abcam

Product datasheet

Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] ab302977





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Overview

Product name Alexa Fluor® 488 Anti-GFAP antibody [EP672Y]

Description Alexa Fluor® 488 Rabbit monoclonal [EP672Y] to GFAP

Host species Rabbit

Conjugation Alexa Fluor® 488. Ex: 495nm, Em: 519nm

Tested applications Suitable for: IHC-P. ICC/IF

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Human cerebrum, astrocytoma; mouse cerebrum; rat cerebrum FFPE tissue sections.

ICC/IF: Rat primary neural/glia cells.

General notes This product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity
- Long-term security of supply
- Animal-free production

For more information see here.

Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**® **patents**.

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outlicensing@thermofisher.com.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C long

term. Avoid freeze / thaw cycle. Store In the Dark.

Storage buffer pH: 7.40

Preservative: 0.02% Sodium azide

Constituents: 68% PBS, 30% Glycerol (glycerin, glycerine), 1% BSA

Purity Protein A purified

Clonality Monoclonal
Clone number EP672Y

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab302977 in the following tested applications.

Table Guarantee overs the use of about 277 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
IHC-P		1/50 - 1/100. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
ICC/IF		1/50.

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Function GFAP, a class-Ill intermediate filament, is a cell-specific marker that, during the development of

the central nervous system, distinguishes astrocytes from other glial cells.

Tissue specificity Expressed in cells lacking fibronectin.

Involvement in disease Defects in GFAP are a cause of Alexander disease (ALEXD) [MIM:203450]. Alexander disease

is a rare disorder of the central nervous system. It is a progressive leukoencephalopathy whose hallmark is the widespread accumulation of Rosenthal fibers which are cytoplasmic inclusions in astrocytes. The most common form affects infants and young children, and is characterized by progressive failure of central myelination, usually leading to death usually within the first decade. Infants with Alexander disease develop a leukoencephalopathy with macrocephaly, seizures, and psychomotor retardation. Patients with juvenile or adult forms typically experience ataxia, bulbar

signs and spasticity, and a more slowly progressive course.

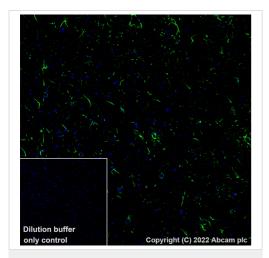
Sequence similarities Belongs to the intermediate filament family.

Post-translational Phosphorylated by PKN1.

modifications

Cellular localization Cytoplasm. Associated with intermediate filaments.

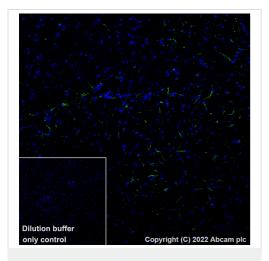
Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] (ab302977)

Immunohistochemical analysis of paraffin-embedded rat cerebrum tissue labeling GFAP with ab302977 at 1/50 dilution (10.0 μ g/mL). Positive staining is observed on astrocytes of rat cerebrum. The section was incubated with ab302977 for 60 mins at room temperature (shown in green). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

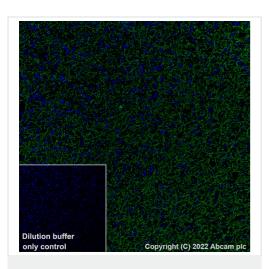
Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 40 mins was used.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] (ab302977)

Immunohistochemical analysis of paraffin-embedded mouse cerebrum tissue labeling GFAP with ab302977 at 1/50 dilution (10.0 μ g/mL). Positive staining is observed on astrocytes of mouse cerebrum. The section was incubated with ab302977 for 60 mins at room temperature (shown in green). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

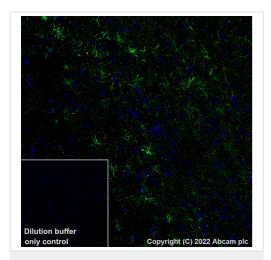
Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 40 mins was used.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] (ab302977)

Immunohistochemical analysis of paraffin-embedded human astrocytoma tissue labeling GFAP with ab302977 at 1/100 dilution (5.0 µg/mL). Positive staining is observed on human astrocytoma. The section was incubated with ab302977 for 60 mins at room temperature (shown in green). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

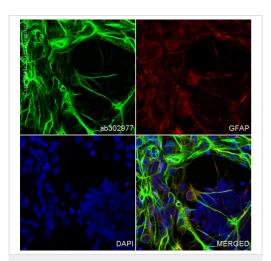
Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 40 mins was used.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] (ab302977)

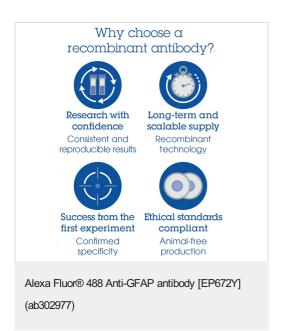
Immunohistochemical analysis of paraffin-embedded human cerebrum tissue labeling GFAP with ab302977 at 1/100 dilution (5.0 μ g/mL). Positive staining is observed on astrocytes of human cerebrum. The section was incubated with ab302977 for 60 mins at room temperature (shown in green). Nuclear DNA was labeled with DAPI (shown in blue). The section was then mounted using Fluoromount[®]. The immunostaining was performed on a Leica Biosystems BOND[®] RX instrument. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8).

Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0, epitope retrieval solution 2) for 40 mins was used.



Immunocytochemistry/ Immunofluorescence - Alexa Fluor® 488 Anti-GFAP antibody [EP672Y] (ab302977)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized rat primary neural/glia cells labeling GFAP with ab302977 at 1/50 dilution (10 µg/ml) (Green). Confocal image showing positive staining in rat primary glia cell. Confocal scanning Z step was set as 0.3 µm followed by image processing with maximum Z projection. Image was taken with a confocal microscope (Leica-Microsystems, TCS SP8). Anti-Glial Fibrillary Acidic Protein (GFAP) mouse monoclonal antibody was used as counterstain at 1/100 (1ug/ml) dilution (Red). The nuclear counterstain was DAPI (Blue).



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