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MAPK15 Protein (AA 1-544) (Strep Tag)





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Quantity:	1 mg
Target:	MAPK15
Protein Characteristics:	AA 1-544
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This MAPK15 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Sequence:

MCTVVDPRIV RRYLLRRQLG QGAYGIVWKA VDRRTGEVVA IKKIFDAFRD KTDAQRTFRE
ITLLQEFGDH PNIISLLDVI RAENDRDIYL VFEFMDTDLN AVIRKGGLLQ DVHVRSIFYQ
LLRATRFLHS GHVVHRDQKP SNVLLDANCT VKLCDFGLAR SLGDLPEGPE DQAVTEYVAT
RWYRAPEVLL SSHRYTLGVD MWSLGCILGE MLRGRPLFPG TSTLHQLELI LETIPPPSEE
DLLALGSGCR ASVLHQLGSR PRQTLDALLP PDTSPEALDL LRRLLVFAPD KRLSATQALQ
HPYVQRFHCP SDEWAREADV RPRAHEGVQL SVPEYRSRVY QMILECGGSS GTSREKGPEG
VSPSQAHLHK PRADPQLPSR TPVQGPRPRP QSSPGHDPAE HESPRAAKNV PRQNSAPLLQ
TALLGNGERP PGAKEAPPLT LSLVKPSGRG AAPSLTSQAA AQVANQALIR GDWNRGGGVR
VASVQQVPPR LPPEARPGRR MFSTSALQGA QGGARALLGG YSQAYGTVCH SALGHLPLLE GHHV

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 by multi-step, protein-specific process to ensure correct folding and modification.
- 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 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 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®):

1. 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: >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)

Grade: Crystallography grade

Target Details

Target: MAPK15

Alternative Name: MAPK15 (MAPK15 Products)

Background: Mitogen-activated protein kinase 15 (MAP kinase 15) (MAPK 15) (EC 2.7.11.24) (Extracellular

signal-regulated kinase 7) (ERK-7) (Extracellular signal-regulated kinase 8) (ERK-8),FUNCTION: Atypical MAPK protein that regulates several process such as autophagy, ciliogenesis, protein trafficking/secretion and genome integrity, in a kinase activity-dependent manner (PubMed:22948227, PubMed:24618899, PubMed:29021280, PubMed:21847093,

interaction with GABARAP, MAP1LC3B and GABARAPL1 leading to autophagosome formation, SQSTM1 degradation and reduced MAP1LC3B inhibitory phosphorylation (PubMed:22948227).

PubMed:20733054). Controls both, basal and starvation-induced autophagy throught its

Regulates primary cilium formation and the localization of ciliary proteins involved in cilium

structure, transport, and signaling (PubMed:29021280). Prevents the relocation of the sugaradding enzymes from the Golgi to the endoplasmic reticulum, thereby restricting the production

of sugar-coated proteins (PubMed:24618899). Upon amino-acid starvation, mediates

transitional endoplasmic reticulum site disassembly and inhibition of secretion

(PubMed:21847093). Binds to chromatin leading to MAPK15 activation and interaction with

PCNA, that which protects genomic integrity by inhibiting MDM2-mediated degradation of

PCNA (PubMed:20733054). Regulates DA transporter (DAT) activity and protein expression via

activation of RhoA (PubMed:28842414). In response to H(2)O(2) treatment phosphorylates ELAVL1, thus preventing it from binding to the PDCD4 3'UTR and rendering the PDCD4 mRNA

accessible to miR-21 and leading to its degradation and loss of protein expression

(PubMed:26595526). Also functions in a kinase activity-independent manner as a negative

regulator of growth (By similarity). Phosphorylates in vitro FOS and MBP (PubMed:11875070,

PubMed:16484222, PubMed:20638370, PubMed:19166846). During oocyte maturation, plays a

key role in the microtubule organization and meiotic cell cycle progression in oocytes, fertilized

eggs, and early embryos (By similarity). Interacts with ESRRA promoting its re-localization from

Buffer:

Handling Advice:

D. ((
Format:	Liquid
Handling	
Restrictions:	For Research Use only
	needed is the DNA that codes for the desired protein!
	something that functions like a cell, but without the constraints of a living system - all that's
	components needed for protein production (amino acids, cofactors, etc.) are added to produce
	mitochondria to drive the reaction. During our lysate completion steps, the additional
	protein production are removed, leaving only the protein production machinery and the
	During lysate production, the cell wall and other cellular components that are not required for
	modifications.
	even the most difficult-to-express proteins, including those that require post-translational
	Nicotiana tabacum c.v This contains all the protein expression machinery needed to produce
Comment:	ALiCE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from
	guarantee though.
•	as well. As the protein has not been tested for functional studies yet we cannot offer a
Application Notes:	In addition to the applications listed above we expect the protein to work for functional studies
Application Details	
UniProt:	Q8TD08
Molecular Weight:	59.8 kDa
	ECO:0000269 PubMed:28842414, ECO:0000269 PubMed:29021280}.
	ECO:0000269 PubMed:24618899, ECO:0000269 PubMed:26595526,
	ECO:0000269 PubMed:21847093, ECO:0000269 PubMed:22948227,
	ECO:0000269 PubMed:20733054, ECO:0000269 PubMed:21190936,
	ECO:0000269 PubMed:19166846, ECO:0000269 PubMed:20638370,
	ECO:0000269 PubMed:11875070, ECO:0000269 PubMed:16484222,
	{ECO:0000250 UniProtKB:Q80Y86, ECO:0000250 UniProtKB:Q9Z2A6,
	the nucleus to the cytoplasm and then prevents its transcriptional activity (PubMed:21190936)

please contact us.

Avoid repeated freeze-thaw cycles.

The buffer composition is at the discretion of the manufacturer. If you have a special request,

Handling

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

Images

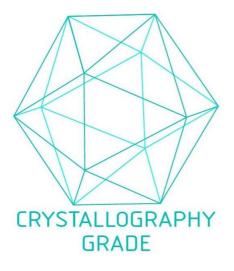


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process