

Datasheet for ABIN3134932

TRIM72 Protein (AA 1-477) (Strep Tag)



Overview

Quantity:	1 mg
Target:	TRIM72
Protein Characteristics:	AA 1-477
Origin:	Mouse
Source:	Cell-free protein synthesis (CFPS)
Protein Type:	Recombinant
Purification tag / Conjugate:	This TRIM72 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details	
Brand:	AliCE®
Sequence:	MSAAPGLLRQ ELSCPLCLQL FDAPVTAECG HSFCRACLIR VAGEPAADGT VACPCCQAPT
	RPQALSTNLQ LSRLVEGLAQ VPQGHCEEHL DPLSIYCEQD RTLVCGVCAS LGSHRGHRLL
	PAAEAQARLK TQLPQQKMQL QEACMRKEKT VAVLEHQLVE VEETVRQFRG AVGEQLGKMR
	MFLAALESSL DREAERVRGD AGVALRRELS SLNSYLEQLR QMEKVLEEVA DKPQTEFLMK
	FCLVTSRLQK ILSESPPPAR LDIQLPVISD DFKFQVWKKM FRALMPALEE LTFDPSSAHP
	SLVVSSSGRR VECSDQKAPP AGEDTRQFDK AVAVVAQQLL SQGEHYWEVE VGDKPRWALG
	VMAADASRRG RLHAVPSQGL WLLGLRDGKI LEAHVEAKEP RALRTPERPP ARIGLYLSFA
	DGVLAFYDAS NPDVLTPIFS FHERLPGPVY PIFDVCWHDK GKNAQPLLLV GPEQEQA
	Sequence without tag. The proposed Strep-Tag is based on experience s with the expression
	system, a different complexity of the protein could make another tag necessary. In case you
	have a special request, please contact us.

Characteristics:

Key Benefits:

- Made in Germany from design to production by highly experienced protein experts.
- · Protein expressed with ALiCE® and purified in one-step affinity chromatography
- 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 try to ensure that you receive soluble 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 posttranslational 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 against its specific reference buffer.
- We use the Expasy's ProtParam tool to determine the absorption coefficient of each protein.

Purification:	One-step Strep-tag purification of proteins expressed in Almost Living Cell-Free Expression System (AliCE®).
Purity:	> 70-80 % as determined by SDS PAGE, Western Blot and analytical SEC (HPLC).
Grade:	custom-made

Target Details

Target:	TRIM72
Alternative Name:	Trim72 (TRIM72 Products)
Background:	Tripartite motif-containing protein 72 (Mitsugumin-53) (Mg53),FUNCTION: Muscle-specific
	protein that plays a central role in cell membrane repair by nucleating the assembly of the repair
	machinery at injury sites. Specifically binds phosphatidylserine. Acts as a sensor of oxidation:
	upon membrane damage, entry of extracellular oxidative environment results in disulfide bond
	formation and homooligomerization at the injury site. This oligomerization acts as a nucleation
	site for recruitment of TRIM72-containing vesicles to the injury site, leading to membrane patch
	formation. Probably acts upstream of the Ca(2+)-dependent membrane resealing process.
	Required for transport of DYSF to sites of cell injury during repair patch formation. Regulates
	membrane budding and exocytosis. May be involved in the regulation of the mobility of KCNB1
	containing endocytic vesicles. {ECO:0000269 PubMed:19029292,
	ECO:0000269 PubMed:19043407, ECO:0000269 PubMed:19202355,
	ECO:0000269 PubMed:19380584}.
Molecular Weight:	52.8 kDa
UniProt:	Q1XH17
Application Dataila	
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
	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. Standard Storage Buffer: PBS pH 7.4, 10 % Glycerol Might differ depending on protein.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C
Storage Comment:	Store at -80°C.
Expiry Date:	12 months