#### **Product Datasheet**

# Claudin 5 (Phospho-Tyr217) Antibody

Catalog No: #11792

Package Size: #11792-1 50ul #11792-2 100ul



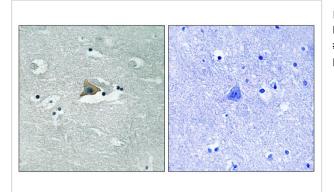
Orders: order@signalwayantibody.com Support: tech@signalwayantibody.com

Host Species Rabbit Clonality Polyclonal Purification Antibodies were produced by immunizing rabbits with synthetic phosphopeptide and KLH conjugates. Antibodies were purified by affinity-chromatography using epitope-specific phosphopeptide. Non-phospho specific antibodies were removed by chromatogramphy using non-phosphopeptide. Non-phospho specific antibodies were removed by chromatogramphy using non-phosphopeptide.  Applications WB IHC IF  Species Reactivity Hu Ms Rt  Specificity The antibody detects endogenous levels of Claudin 5 only when phosphorylated at tyrosine 217.  Immunogen Type Peptide-KLH  Immunogen Description Peptide sequence around phosphorylation site of tyrosine 217 (K-K-N-Y(p)-V) derived from Human Claudin 5.  Target Name Claudin 5  Modification Phospho  Other Names CLD5; CLDN5; TMDVCF;  Accession No. Swiss-Prot#: 000501; NCBI Gene#: 7122; NCBI Protein#: NP_001124333.1.  Uniprot 000501  GeneID 7122;  SDS-PAGE MW 23kd  Concentration 1.0mg/ml  Formulation Rabbit IgG in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.	Product Name	Claudin 5 (Phospho-Tyr217) Antibody				
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and 50% glycerol.	Formulation	Rabbit IgG in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl, 0.02% sodium azide				
		and 50% glycerol.				

## Application Details

WB 1:500-2000			
IHC 1:100-1:300			
IF 1:50-200			

#### **Images**



Immunohistochemical analysis of paraffin-embedded human brain tissue using Claudin 5 (Phospho-Tyr217) antibody #11792 (left)or the same antibody preincubated with blocking peptide (right).

### Background

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 solutes and water from passing freely through the paracellular space between epithelial or endothelial cell sheets. Mutations in this gene have been found in patients with velocardiofacial syndrome.

Sirotkin H., Genomics 42:245-251(1997).

Collins J.E., Genome Biol. 5:RESEARCH84.1-RESEARCH84.11(2004).

The MGC Project Team; Genome Res. 14:2121-2127(2004).

Note: This product is for in vitro research use only