GLRA1 Antibody

Catalog No: #37329

Description



Orders: order@signalwayantibody.com

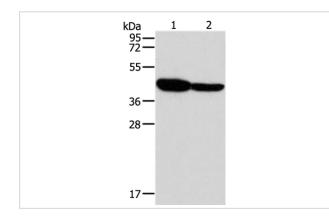
Support: tech@signalwayantibody.com

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|-----------------------|---|
| Product Name | GLRA1 Antibody |
| Host Species | Rabbit |
| Clonality | Polyclonal |
| Purification | Antigen affinity purification. |
| Applications | WB IHC |
| Species Reactivity | Hu Ms Rt |
| Specificity | The antibody detects endogenous levels of total GLRA1 protein. |
| Immunogen Type | Peptide |
| Immunogen Description | Synthetic peptide corresponding to a region derived from internal residues of human Glycine receptor, alpha 1 |
| Target Name | GLRA1 |
| Other Names | STHE; HKPX1 |
| Accession No. | Swiss-Prot#: P23415NCBI Gene ID: 2741Gene Accssion: NP_000162 |
| Uniprot | P23415 |
| GenelD | 2741; |
| SDS-PAGE MW | 53kd |
| Concentration | 1.1mg/ml |
| Formulation | Rabbit IgG in pH7.4 PBS, 0.05% NaN3, 40% Glycerol. |
| Storage | Store at -20°C |
| | |

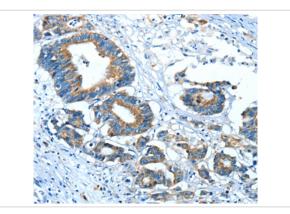
Application Details

Western blotting: 1:500-1:2000 Immunohistochemistry: 1:25-1:100

Images



Gel: 10%SDS-PAGE Lysates (from left to right): Mouse brain and human fetal brain tissue Amount of lysate: 40ug per lane Primary antibody: 1/500 dilution Secondary antibody dilution: 1/8000 Exposure time: 1 minute



Immunohistochemical analysis of paraffin-embedded Human colon cancer tissue using #37329 at dilution 1/30.

Background

The protein encoded by this gene is a subunit of a pentameric inhibitory glycine receptor. The receptor mediates postsynaptic inhibition in the central nervous system. Defects in this gene are a cause of startle disease (STHE), also known as hereditary hyperekplexia or congenital stiff-person syndrome. Two transcript variants encoding different isoforms have been found for this gene.

Note: This product is for in vitro research use only