

GFAP (ABT331) mouse mAb

Catalog No: YM6951

Reactivity: Human; Rat;

Applications: IHC;ELISA

Target: GFAP

Fields: >>JAK-STAT signaling pathway

Gene Name: GFAP

Protein Name: GFAP

Human Gene Id: 2670

Human Swiss Prot

No:

Immunogen: Synthesized peptide derived from human GFAP AA range: 300-432

Specificity: The antibody can specifically recognize human GFAP protein.

Formulation: Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Source: Mouse, Monoclonal/lgG1, Kappa

P14136

Dilution : IHC 1:200-400, ELISA 1:5000-20000

Purification: The antibody was affinity-purified from mouse ascites by affinity-

chromatography using specific immunogen.

Storage Stability: -15°C to -25°C/1 year(Do not lower than -25°C)

Background: This gene encodes one of the major intermediate filament proteins of mature

astrocytes. It is used as a marker to distinguish astrocytes from other glial cells during development. Mutations in this gene cause Alexander disease, a rare disorder of astrocytes in the central nervous system. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Oct

1/2



2008],

Function:

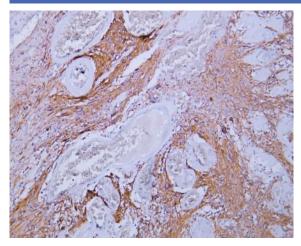
alternative products:Isoforms differ in the C-terminal region which is encoded by alternative exons, disease:Defects in GFAP are a cause of Alexander disease (ALEXD) [MIM:203450]. Alexander disease is a rare disorder of the central nervous system. It is a progressive leukoencephalopathy whose hallmark 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-spe

Subcellular Location :

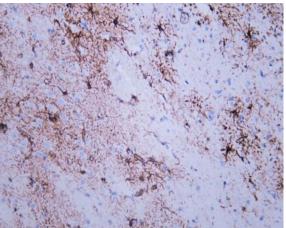
Cytoplasmic

Expression: Expressed in cells lacking fibronectin.

Products Images



Human astrocytoma tissue was stained with Anti-Glial Fibrillary Acidic Protein (GFAP) (ABT331) Antibody



Human cerebrum tissue was stained with Anti-Glial Fibrillary Acidic Protein (GFAP) (ABT331) Antibody