

Recombinant Human GFAP protein ab114149

5 References [画像数 2](#)

製品の詳細

製品名	Recombinant Human GFAP protein	
発現系	Wheat germ	
アクセッション番号	P14136	
タンパク質長	Full length protein	
Animal free	No	
由来	Recombinant	
生物種	Human	
配列		<p>MERRRITSAARRSYVSSGEMMVGGLAPGRRLGPGTRLSLARM PPPLPTRV DFSLAGALNAGFKETRASERAEMMELNDRFASYIEKVRFLEQ QNKALAAE LNQLRAKEPTKLADVYQAEELRELRLRLDQLTANSARLEVERD NLAQDLAT VRQKLQDETNLRLAENNLAAAYRQEADEATLARLDLERKIES LEEEIRFL RKIHEEEVRELQEQLARQQVHVVELDVAKPDLTAALKEIRTQY EAMASSNM HEAEEWYRSKFADLTDAARNAELLRQAKHEANDYRRQLQSL TCDLESRL GTNESLERQMREQEERHVREAASYQEALARLEEEGQSLKDEM ARHLQEYQ DLLNVKLALDIEIATYRKLLERGEENRITIPVQTFSNLQIRET SLDTKSVS EGHLLKRNIVVKTVEMRDGEVIKESKQEHKDVM</p>
予測される分子量	76 kDa including tags	
領域	1 to 432	
タグ	GST tag N-Terminus	

特性

Our **Abpromise guarantee** covers the use of **ab114149** in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

アプリケーション ELISA

Western blot

SDS-PAGE

製品の状態

Liquid

備考

前処理および保存

保存方法および安定性

Shipped on dry ice. Upon delivery aliquot and store at -80°C. Avoid freeze / thaw cycles.

pH: 8.00

Constituents: 0.3% Glutathione, 0.79% Tris HCl

関連情報

機能

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.

組織特異性

Expressed in cells lacking fibronectin.

関連疾患

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.

配列類似性

Belongs to the intermediate filament family.

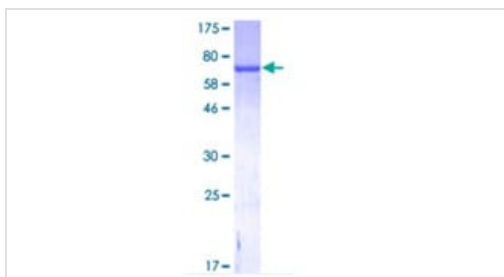
翻訳後修飾

Phosphorylated by PKN1.

細胞内局在

Cytoplasm. Associated with intermediate filaments.

画像



12.5% SDS-PAGE analysis of ab114149, stained with Coomassie Blue.

SDS-PAGE - Recombinant Human GFAP protein (ab114149)



Western blot - Recombinant Human GFAP protein (ab114149)

Anti-GFAP antibody - Astrocyte Marker (**ab48050**) at 1 µg/ml +
Recombinant Human GFAP protein (ab114149) at 0.1 µg

Secondary

Goat Anti-Rabbit IgG H&L (HRP) preadsorbed (**ab97080**) at 1/5000 dilution

Developed using the ECL technique.

Performed under reducing conditions.

Exposure time: 10 seconds

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