

# Glial Fibrillary Acidic Protein (GFAP) (PT0129R) Rabbit mAb (Ready to Use)

CatalogNo: YM7125R Recombinant R

## **Key Features**

**Host Species** 

Rabbit

Reactivity

· Human, Rat,

**Applications** 

IHC

Isotype

• IgG1,Kappa

#### **Recommended Dilution Ratios**

Ready to use for IHC

## Storage

Storage\* 2°C to 8°C/1 year

**Formulation** The prediluted ready-to-use antibody is diluted in phosphate buffer saline containing

stabilizing protein and 0.05% Proclin 300

#### **Basic Information**

**Clonality** Monoclonal

Clone Number PT0129R

# Immunogen Information

Immunogen Synthesized peptide derived from human Glial Fibrillary Acidic Protein (GFAP) AA

range:300-432

**Specificity** This antibody detects endogenous levels of GFAP

# | Target Information

Gene name GFAP

**Protein Name** Glial fibrillary acidic protein (GFAP)

Organism Gene ID UniProt ID

Human <u>2670;</u> <u>P14136;</u>

Cellular Localization Cytoplasmic

**Tissue specificity** Expressed in cells lacking fibronectin.

**Function** alternative products:Isoforms differ in the C-terminal region which is encoded by alternative

exons, 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.,function:GFAP, a class-III intermediate filament, is a cell-specific marker that, during the development of the central nervous system, distinguishes astrocytes from other glial cells.,online information:GFAP entry,similarity:Belongs to the intermediate filament family.,subcellular location:Associated with intermediate

filaments., subunit: Interacts with SYNM (By similarity). Isoform 3 interacts with PSEN1 (via N-

terminus)., tissue specificity: Expressed in cells lacking fibronectin.,

## | Validation Data

#### Contact information

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