

## Claudin-4 (Phospho Tyr208) Rabbit pAb

CatalogNo: YP1088

### Key Features

**Host Species**

- Rabbit

**Reactivity**

- Human,Rat,Mouse,

**Applications**

- IHC,IF,ELISA

**MW**

- 22kD (Calculated)

**Isotype**

- IgG

### Recommended Dilution Ratios

**IHC 1:100-1:300****ELISA 1:5000****IF 1:50-200**

### Storage

**Storage\***

-15°C to -25°C/1 year(Do not lower than -25°C)

**Formulation**

Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

### Basic Information

**Clonality**

Polyclonal

### Immunogen Information

**Immunogen**

The antiserum was produced against synthesized peptide derived from human Claudin 4 around the phosphorylation site of Tyr208. AA range:160-209

**Specificity**

Phospho-Claudin-4 (Y208) Polyclonal Antibody detects endogenous levels of Claudin-4 protein only when phosphorylated at Y208. The name of modified sites may be influenced by many factors, such as species (the modified site was not originally found in human samples) and the change of protein sequence (the previous protein sequence is incomplete, and the protein sequence may be prolonged with the development of protein sequencing technology). When naming, we will use the "numbers" in historical reference to keep the sites consistent with the reports. The antibody binds to the following modification sequence (lowercase letters are modification sites):ASNyV

## | Target Information

**Gene name** CLDN4

**Protein Name** Claudin-4

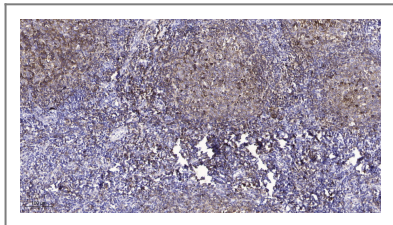
Organism	Gene ID	UniProt ID
Human	<a href="#">1364;</a>	<a href="#">O14493;</a>
Mouse		<a href="#">O35054;</a>

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

**Tissue specificity** Colon,Fetal brain,Trachea,

**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.,

## | Validation Data



Immunohistochemical analysis of paraffin-embedded human tonsil. 1, Antibody was diluted at 1:200(4° overnight). 2, Tris-EDTA,pH9.0 was used for antigen retrieval. 3,Secondary antibody was diluted at 1:200(room temperature, 45min).

## | Contact information

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