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Datasheet for ABIN3120620 **Fat Storage-Inducing Transmembrane Protein 2 (FITM2) (AA 1-262) protein (Strep Tag)**



Overview

e-Inducing Transmembrane Protein 2 (FITM2) Nicotiana tabacum)
Nicotiana tabacum)
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ant
S-PAGE (SDS), Western Blotting (WB)
AWF LRGTLVRATV RRHLPWALVA AMLAGSVVKE LSPLPESYLS NKRNVLNVYF
TVC LLLPFIALTN YHLTGKTSLV LRRLSTLLVG TAIWYICTAL FSNIEHYTGS
GI RQEHRSKQQC HREGGFWHGF DISGHSFLLT FCALMIVEEM AVLHEVKTDR
ITT LVVALGFLTF IWVWMFLCTA VYFHDLTQKV FGTMFGLLGW YGTYGYWYLK
PQS CSLTLKRDTY KK
without tag. The proposed Strep-Tag is based on experience s with the expression
different complexity of the protein could make another tag necessary. In case you
cial request, please contact us.
ts:
Germany - from design to production - by highly experienced protein experts.
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- These proteins are normally active (enzymatically functional) as our customers have reported (not tested by us and not guaranteed).
- State-of-the-art algorithm used for plasmid design (Gene synthesis).

This protein is a **made-to-order protein** and will be made for the first time for your order. Our experts in the lab will ensure that you receive a correctly folded protein. The big advantage of ordering our **made-to-order proteins** in comparison to ordering custom made proteins from other companies is that there is no financial obligation in case the protein cannot be expressed or purified.

Expression System:

- ALICE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from Nicotiana tabacum c.v.. This contains all the protein expression machinery needed to produce even the most difficult-to-express proteins, including those that require post-translational modifications.
- During lysate production, the cell wall and other cellular components that are not required for
 protein production are removed, leaving only the protein production machinery and the
 mitochondria to drive the reaction. During our lysate completion steps, the additional
 components needed for protein production (amino acids, cofactors, etc.) are added to
 produce something that functions like a cell, but without the constraints of a living system all that's needed is the DNA that codes for the desired protein!

Concentration:

- The concentration of our recombinant proteins is measured using the absorbance at 280nm.
- The protein's absorbance will be measured in several dilutions and is measured against its specific reference buffer.
- We use the Expasy's protparam tool to determine the absorption coefficient of each protein.

Purification:	Two step purification of proteins expressed in Almost Living Cell-Free Expression System (ALiCE®):
	 In a first purification step, the protein is purified from the cleared cell lysate using StrepTag capture material. Eluate fractions are analyzed by SDS-PAGE. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatography. Eluate fractions are analyzed by SDS-PAGE and Western blot.
Purity:	\ge 80 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)

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Product Details

Grade:

Crystallography grade

Target Details

Target:	Fat Storage-Inducing Transmembrane Protein 2 (FITM2)
Alternative Name:	Fitm2 (FITM2 Products)
Background:	Acyl-coenzyme A diphosphatase FITM2 (EC 3.6.1) (Fat storage-inducing transmembrane protein 2) (Fat-inducing protein 2),FUNCTION: Fatty acyl-coenzyme A (CoA) diphosphatase tha hydrolyzes fatty acyl-CoA to yield acyl-4'-phosphopantetheine and adenosine 3',5'-bisphosphata (By similarity). Preferentially hydrolyzes unsaturated long-chain acyl-CoA substrates such as oleoyl-CoA/(9Z)-octadecenoyl-CoA and arachidonoyl-CoA/(5Z,8Z,11Z,14Z)-eicosatetraenoyl-CoA in the endoplasmic reticulum (ER) lumen (By similarity). This catalytic activity is required for maintaining ER structure and for lipid droplets (LDs) biogenesis, which are lipid storage organelles involved in maintaining lipid and energy homeostasis (PubMed:18160536, PubMed:26504167) (By similarity). Directly binds to diacylglycerol (DAGs) and triacylglycerol, which is also important for LD biogenesis (PubMed:22106267) (By similarity). May support directional budding of nacent LDs from the ER into the cytosol by reducing DAG levels at sites of LD formation (By similarity). Plays a role in the regulation of cell morphology and cytoskeleta organization (By similarity). {ECO:0000255 HAMAP-Rule:MF_03230, ECO:0000269 PubMed:26504167}.
Molecular Weight:	30.0 kDa
UniProt:	P59266
Application Details	
Application Notes:	In addition to the applications listed above we expect the protein to work for functional studies as well. As the protein has not been tested for functional studies yet we cannot offer a guarantee though.
Comment:	ALICE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from Nicotiana tabacum c.v This contains all the protein expression machinery needed to produce even the most difficult-to-express proteins, including those that require post-translational modifications.

During lysate production, the cell wall and other cellular components that are not required for protein production are removed, leaving only the protein production machinery and the

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Application Details		
	mitochondria to drive the reaction. During our lysate completion steps, the additional components needed for protein production (amino acids, cofactors, etc.) are added to produce something that functions like a cell, but without the constraints of a living system - all that's needed is the DNA that codes for the desired protein!	
Restrictions:	For Research Use only	
Handling		
Format:	Liquid	
Buffer:	The buffer composition is at the discretion of the manufacturer. If you have a special request, please contact us.	
Handling Advice:	Avoid repeated freeze-thaw cycles.	

Storage:-80 °CStorage Comment:Store at -80°C.Expiry Date:Unlimited (if stored properly)