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EPM2A Protein (AA 1-331) (Strep Tag)





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Overview

Quantity:	1 mg
Target:	EPM2A
Protein Characteristics:	AA 1-331
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This EPM2A protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Sequence:

MRFRFGVVVP PAVAGARPEL LVVGSRPELG RWEPRGAVRL RPAGTAAGDG ALALQEPGLW
LGEVELAAEE AAQDGAEPGR VDTFWYKFLK REPGGELSWE GNGPHHDRCC TYNENNLVDG
VYCLPIGHWI EATGHTNEMK HTTDFYFNIA GHQAMHYSRI LPNIWLGSCP RQVEHVTIKL
KHELGITAVM NFQTEWDIVQ NSSGCNRYPE PMTPDTMIKL YREEGLAYIW MPTPDMSTEG
RVQMLPQAVC LLHALLEKGH IVYVHCNAGV GRSTAAVCGW LQYVMGWNLR KVQYFLMAKR
PAVYIDEEAL ARAQEDFFQK FGKVRSSVCS L

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.

Product Details		
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)	
Grade:	Crystallography grade	
Target Details		
Target:	EPM2A	
Alternative Name:	EPM2A (EPM2A Products)	
Background:	Laforin (EC 3.1.3) (EC 3.1.3.16) (EC 3.1.3.48) (Glucan phosphatase) (Glycogen phosphatase)	
	(Lafora PTPase) (LAFPTPase), FUNCTION: Plays an important role in preventing glycogen	
	hyperphosphorylation and the formation of insoluble aggregates, via its activity as glycogen	
	phosphatase, and by promoting the ubiquitination of proteins involved in glycogen metabolism	
	via its interaction with the E3 ubiquitin ligase NHLRC1/malin. Shows strong phosphatase	
	activity towards complex carbohydrates in vitro, avoiding glycogen hyperphosphorylation which	
	is associated with reduced branching and formation of insoluble aggregates	
	(PubMed:16901901, PubMed:23922729, PubMed:26231210, PubMed:25538239,	
	PubMed:25544560). Dephosphorylates phosphotyrosine and synthetic substrates, such as	
	para-nitrophenylphosphate (pNPP), and has low activity with phosphoserine and	
	phosphothreonine substrates (in vitro) (PubMed:11001928, PubMed:11220751,	
	PubMed:11739371, PubMed:14532330, PubMed:16971387, PubMed:18617530,	

PubMed:22036712, PubMed:23922729, PubMed:14722920). Has been shown to dephosphorylate MAPT (By similarity). Forms a complex with NHLRC1/malin and HSP70, which suppresses the cellular toxicity of misfolded proteins by promoting their degradation through the ubiquitin-proteasome system (UPS). Acts as a scaffold protein to facilitate PPP1R3C/PTG ubiquitination by NHLRC1/malin (PubMed:23922729). Also promotes proteasome-independent protein degradation through the macroautophagy pathway (PubMed:20453062). {ECO:0000250|UniProtKB:Q9WUA5, ECO:0000269|PubMed:11001928, ECO:0000269|PubMed:11220751, ECO:0000269|PubMed:11739371, ECO:0000269|PubMed:14532330, ECO:0000269|PubMed:14722920, ECO:0000269|PubMed:16901901, ECO:0000269|PubMed:16971387, ECO:0000269|PubMed:18070875, ECO:0000269|PubMed:18617530, ECO:0000269|PubMed:19036738, ECO:0000269|PubMed:20453062, ECO:0000269|PubMed:22036712, ECO:0000269|PubMed:23624058, ECO:0000269|PubMed:23922729, ECO:0000269|PubMed:25538239, ECO:0000269|PubMed:25544560, ECO:0000269|PubMed:26231210}., FUNCTION: [Isoform 2]: Does not bind to glycogen (PubMed:18617530). Lacks phosphatase activity and might function

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	as a dominant-negative regulator for the phosphatase activity of isoform 1 and isoform 7 (PubMed:18617530, PubMed:22036712). {ECO:0000269 PubMed:18617530, ECO:0000269 PubMed:22036712}., FUNCTION: [Isoform 7]: Has phosphatase activity (in vitro). {ECO:0000269 PubMed:22036712}.
Molecular Weight:	37.2 kDa
UniProt:	095278
Pathways:	Cellular Glucan Metabolic Process
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. If you have a special request, please contact us.
Handling Advice:	Avoid repeated freeze-thaw cycles.
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