CoraLite®594-conjugated GFAP Monoclonal antibody

Catalog Number:CL594-60190 1 Publications



Basic Information	Catalog Number: CL594-60190	GenBank Accession Number: BC013596	Purification Method: Protein A purification	
	Size:	GenelD (NCBI):	CloneNo.:	
	1000 µg/ml		4B2E10	
	Mouse	P14136	IF 1:50-1:500	
	lsotype: IgG2a	Full Name: glial fibrillary acidic protein	Excitation/Emission maxima wavelengths:	
	Immunogen Catalog Number: AG10452	Calculated MW: 432 aa, 50 kDa	588 nm / 604 nm	
Applications	Tested Applications:	Positive Controls: IF : mouse brain tissue,		
	Cited Applications: IF			
	Species Specificity: human, mouse, rat, pig			
	Cited Species: mouse			
Background Informatic	GFAP Function GFAP (Glial fibril central nervous system (CNS). Gf astrocytes and has been propose strength, and in mitosis. Tissue s astrocytes. GFAP is commonly us non-CNS cells, including fibrobla Involvement in disease Mutatio CNS disorder. The mutations pres GFAP is a hallmark of reactive as astrogliosis is present in many n (including Alzheimer's and Parki isoforms of GFAP that differ in th size. Isoform expression varies d are upregulated in reactive astro regulated by phosphorylation. Si which are reported to control fila intermediate filaments and stair Ex/Em 593 nm/614 nm.	GFAP Function GFAP (Glial fibrillary acidic protein) is a type III intermediate filament (IF) protein specific to the central nervous system (CNS). GFAP is one of the main components of the intermediate filament network in astrocytes and has been proposed as playing a role in cell migration, cell motility, maintaining mechanical strength, and in mitosis. Tissue specificity GFAP is expressed in central nervous system cells, predominantly in astrocytes. GFAP is commonly used as an astrocytes, and liver stellate cells (PMID: 21219963). Involvement in disease Mutations in GFAP lead to Alexander disease (OMIM: 203450), an autosomal dominant CNS disorder. The mutations present in affected individuals are thought to be gain-of-function. Upregulation of GFAP is a hallmark of reactive astrocytes, in which GFAP is present in hypertrophic cellular processes. Reactive astrogliosis is present in many neurological disorders, such as stroke, various neurodegenerative diseases (including Alzheimer's and Parkinson's disease), and neurotrauma. Isoforms Astrocytes express 10 different isoforms of GFAP that differ in the rod and tail domains (PMID: 25726916), which means that they differ in molecular size. Isoform expression varies during the development and across different subtypes of astrocytes. Not all isoforms are upregulated in reactive astrocytes. Post-translational modifications Intermediate filament proteins are regulated by phosphorylation. Six phosphorylation sites have been identified in GFAP localizes to intermediate filaments and stains well in astrocyte cellular processes. The antibody is conjugated with CL594, Ex/Em 593 nm/614 nm.		
Notable Publications	Author	Pubmed ID Journal	Application	
	Yue Wan	36598105 Glia	IF	
Storage	Storage: Store at -20°C. Avoid exposure to Storage Buffer: PBS with 50% Glycerol, 0.05% Pi Aliquoting is unnecessary for -20	light. Stable for one year after shipme roclin300, 0.5% BSA, pH 7.3. °C storage	int.	

For technical support and original validation data for this product please contact: T: 4006900926 E: Proteintech-CN@ptglab.com W: ptgcn.com This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

Selected Validation Data





Immunofluorescent analysis of (4% PFA) fixed paraffin-embedded mouse brain tissue using CoraLite® 594 GFAP antibody (CL594-60190, Clone: 4B2E10) at dilution of 1:100. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0). Immunofluorescent analysis of (4% PFA) fixed paraffin-embedded mouse brain tissue using CoraLite® 594 GFAP antibody (CL594-60190, Clone: 4B2E10) at dilution of 1:100. Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).