

Product datasheet

Recombinant Human LAMTOR2 protein ab101637

1 Image

Description

Product name	Recombinant Human LAMTOR2 protein
Purity	> 95 % SDS-PAGE. ab101637 was purified using conventional chromatography techniques.
Expression system	Escherichia coli
Accession	<u>Q9Y2Q5</u>
Protein length	Full length protein
Animal free	No
Nature	Recombinant
Species	Human
Sequence	MGSSHHHHHHSSGLVPRGSHMGSHMLRPKALTQVLSQ ANTGGVQSTLLLN NEGSLLAGSYGDTDARVTAAIASNIWAAYDRNGNQAFNE DNLKFILMDC MEGRVAITRVANLLLCMYAKETVGFGMLKAKAQALVQYLE EPLTQVAAS
Predicted molecular weight	16 kDa including tags
Amino acids	1 to 125
Tags	His tag N-Terminus

Specifications

Our **Abpromise guarantee** covers the use of **ab101637** 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 Mass Spectrometry
Mass spectrometry	MALDI-TOF
Form	Liquid

Preparation and Storage

Stability and Storage	Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -
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80°C. Avoid freeze / thaw cycle.

pH: 8.00

Constituents: 0.0308% DTT, 0.316% Tris HCl, 10% Glycerol (glycerin, glycerine), 1.16% Sodium chloride

General Info

Function

As part of the Ragulator complex it is involved in amino acid sensing and activation of mTORC1, a signaling complex promoting cell growth in response to growth factors, energy levels, and amino acids. Activated by amino acids through a mechanism involving the lysosomal V-ATPase, the Ragulator functions as a guanine nucleotide exchange factor activating the small GTPases Rag. Activated Ragulator and Rag GTPases function as a scaffold recruiting mTORC1 to lysosomes where it is in turn activated. Adapter protein that enhances the efficiency of the MAP kinase cascade facilitating the activation of MAPK2.

Involvement in disease

Defects in LAMTOR2 are the cause of immunodeficiency due to defect in MAPBP-interacting protein (ID-MAPBPIP) [MIM:610798]. This form of primary immunodeficiency syndrome includes congenital neutropenia, partial albinism, short stature and B-cell and cytotoxic T-cell deficiency.

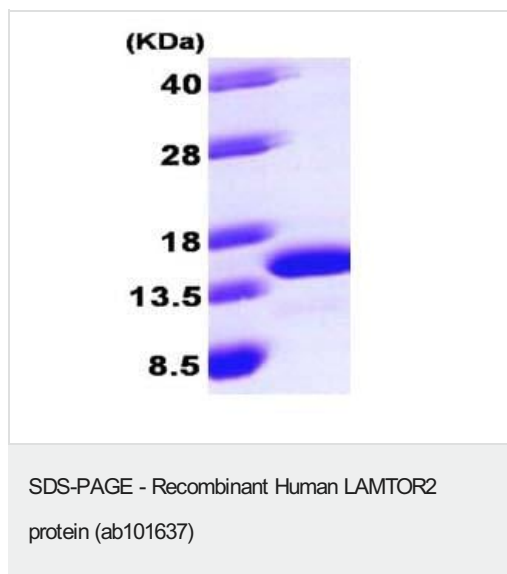
Sequence similarities

Belongs to the GAMAD family.

Cellular localization

Late endosome membrane. Lysosome membrane.

Images



ab101637 (3 µg) analyzed by 15% SDS PAGE.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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