antibodies

# Datasheet for ABIN3084529 NMNAT2 Protein (AA 1-307) (Strep Tag)





### Overview

Quantity:	1 mg
Target:	NMNAT2
Protein Characteristics:	AA 1-307
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This NMNAT2 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

### Product Details

Sequence:	MTETTKTHVI LLACGSFNPI TKGHIQMFER ARDYLHKTGR FIVIGGIVSP VHDSYGKQGL
	VSSRHRLIMC QLAVQNSDWI RVDPWECYQD TWQTTCSVLE HHRDLMKRVT GCILSNVNTP
	SMTPVIGQPQ NETPQPIYQN SNVATKPTAA KILGKVGESL SRICCVRPPV ERFTFVDENA
	NLGTVMRYEE IELRILLLCG SDLLESFCIP GLWNEADMEV IVGDFGIVVV PRDAADTDRI
	MNHSSILRKY KNNIMVVKDD INHPMSVVSS TKSRLALQHG DGHVVDYLSQ PVIDYILKSQ
	LYINASG
	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

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- 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 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!

#### 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®):
	<ol> <li>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.</li> <li>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.</li> </ol>
Purity:	>80 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.

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Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)
Grade:	Crystallography grade
Target Details	
Target:	NMNAT2
Alternative Name:	NMNAT2 (NMNAT2 Products)
Background:	Nicotinamide/nicotinic acid mononucleotide adenylyltransferase 2 (NMN/NaMN
	adenylyltransferase 2) (EC 2.7.7.1) (EC 2.7.7.18) (Nicotinamide mononucleotide
	adenylyltransferase 2) (NMN adenylyltransferase 2) (Nicotinate-nucleotide adenylyltransferase
	2) (NaMN adenylyltransferase 2),FUNCTION: Nicotinamide/nicotinate-nucleotide
	adenylyltransferase that acts as an axon maintenance factor (By similarity). Axon survival
	factor required for the maintenance of healthy axons: acts by delaying Wallerian axon
	degeneration, an evolutionarily conserved process that drives the loss of damaged axons (By
	similarity). Catalyzes the formation of NAD(+) from nicotinamide mononucleotide (NMN) and
	ATP (PubMed:16118205, PubMed:17402747). Can also use the deamidated form, nicotinic ac
	mononucleotide (NaMN) as substrate but with a lower efficiency (PubMed:16118205,
	PubMed:17402747). Cannot use triazofurin monophosphate (TrMP) as substrate
	(PubMed:16118205, PubMed:17402747). Also catalyzes the reverse reaction, i.e. the
	pyrophosphorolytic cleavage of NAD(+) (PubMed:16118205, PubMed:17402747). For the
	pyrophosphorolytic activity prefers NAD(+), NADH and NaAD as substrates and degrades
	nicotinic acid adenine dinucleotide phosphate (NHD) less effectively (PubMed:16118205,
	PubMed:17402747). Fails to cleave phosphorylated dinucleotides NADP(+), NADPH and
	NaADP(+) (PubMed:16118205, PubMed:17402747). Also acts as an activator of ADP-
	ribosylation by supporting the catalytic activity of PARP16 and promoting mono-ADP-
	ribosylation of ribosomes by PARP16 (PubMed:34314702). {ECO:0000250 UniProtKB:Q8BNJ3
	EC0:0000269 PubMed:16118205, EC0:0000269 PubMed:17402747,

Molecular Weight: 34.4 kDa

UniProt:

Q9BZQ4

ECO:0000269|PubMed:34314702}.

## 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

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Application Details	
	guarantee though.
Comment:	<ul> <li>ALiCE®, our Almost Living Cell-Free Expression System is based on a lysate obtained from</li> <li>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.</li> <li>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!</li> </ul>
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

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