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Datasheet for ABIN3134814 ATP5A1 Protein (AA 44-553) (His tag)



Overview

Quantity:	1 mg
Target:	ATP5A1
Protein Characteristics:	AA 44-553
Origin:	Mouse
Source:	Escherichia coli (E. coli)
Protein Type:	Recombinant
Purification tag / Conjugate:	This ATP5A1 protein is labelled with His tag.
Application:	ELISA, SDS-PAGE (SDS), Western Blotting (WB), Crystallization (Crys)

Product Details

Sequence:	QKTGTAEMSS ILEERILGAD TSVDLEETGR VLSIGDGIAR VHGLRNVQAE EMVEFSSGLK
	GMSLNLEPDN VGVVVFGNDK LIKEGDVVKR TGAIVDVPVG EELLGRVVDA LGNAIDGKGP
	IGSKTRRRVG LKAPGIIPRI SVREPMQTGI KAVDSLVPIG RGQRELIIGD RQTGKTSIAI DTIINQKRFN
	DGTDEKKKLY CIYVAIGQKR STVAQLVKRL TDADAMKYTI VVSATASDAA PLQYLAPYSG
	CSMGEYFRDN GKHALIIYDD LSKQAVAYRQ MSLLLRRPPG REAYPGDVFY LHSRLLERAA
	KMNDSFGGGS LTALPVIETQ AGDVSAYIPT NVISITDGQI FLETELFYKG IRPAINVGLS
	VSRVGSAAQT RAMKQVAGTM KLELAQYREV AAFAQFGSDL DAATQQLLSR GVRLTELLKQ
	GQYSPMAIEE QVAVIYAGVR GYLDKLEPSK ITKFENAFLS HVISQHQSLL GNIRSDGKIS
	EQSDAKLKEI VTNFLAGFEP
	Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a
	special request, please contact us.

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 Characteristics: Made in Germany - from design to production - by highly experienced protein exp Mouse Atp5a1 Protein (raised in E. Coli) purified by multi-step, protein-specific pr ensure crystallization grade. 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 o	der. 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 orderin	g custom
made proteins from other companies is that there is no financial obligation in case	the protein
cannot be expressed or purified.	
In the unlikely event that the protein cannot be expressed or purified we do not char	ge anything
(other companies might charge you for any performed steps in the expression proc	ess for
custom-made proteins, e.g. fees might apply for the expression plasmid, the first ex	pression
experiments or purification optimization).	
When you order this made-to-order protein you will only pay upon receival of the co	rrectly
folded protein. With no financial risk on your end you can rest assured that our expe	rienced
protein experts will do everything to make sure that you receive the protein you orde	ered.
The concentration of our recombinant proteins is measured using the absorbance a	at 280nm.
The protein's absorbance will be measured in several dilutions and is measured aga	ainst its
specific reference buffer.	
The concentration of the protein is calculated using its specific absorption coefficie	nt. We use
the Expasy's protparam tool to determine the absorption coefficient of each protein	I.
Purification: Two step purification of proteins expressed in bacterial culture:	
1. In a first purification step, the protein is purified from the cleared cell lysate using different His-tag capture materials: high yield, EDTA resistant, or DTT resistant. E fractions are analyzed by SDS-PAGE.	luate
 Protein containing fractions of the best purification are subjected to second purif through size exclusion chromatography. Eluate fractions are analyzed by SDS-PA Western blot. 	•
Purity: >95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western B	lot.
Sterility: 0.22 µm filtered	
Endotoxin Level: Endotoxin has not been removed. Please contact us if you require endotoxin remov	al.
Grade: Crystallography grade	

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Target Details

Target:	ATP5A1
Alternative Name:	Atp5a1 (ATP5A1 Products)
Background:	Mitochondrial membrane ATP synthase (F(1)F(0) ATP synthase or Complex V) produces ATP
	from ADP in the presence of a proton gradient across the membrane which is generated by
	electron transport complexes of the respiratory chain. F-type ATPases consist of two structural
	domains, $F(1)$ - containing the extramembraneous catalytic core, and $F(0)$ - containing the
	membrane proton channel, linked together by a central stalk and a peripheral stalk. During
	catalysis, ATP synthesis in the catalytic domain of $F(1)$ is coupled via a rotary mechanism of
	the central stalk subunits to proton translocation. Subunits alpha and beta form the catalytic
	core in F(1). Rotation of the central stalk against the surrounding alpha(3)beta(3) subunits leads
	to hydrolysis of ATP in three separate catalytic sites on the beta subunits. Subunit alpha does
	not bear the catalytic high-affinity ATP-binding sites (By similarity). {ECO:0000250}.
Molecular Weight:	56.3 kDa Including tag.
UniProt:	Q03265
Pathways:	Proton Transport, 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 gurantee
	though.
Comment:	Protein has not been tested for activity yet. In cases in which it is highly likely that the
	recombinant protein with the default tag will be insoluble our protein lab may suggest a higher
	molecular weight tag (e.g. GST-tag) instead to increase solubility. We will discuss all possible
	options with you in detail to assure that you receive your protein of interest.
Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	100 mM NaCL, 20 mM Hepes, 10% glycerol. pH value is at the discretion of the manufacturer.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C

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Storage Comment:	Store at -80°C.

Expiry Date:

Unlimited (if stored properly)

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