## GFAP (GA5) Mouse mAb



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IHC-P, IF-F, IF-IC, FC-FP	Reactivity: H M R	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 50	Source/Isotype: Mouse IgG1 kappa	<b>UniProt ID:</b> #P14136	Entrez-Gene Id: 2670
Product Usage Information		Application			Dilution	
		Western Blotting Immunohistochemistry (Paraffin)			1:1000 1:50 - 1:200	
			•			
		Immunofluorescence Immunofluorescence	,	1:400 - 1:800 1:800 - 1:1600		
		Flow Cytometry (Fixed	•	iisti y)		0 - 1:1600 0 - 1:1600
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 μg/ml BSA, 50% glycerol and less than 0.02% sodium azide. Store at –20°C. Do not aliquot the antibody.				
		For a carrier-free (BSA and azide free) version of this product see product #56522.				
Specificity/Sensitivity		GFAP (GA5) Mouse mAb detects endogenous levels of total GFAP protein.				
Source / Purification		Monoclonal antibody is produced by immunizing animals with native GFAP purified from pig spinal cord.				
Background		The cytoskeleton consists of three types of cytosolic fibers: microfilaments (actin filaments), intermediate filaments, and microtubules. Major types of intermediate filaments are specifically expressed in particular cell types: cytokeratins in epithelial cells, glial fibrillary acidic protein (GFAP) in glial cells, desmin in skeletal, visceral, and certain vascular smooth muscle cells, vimentin in cells of mesenchymal origin, and neurofilaments in neurons. GFAP and vimentin form intermediate filaments in astroglial cells and modulate their motility and shape (1). In particular, vimentin filaments are present at early developmental stages, while GFAP filaments are characteristic of differentiated and mature brain astrocytes. Thus, GFAP is commonly used as a marker for intracranial and intraspinal tumors arising from astrocytes (2). In addition, GFAP intermediate filaments are also present in nonmyelin-forming Schwann cells in the peripheral nervous system (3).				
Background References		1. Eng, L.F. et al. (2000) <i>Neurochem. Res.</i> 25, 1439-51. 2. Goebel, H.H. et al. (1987) <i>Acta. Histochem. Suppl.</i> 34, 81-93. 3. Jessen, K.R. et al. (1990) <i>Development</i> 109, 91-103.				
Species Reactiv	rity	Species reactivity is d	etermined by testir	g in at least one approve	d application (e.g.	, western blot).

**Western Blot Buffer** IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat

dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

W: Western Blotting IHC-P: Immunohistochemistry (Paraffin) IF-F: Immunofluorescence (Frozen) IF-IC: **Applications Key** 

Immunofluorescence (Immunocytochemistry) FC-FP: Flow Cytometry (Fixed/Permeabilized)

**Cross-Reactivity Key** H: Human M: Mouse R: Rat

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