



Product Information Sheet

Monoclonal Anti-Glial Fibrillary Acidic Protein, *GFAP*

Catalogue No. MA1045

Immunogen

GFAP from pig spinal cord.

Lot No. 08A12

Purification

Purified by the goat anti-mouse IgG affinity chromatography.

Clone: GA-8

Ig type: mouse IgG1

Application

Western blot

Size: 100µg/vial

At 0.5-1µg/ml with the appropriate system to detect GFAP in cells and tissues.

Specificity

Human, mouse, rat.

No cross reactivity with other proteins.

Immunohistochemistry(P)

At 0.4-1µg/ml to detect GFAP in formalin fixed and paraffin embedded tissues.

Immunohistochemistry(F)

At 0.5-1µg/ml to detect GFAP in formalin or acetone fixed tissues.

Other applications have not been tested.

Optimal dilutions should be determined by end user.

Recommended application

Western blot

Immunohistochemistry(P)

Immunohistochemistry(F)

Formulation

Lyophilized from 1.2% sodium acetate, with 2mg BSA and 0.01mg NaN₃ as preservative.

Reconstitution

1.2% sodium acetate or neutral PBS. If 1ml of PBS is used, the antibody concentration will be 100µg/ml.

To reorder contact us at:

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Storage

At -20°C for one year. After reconstitution, at 4°C for one month. It can also be aliquotted and stored frozen at -20°C for longer time.

BACKGROUND

Glial fibrillary acidic protein(GFAP) is an intermediate filament protein of 52Kda. GFAP gene is mapped to human 17q21. GFAP is a useful marker of astroglia in the brain. Mutations in GFAP, encoding glial fibrillary acidic protein, are associated with Alexander disease

REFERENCE

- 1 Brenner, M.; Johnson, A. B.; Boespflug-Tanguy, O.; Rodriguez, D.; Goldman, J. E.; Messing, A. : Mutations in GFAP, encoding glial fibrillary acidic protein, are associated with Alexander disease. *Nature Genet.* 27: 117-120, 2001.
- 2 Rodriguez, D.; Gauthier, F.; Bertini, E.; Bugiani, M.; Brenner, M.; N'guyen, S.; Goizet, C.; Gelot, A.; Surtees, R.; Pedespan, J.-M.; Hernandorena, X.; Troncoso, M.; Uziel, G.; Messing, A.; Ponsot, G.; Pham-Dinh, D.; Dautigny, A.; Boespflug-Tanguy, O. : Infantile Alexander disease: spectrum of GFAP mutations and genotype-phenotype correlation. *Am. J. Hum. Genet.* 69: 1134-1140, 2001. Note: Erratum: *Am. J. Hum. Genet.* 69: 1413 only, 2001.