antibodies

# Datasheet for ABIN3088362 ADAL Protein (AA 1-355) (Strep Tag)





### Overview

Quantity:	1 mg
Target:	ADAL
Protein Characteristics:	AA 1-355
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This ADAL protein is labelled with Strep Tag.
Application:	ELISA, SDS-PAGE (SDS), Western Blotting (WB)

## Product Details

Sequence:	MIEAEEQQPC KTDFYSELPK VELHAHLNGS ISSHTMKKLI AQKPDLKIHD QMTVIDKGKK
	RTLEECFQMF QTIHQLTSSP EDILMVTKDV IKEFADDGVK YLELRSTPRR ENATGMTKKT
	YVESILEGIK QSKQENLDID VRYLIAVDRR GGPLVAKETV KLAEEFFLST EGTVLGLDLS
	GDPTVGQAKD FLEPLLEAKK AGLKLALHLS EIPNQKKETQ ILLDLLPDRI GHGTFLNSGE
	GGSLDLVDFV RQHRIPLELC LTSNVKSQTV PSYDQHHFGF WYSIAHPSVI CTDDKGVFAT
	HLSQEYQLAA ETFNLTQSQV WDLSYESINY IFASDSTRSE LRKKWNHLKP RVLHI
	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> </ol>
	2. 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.

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Product Details	
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)
Grade:	Crystallography grade

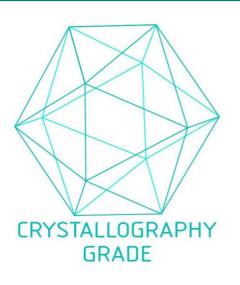
## Target Details

Target:	ADAL
Alternative Name:	ADAL (ADAL Products)
Background:	Adenosine deaminase-like protein (EC 3.5.4) (Adenosine deaminase-like protein isoform 1)
	(N6-mAMP deaminase) (HsMAPDA) (N6-methyl-AMP aminohydrolase),FUNCTION: Catalyzes
	the hydrolysis of the free cytosolic methylated adenosine nucleotide N(6)-methyl-AMP (N6-
	mAMP) to produce inositol monophosphate (IMP) and methylamine (PubMed:21755941,
	PubMed:29884623). Is required for the catabolism of cytosolic N6-mAMP, which is derived
	from the degradation of mRNA containing N6-methylated adenine (m6A) (PubMed:21755941,
	PubMed:29884623). Catalyzes the removal of different alkyl groups not only from N6-
	substituted purine or 2-aminopurine nucleoside monophosphates but also from O6-substituted
	compounds in vitro (PubMed:21755941). {ECO:0000269 PubMed:21755941,
	ECO:0000269 PubMed:29884623}.
Molecular Weight:	40.3 kDa
UniProt:	Q6DHV7
Pathways:	Ribonucleoside Biosynthetic 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

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Application Details	
	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

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