

## TRPM5 Antibody

Catalog No: #37284

Orders: order@signalwayantibody.com

Support: tech@signalwayantibody.com

## Description

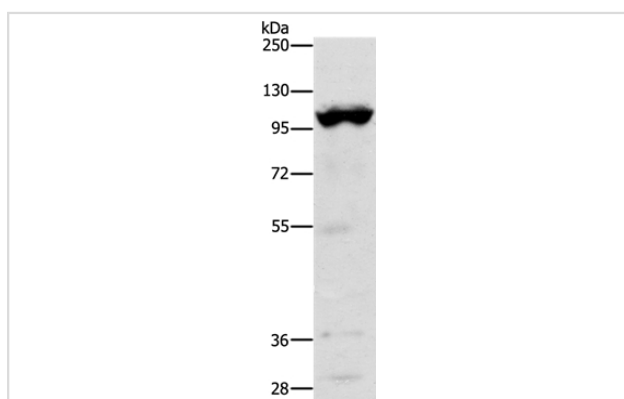
Product Name	TRPM5 Antibody
Host Species	Rabbit
Clonality	Polyclonal
Purification	Antigen affinity purification.
Applications	WB IHC
Species Reactivity	Hu Ms
Specificity	The antibody detects endogenous levels of total TRPM5 protein.
Immunogen Type	Peptide
Immunogen Description	Synthetic peptide corresponding to a region derived from internal residues of human transient receptor potential cation channel, subfamily M, member 5
Target Name	TRPM5
Other Names	MTR1; LTRPC5
Accession No.	Swiss-Prot#: Q9NZQ8NCBI Gene ID: 29850Gene Accssion: NP_055370
Uniprot	Q9NZQ8
GeneID	29850;
SDS-PAGE MW	131kd
Concentration	2.4mg/ml
Formulation	Rabbit IgG in pH7.4 PBS, 0.05% NaN <sub>3</sub> , 40% Glycerol.
Storage	Store at -20°C

## Application Details

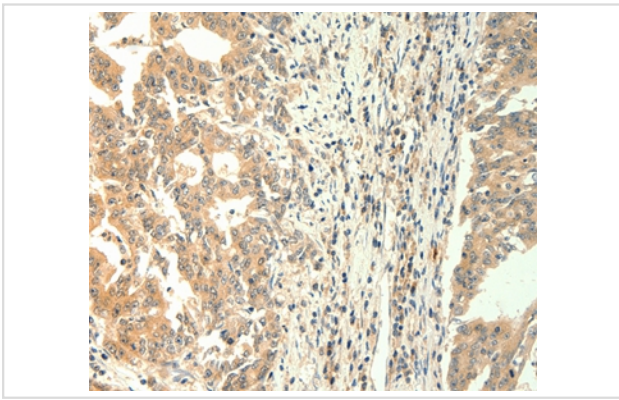
Western blotting: 1:500-1:2000

Immunohistochemistry: 1:50-1:200

## Images



Gel: 10%SDS-PAGE  
 Lysates (from left to right): Mouse heart tissue  
 Amount of lysate: 40ug per lane  
 Primary antibody: 1/1200 dilution  
 Secondary antibody dilution: 1/8000  
 Exposure time: 20 seconds



Immunohistochemical analysis of paraffin-embedded Human gastric cancer tissue using #37284 at dilution 1/60.

## Background

This gene encodes a member of the transient receptor potential (TRP) protein family, which is a diverse group of proteins with structural features typical of ion channels. This protein plays an important role in taste transduction, and has characteristics of a calcium-activated, non-selective cation channel that carries Na<sup>+</sup>, K<sup>+</sup>, and Cs<sup>+</sup> ions equally well, but not Ca<sup>2+</sup> ions. It is activated by lower concentrations of intracellular Ca<sup>2+</sup>, and inhibited by higher concentrations. It is also a highly temperature-sensitive, heat activated channel showing a steep increase of inward currents at temperatures between 15 and 35 degrees Celsius. This gene is located within the Beckwith-Wiedemann syndrome critical region-1 on chromosome 11p15.5, and has been shown to be imprinted, with exclusive expression from the paternal allele.

Note: This product is for in vitro research use only