CDH23 antibody

Catalog No: #38463

Package Size: #38463-1 50ul #38463-2 100ul



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

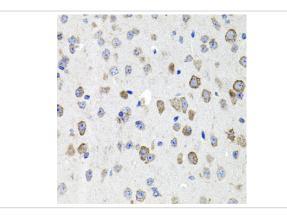
| Description |  |
|-------------|--|
|             |  |

| Product Name          | CDH23 antibody   |
|-----------------------|--|
| Host Species          | Rabbit   |
| Clonality             | Polyclonal   |
| Purification          | Antibodies were purified by affinity purification using immunogen.                                   |
| Applications          | WB,IHC   |
| Species Reactivity    | Human,Mouse  |
| Specificity           | The antibody detects endogenous level of total CDH23 protein.  |
| Immunogen Type        | Protein  |
| Immunogen Description | Fusion protein of human CDH23.   |
| Target Name           | CDH23  |
| Other Names           | DFNB12; DKFZp434P2350; FLJ00233; FLJ36499; KIAA1774; KIAA1812; M GC102761; USH1D; CDHR23;            |
| Accession No.         | Swiss-Prot#: Q9H251NCBI Gene ID: 64072   |
| Uniprot               | Q9H251   |
| GenelD                | 64072;   |
| SDS-PAGE MW           | 45kd   |
| Concentration         | 1.0mg/ml   |
| Formulation           | Supplied at 1.0mg/mL in phosphate buffered saline (without Mg2+ and Ca2+), pH 7.4, 150mM NaCl, 0.02% |
|                       | sodium azide and 50% glycerol.   |
| Storage               | Store at -20°C   |
|                       |  |

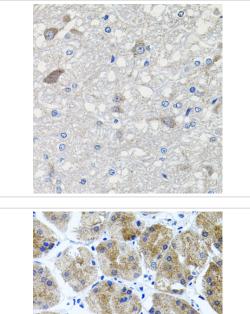
## Application Details

WB 1:500 - 1:2000IHC 1:100 - 1:200

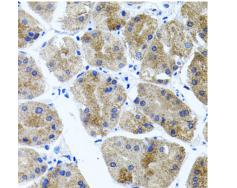
## Images



Immunohistochemistry of paraffin-embedded mouse brain using CDH23 at dilution of 1:100 (40x lens).



Immunohistochemistry of paraffin-embedded rat brain using CDH23 at dilution of 1:100 (40x lens).



Immunohistochemistry of paraffin-embedded human stomach using CDH23 at dilution of 1:100 (40x lens).

## Background

This gene is a member of the cadherin superfamily, whose genes encode calcium dependent cell-cell adhesion glycoproteins. The encoded protein is thought to be involved in stereocilia organization and hair bundle formation. The gene is located in a region containing the human deafness loci DFNB12 and USH1D. Usher syndrome 1D and nonsyndromic autosomal recessive deafness DFNB12 are caused by allelic mutations of this cadherin-like gene. Alternative splice variants encoding different isoforms have been described.

Note: This product is for in vitro research use only