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CAMKII gamma Protein (AA 1-558) (Strep Tag)



Image



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Overview

Quantity:	1 mg
Target:	CAMKII gamma (CAMK2G)
Protein Characteristics:	AA 1-558
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This CAMKII gamma protein is labelled with Strep Tag.
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA

Product Details

Sequence:

MATTATCTRF TDDYQLFEEL GKGAFSVVRR CVKKTSTQEY AAKIINTKKL SARDHQKLER EARICRLLKH PNIVRLHDSI SEEGFHYLVF DLVTGGELFE DIVAREYYSE ADASHCIHQI LESVNHIHQH DIVHRDLKPE NLLLASKCKG AAVKLADFGL AIEVQGEQQA WFGFAGTPGY LSPEVLRKDP YGKPVDIWAC GVILYILLVG YPPFWDEDQH KLYQQIKAGA YDFPSPEWDT VTPEAKNLIN QMLTINPAKR ITADQALKHP WVCQRSTVAS MMHRQETVEC LRKFNARRKL KGAILTTMLV SRNFSAAKSL LNKKSDGGVK KRKSSSSVHL MPQSNNKNSL VSPAQEPAPL QTAMEPQTTV VHNATDGIKG STESCNTTTE DEDLKGRVPE GRSSRDRTAP SAGMQPQPSL CSSAMRKQEI IKITEQLIEA INNGDFEAYT KICDPGLTSF EPEALGNLVE GMDFHKFYFE NLLSKNSKPI HTTILNPHVH VIGEDAACIA YIRLTQYIDG QGRPRTSQSE ETRVWHRRDG KWLNVHYHCS GAPAAPLQ

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.

Application Notes:	In addition to the applications listed above we expect the protein to work for functional studies
Application Details	
Pathways:	WNT Signaling, Interferon-gamma Pathway, Hormone Transport, Myometrial Relaxation and Contraction, Regulation of long-term Neuronal Synaptic Plasticity
UniProt:	Q13555 WNT Signaling Interferon gamma Dathway Harmona Transport Myometrial Polavation and
Molecular Weight:	
Molocular Weight	62.6 kDa
	signaling pathway (By similarity). {ECO:0000250 UniProtKB:Q923T9, ECO:0000269 PubMed:16690701, ECO:0000269 PubMed:30184290}.
	(IFN-gamma) stimulation, catalyzes phosphorylation of STAT1, stimulating the JAK-STAT
	potentiation (LTP) and hippocampus-dependent learning. In response to interferon-gamma
	spine and synapse formation and maintenance of synaptic plasticity which enables long-term
	formation and arborization (PubMed:30184290). It may participate in the promotion of dendrit
Buonground.	(PubMed:16690701). In the central nervous system, it is involved in the regulation of neurite
	release from the SR through phosphorylation of the ryanodine receptor-coupling factor triadin
	reticulum (SR) Ca(2+) transport and in fast-twitch muscle participates in the control of Ca(2+)
	(PubMed:16690701). In slow-twitch muscles, is involved in regulation of sarcoplasmic
	muscle and may function in dendritic spine and synapse formation and neuronal plasticity
	autophosphorylation, and is involved in sarcoplasmic reticulum Ca(2+) transport in skeletal
	protein kinase that functions autonomously after Ca(2+)/calmodulin-binding and
	gamma) (CaMK-II subunit gamma) (EC 2.7.11.17), FUNCTION: Calcium/calmodulin-dependent
Background:	Calcium/calmodulin-dependent protein kinase type II subunit gamma (CaM kinase II subunit
Alternative Name:	CAMK2G (CAMK2G Products)
Target:	CAMKII gamma (CAMK2G)
Target Details	
Grade:	Crystallography grade
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)
Purity:	>80 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.
	through size exclusion chromatography. Eluate fractions are analyzed by SDS-PAGE and Western blot.

Application Details

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	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)



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