedModificationPhosphorylated

**Isotype** IgG

ClonalityPolyclonalFormLiquidConcentration1mg/ml

**Storage** Aliquot and store at -20°C (valid for 12 months). Avoid freeze/thaw cycles.

**Shipping** Ice bags

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% New type **Buffer** 

preservative N.

**Purification** Affinity purification

#### **Application**

**Dilution Ratio** WB 1:500-1:2000,IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:5000-1:10000

Molecular Weight 28kDa

### **Antigen Information**

Gene Name CLDN3

CLDN3; C7orf1; CPETR2; Claudin-3; Clostridium perfringens enterotoxin receptor 2; CPE-R 2;

Alternative Names

CPE-receptor 2; Rat ventral prostate.1 protein homolog; hRVP1

**Gene ID** 1365.0 **SwissProt ID** O15551

The antiserum was produced against synthesized peptide derived from human Claudin 3 Immunogen

around the phosphorylation site of Tyr219. AA range:171-220

# **Background**

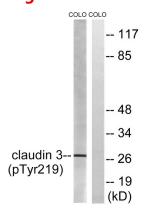


Tight junctions represent one mode of cell-to-cell adhesion in epithelial or endothelial cell sheets, forming continuous seals around cells and serving as a physical barrier to prevent solutes and water from passing freely through the paracellular space. These junctions are comprised of sets of continuous networking strands in the outwardly facing cytoplasmic leaflet, with complementary grooves in the inwardly facing extracytoplasmic leaflet. The protein encoded by this intronless gene, a member of the claudin family, is an integral membrane protein and a component of tight junction strands. It is also a low-affinity receptor for Clostridium perfringens enterotoxin, and shares as sequence similarity with a putative apoptosis-related protein found in rat. [provided by RefSeq, Jul 2008], disease:Haploinsufficiency of CLDN3 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, through calcium-independent cell-adhesion activity., similarity:Belongs to the claudin family., subunit:Can form homo- and heteropolymers with other CLDN. Homopolymers interact with CLDN1 and CLDN2 homopolymers. Directly interacts with TJP1/ZO-1, TJP2/ZO-2 and TJP3/ZO-3.,

#### **Research Area**

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

# **Image Data**



Western blot analysis of lysates from COLO205 cells treated with EGF 200ng/ml 30 ', using Claudin 3 (Phospho-Tyr219) Antibody. The lane on the right is blocked with the phospho peptide.

Web: https://www.enkilife.com E-mail: order@enkilife.com techsupport@enkilife.com Tel: 0086-27-87002838