

Product Data Sheet

Anti-GFAP Antibody

Catalog #	Source	Reactivity	Applications		
CQA3697	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	bbit polyclonal antibody	/ to GFAP		
Immunogen	Re	combinant protein of h	uman GFAP. The exact sequence is proprietary.		
Purification	Th	e antibody was purified	by immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous le	vels of GFAP protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Lic	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	d 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/	100 - 1⁄200), IF/IC (1⁄100 - 1⁄500)		
Gene Symbol	GF	AP			
Alternative Na	ames Gli	ial fibrillary acidic protei	n; GFAP		
Entrez Gene	26	70 (Human); 14580 (Mc	ouse); 24387 (Rat)		
SwissProt		P14136 (Human); P03995 (Mouse); P47819 (Rat)			
Storage/Stabi	lity Sh	ipped at 4°C. Upon deliv	very aliquot and store at -20°C for one year. Avoid		
	fre	eze/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

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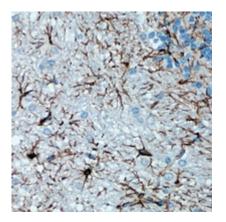
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For research purposes only, not for human use

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Western blot analysis of GFAP expression in mouse brain (A), rat brain (B) whole cell lysates. (Predicted band size: 49; 50 kD; Observed band size: 45-50 kD)



Immunohistochemical analysis of GFAP staining in human brain formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of GFAP staining in U251MG cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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