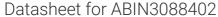
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PRKAG3 Protein (AA 1-489) (Strep Tag)



Image



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Overview

Quantity:	1 mg
Target:	PRKAG3
Protein Characteristics:	AA 1-489
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This PRKAG3 protein is labelled with Strep Tag.
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA

Product Details

Sequence:

MEPGLEHALR RTPSWSSLGG SEHQEMSFLE QENSSSWPSP AVTSSSERIR GKRRAKALRW

TRQKSVEEGE PPGQGEGPRS RPAAESTGLE ATFPKTTPLA QADPAGVGTP PTGWDCLPSD

CTASAAGSST DDVELATEFP ATEAWECELE GLLEERPALC LSPQAPFPKL GWDDELRKPG

AQIYMRFMQE HTCYDAMATS SKLVIFDTML EIKKAFFALV ANGVRAAPLW DSKKQSFVGM

LTITDFILVL HRYYRSPLVQ IYEIEQHKIE TWREIYLQGC FKPLVSISPN DSLFEAVYTL IKNRIHRLPV

LDPVSGNVLH ILTHKRLLKF LHIFGSLLPR PSFLYRTIQD LGIGTFRDLA VVLETAPILT

ALDIFVDRRV SALPVVNECG QVVGLYSRFD VIHLAAQQTY NHLDMSVGEA LRQRTLCLEG

VLSCQPHESL GEVIDRIARE QVHRLVLVDE TQHLLGVVSL SDILQALVLS PAGIDALGA

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.

Product Details

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

- arget Details		
Target:	PRKAG3	
Alternative Name:	PRKAG3 (PRKAG3 Products)	
Background:	5'-AMP-activated protein kinase subunit gamma-3 (AMPK gamma3) (AMPK subunit gamma-	
	3),FUNCTION: AMP/ATP-binding subunit of AMP-activated protein kinase (AMPK), an energy	
	sensor protein kinase that plays a key role in regulating cellular energy metabolism	
	(PubMed:14722619, PubMed:24563466, PubMed:17878938). In response to reduction of	
	intracellular ATP levels, AMPK activates energy-producing pathways and inhibits energy-	
	consuming processes: inhibits protein, carbohydrate and lipid biosynthesis, as well as cell	
	growth and proliferation. AMPK acts via direct phosphorylation of metabolic enzymes, and by	
	longer-term effects via phosphorylation of transcription regulators. AMPK also acts as a	
	regulator of cellular polarity by remodeling the actin cytoskeleton, probably by indirectly	
	activating myosin. The AMPK gamma3 subunit is a non-catalytic subunit with a regulatory role	
	in muscle energy metabolism (PubMed:17878938). It mediates binding to AMP, ADP and ATP,	
	leading to AMPK activation or inhibition: AMP-binding results in allosteric activation of alpha	
	catalytic subunit (PRKAA1 or PRKAA2) both by inducing phosphorylation and preventing	
	dephosphorylation of catalytic subunits. ADP also stimulates phosphorylation, without	
	stimulating already phosphorylated catalytic subunit. ATP promotes dephosphorylation of	
	catalytic subunit, rendering the AMPK enzyme inactive. {ECO:0000269 PubMed:14722619,	
	ECO:0000269 PubMed:17878938, ECO:0000269 PubMed:24563466}.	
Molecular Weight:	54.3 kDa	
UniProt:	Q9UGI9	
Pathways:	AMPK Signaling, Cellular Glucan Metabolic Process, Warburg Effect	

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.

Application Details

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

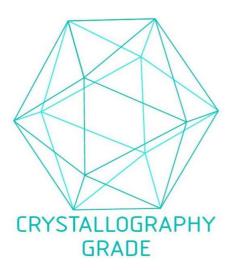


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process