AGA antibody

Catalog No: #38840

SAB Signalway Antibody

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

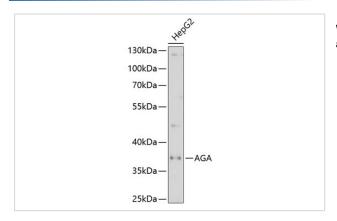
Package Size: #38840-1 50ul #38840-2 100ul

| Description | |
|-----------------------|--|
| Product Name | AGA antibody |
| Host Species | Rabbit |
| Clonality | Polyclonal |
| Purification | Antibodies were purified by affinity purification using immunogen. |
| Applications | WB,IF |
| Species Reactivity | Human,Mouse |
| Specificity | The antibody detects endogenous level of total AGA protein. |
| Immunogen Type | Recombinant Protein |
| Immunogen Description | Recombinant protein of human AGA. |
| Target Name | AGA |
| Other Names | GA; AGU; ASRG; |
| Accession No. | Swiss-Prot#: P20933NCBI Gene ID: 175 |
| Uniprot | P20933 |
| GeneID | 175; |
| SDS-PAGE MW | 37kd |
| 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 - 1:2000IF 1:50 - 1:100

Images



Western blot analysis of extracts of HepG2 cells, using AGA at 1:1000 dilution.

Background

Aspartylglucosaminidase is involved in the catabolism of N-linked oligosaccharides of glycoproteins. It cleaves asparagine from N-acetylglucosamines as one of the final steps in the lysosomal breakdown of glycoproteins. The lysosomal storage disease aspartylglycosaminuria is caused by a deficiency in the AGA enzyme. Alternatively spliced transcript variants have been identified.

Note: This product is for in vitro research use only