antibodies

Datasheet for ABIN3093339 CAMK1 Protein (AA 1-370) (Strep Tag)





Overview

	1
Quantity:	1 mg
Target:	CAMK1
Protein Characteristics:	AA 1-370
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This CAMK1 protein is labelled with Strep Tag.
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA

Product Details

Sequence:	MLGAVEGPRW KQAEDIRDIY DFRDVLGTGA FSEVILAEDK RTQKLVAIKC IAKEALEGKE
	GSMENEIAVL HKIKHPNIVA LDDIYESGGH LYLIMQLVSG GELFDRIVEK GFYTERDASR
	LIFQVLDAVK YLHDLGIVHR DLKPENLLYY SLDEDSKIMI SDFGLSKMED PGSVLSTACG
	TPGYVAPEVL AQKPYSKAVD CWSIGVIAYI LLCGYPPFYD ENDAKLFEQI LKAEYEFDSP
	YWDDISDSAK DFIRHLMEKD PEKRFTCEQA LQHPWIAGDT ALDKNIHQSV SEQIKKNFAK
	SKWKQAFNAT AVVRHMRKLQ LGTSQEGQGQ TASHGELLTP VAGGPAAGCC CRDCCVEPGT
	ELSPTLPHQL
	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:

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- · 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.

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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

Target:	CAMK1
Alternative Name:	CAMK1 (CAMK1 Products)
Background:	Calcium/calmodulin-dependent protein kinase type 1 (EC 2.7.11.17) (CaM kinase I) (CaM-KI)
	(CaM kinase I alpha) (CaMKI-alpha),FUNCTION: Calcium/calmodulin-dependent protein kinase
	that operates in the calcium-triggered CaMKK-CaMK1 signaling cascade and, upon calcium
	influx, regulates transcription activators activity, cell cycle, hormone production, cell
	differentiation, actin filament organization and neurite outgrowth. Recognizes the substrate
	consensus sequence [MVLIF]-x-R-x(2)-[ST]-x(3)-[MVLIF]. Regulates axonal extension and growt
	cone motility in hippocampal and cerebellar nerve cells. Upon NMDA receptor-mediated Ca(2+)
	elevation, promotes dendritic growth in hippocampal neurons and is essential in synapses for
	full long-term potentiation (LTP) and ERK2-dependent translational activation. Downstream of
	NMDA receptors, promotes the formation of spines and synapses in hippocampal neurons by
	phosphorylating ARHGEF7/BETAPIX on 'Ser-694', which results in the enhancement of
	ARHGEF7 activity and activation of RAC1. Promotes neuronal differentiation and neurite
	outgrowth by activation and phosphorylation of MARK2 on 'Ser-91', 'Ser-92', 'Ser-93' and 'Ser-
	294'. Promotes nuclear export of HDAC5 and binding to 14-3-3 by phosphorylation of 'Ser-259'
	and 'Ser-498' in the regulation of muscle cell differentiation. Regulates NUMB-mediated
	endocytosis by phosphorylation of NUMB on 'Ser-276' and 'Ser-295'. Involved in the regulation
	of basal and estrogen-stimulated migration of medulloblastoma cells through
	ARHGEF7/BETAPIX phosphorylation (By similarity). Is required for proper activation of cyclin-
	D1/CDK4 complex during G1 progression in diploid fibroblasts. Plays a role in K(+) and ANG2-
	mediated regulation of the aldosterone synthase (CYP11B2) to produce aldosterone in the
	adrenal cortex. Phosphorylates EIF4G3/eIF4GII. In vitro phosphorylates CREB1, ATF1, CFTR,
	MYL9 and SYN1/synapsin I. {ECO:0000250, ECO:0000269 PubMed:11114197,
	ECO:0000269 PubMed:12193581, ECO:0000269 PubMed:14507913,
	ECO:0000269 PubMed:14754892, ECO:0000269 PubMed:17056143,
	EC0:0000269 PubMed:17442826, EC0:0000269 PubMed:18184567,
	EC0:0000269 PubMed:20181577}.

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Target Details	
Molecular Weight:	41.3 kDa
UniProt:	Q14012
Pathways:	Myometrial Relaxation and Contraction, Regulation of Muscle Cell Differentiation, Smooth Muscle Cell Migration
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)

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

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