abcam

Product datasheet

Recombinant Human GLB1/Beta-galactosidase protein ab151890

Description

Product name	Recombinant Human GLB1/Beta-galactosidase protein
Purity	> 95 % SDS-PAGE. ab151890 has greater than 95% purity as determined by SEC-HPLC and reducing SDS-PAGE.
Endotoxin level	< 0.100 Eu/µg
Expression system	HEK 293 cells
Accession	P16278
Protein length	Full length protein
Animal free	No
Nature	Recombinant
Species	Human
Sequence	NATQRMFEIDYSRDSFLKDGQPFRYISGSIHYSRVPRFYWK DRLLKMKMA GLNAIQTYVPWNFHEPWPGQYQFSEDHDVEYFLRLAHEL GLLVILRPGPY ICAEWEMGGLPAWLLEKESILLRSSDPDYLAAVDKWLGV LLPKMKPLLYQ NGGPVITVQVENEYGSYFACDFDYLRFLQKRFRHHLGDDV VLFTTDGAHK TFLKCGALQGLYTTVDFGTGSNITDAFLSQRKCEPKGPLIN SEFYTGWLD HWGQPHSTIKTEAVASSLYDILARGASVNLYMFIGGTNFAY WNGANSPYA AQPTSYDYDAPLSEAGDLTEKYFALRNIIQKFEKVPEGPIP PSTPKFAYG KVTLEKLKTVGAALDILCPSGPIKSLYPLTFIQVKQHYGFVL YRTTLPQD CSNPAPLSSPLNGVHDRAYVAVDGIPQGVLERNNVITLNIT GKAGATLDL LVENMGRVNYGAYINDFKGLVSNLTLSSNILTDWTIFPLDTE DAVRSHLG GWGHRDSGHHDEAWAHNSSNYTLPAFYMGNFSIPSGIPD LPQDTFIQFPG WTKGQVWINGFNLGRYWPARGPQLTLFVPQHILMTSAPNT

ITVLELEWAP
CSSDDPELCAVTFVDRPVIGSSVTYDHPSKPVEKRLMPP
PPQKNKDSWLD HVVDHHHHHH

Predicted molecular weight	74 kDa including tags
Amino acids	24 to 677
Tags	His tag C-Terminus

Specifications

Our **Abpromise guarantee** covers the use of **ab151890** in the following tested applications.

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

Applications	SDS-PAGE HPLC
Form	Liquid

Preparation and Storage

Stability and Storage	Shipped on dry ice. Upon delivery aliquot and store at -80ºC. Avoid freeze / thaw cycles. pH: 8.00 Constituents: 0.32% Tris HCl, 0.88% Sodium chloride

General Info

Function	Cleaves beta-linked terminal galactosyl residues from gangliosides, glycoproteins, and glycosaminoglycans. Isoform 2 has no beta-galactosidase catalytic activity, but plays functional roles in the formation of extracellular elastic fibers (elastogenesis) and in the development of connective tissue. Seems to be identical to the elastin-binding protein (EBP), a major component of the non-integrin cell surface receptor expressed on fibroblasts, smooth muscle cells, chondroblasts, leukocytes, and certain cancer cell types. In elastin producing cells, associates with tropoelastin intracellularly and functions as a recycling molecular chaperone which facilitates the secretions of tropoelastin and its assembly into elastic fibers.
Involvement in disease	Defects in GLB1 are the cause of GM1-gangliosidosis type 1 (GM1G1) [MIM:230500]; also known as infantile GM1-gangliosidosis. GM1-gangliosidosis is an autosomal recessive lysosomal storage disease marked by the accumulation of GM1 gangliosides, glycoproteins and keratan sulfate primarily in neurons of the central nervous system. GM1G1 is characterized by onset within the first three months of life, central nervous system degeneration, coarse facial features, hepatosplenomegaly, skeletal dysmorphology reminiscent of Hurler syndrome, and rapidly progressive psychomotor deterioration. Urinary oligosaccharide levels are high. It leads to death usually between the first and second year of life. Defects in GLB1 are the cause of GM1-gangliosidosis type 2 (GM1G2) [MIM:230600]; also known as late infantile/juvenile GM1-gangliosidosis. GM1G2 is characterized by onset between ages 1 and 5. The main symptom is locomotor ataxia, ultimately leading to a state of decerebration with epileptic seizures. Patients do not display the skeletal changes associated with the infantile form, but they nonetheless excrete elevated amounts of beta-linked galactose-terminal oligosaccharides. Inheritance is autosomal recessive. Defects in GLB1 are the cause of GM1-gangliosidosis type 3 (GM1G3) [MIM:230650]; also

	known as adult or chronic GM1-gangliosidosis. GM1G3 is characterized by a variable phenotype. Patients show mild skeletal abnormalities, dysarthria, gait disturbance, dystonia and visual impairment. Visceromegaly is absent. Intellectual deficit can initially be mild or absent but progresses over time. Inheritance is autosomal recessive. Defects in GLB1 are the cause of mucopolysaccharidosis type 4B (MPS4B) [MIM:253010]; also known as Morquio syndrome B. MPS4B is a form of mucopolysaccharidosis type 4, an autosomal recessive lysosomal storage disease characterized by intracellular accumulation of keratan sulfate and chondroitin-6-sulfate. Key clinical features include short stature, skeletal dysplasia, dental anomalies, and corneal clouding. Intelligence is normal and there is no direct central nervous system involvement, although the skeletal changes may result in neurologic complications. There is variable severity, but patients with the severe phenotype usually do not survive past the second or third decade of life.
Sequence similarities	Belongs to the glycosyl hydrolase 35 family.
Cellular localization	Lysosome and Cytoplasm > perinuclear region. Localized to the perinuclear area of the cytoplasm but not to lysosomes.

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