

Product datasheet

info@arigobio.com

ARG58933 anti-GM2A antibody

Package: 50 μl Store at: -20°C

Summary

Tested Application

Product Description Rabbit Polyclonal antibody recognizes GM2A

WB

Tested Reactivity Hu
Predict Reactivity Gpig

Host Rabbit

Clonality Polyclonal

Isotype IgG

Target Name GM2A
Species Human

Immunogen Synthetic peptide around the N-terminal region of Human GM2A. (within the following region:

SWDNCDEGKDPAVIRSLTLEPDPIIVPGNVTLSVMGSTSVPLSSPLKVDL)

Conjugation Un-conjugated

Alternate Names Ganglioside GM2 activator; SAP-3; Cerebroside sulfate activator protein; GM2-AP; Sphingolipid

activator protein 3

Application Instructions

Predict Reactivity Note Predicted Homology Based On Immunogen Sequence: Guinea pig: 83%

Application table Application Dilution

WB 0.2 - 1 μg/ml

Application Note * The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations

should be determined by the scientist.

Positive Control HeLa

Properties

Form Liquid

Purification Affinity purified.

Buffer PBS, 0.09% (w/v) Sodium azide and 2% Sucrose.

Preservative 0.09% (w/v) Sodium azide

Stabilizer 2% Sucrose

Concentration Batch dependent: 0.5 - 1 mg/ml

Storage instruction For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot

and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed

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before use.

Note

For laboratory research only, not for drug, diagnostic or other use.

Bioinformation

Gene Symbol GM2A

Gene Full Name GM2 ganglioside activator

Background This gene encodes a small glycolipid transport protein which acts as a substrate specific co-factor for

the lysosomal enzyme beta-hexosaminidase A. Beta-hexosaminidase A, together with GM2 ganglioside activator, catalyzes the degradation of the ganglioside GM2, and other molecules containing terminal Nacetyl hexosamines. Mutations in this gene result in GM2-gangliosidosis type AB or the AB variant of Tay-Sachs disease. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Nov

2009]

Function The large binding pocket can accommodate several single chain phospholipids and fatty acids, GM2A

also exhibits some calcium-independent phospholipase activity (By similarity). Binds gangliosides and stimulates ganglioside GM2 degradation. It stimulates only the breakdown of ganglioside GM2 and glycolipid GA2 by beta-hexosaminidase A. It extracts single GM2 molecules from membranes and presents them in soluble form to beta-hexosaminidase A for cleavage of N-acetyl-D-galactosamine and

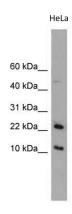
conversion to GM3. [UniProt]

Calculated Mw 21 kDa

PTM The serines in positions 32 and 33 are absent in 80% of the sequenced protein. [UniProt]

Cellular Localization Lysosome. [UniProt]

Images



ARG58933 anti-GM2A antibody WB image

Western blot: HeLa cell lysate stained with ARG58933 anti-GM2A antibody at 0.2 - 1 μ g/ml dilution.