antibodies

Datasheet for ABIN3077056 SNAIL Protein (SNAI1) (AA 1-264) (Strep Tag)



Overview

Image

Quantity:	1 mg
Target:	SNAIL (SNAI1)
Protein Characteristics:	AA 1-264
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This SNAIL protein is labelled with Strep Tag.
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA

Product Details

Sequence:	MPRSFLVRKP SDPNRKPNYS ELQDSNPEFT FQQPYDQAHL LAAIPPPEIL NPTASLPMLI
	WDSVLAPQAQ PIAWASLRLQ ESPRVAELTS LSDEDSGKGS QPPSPPSPAP SSFSSTSVSS
	LEAEAYAAFP GLGQVPKQLA QLSEAKDLQA RKAFNCKYCN KEYLSLGALK MHIRSHTLPC
	VCGTCGKAFS RPWLLQGHVR THTGEKPFSC PHCSRAFADR SNLRAHLQTH SDVKKYQCQA
	CARTFSRMSL LHKHQESGCS GCPR
	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.
	· Made in demany - norn design to production - by highly experienced protein expense.
	 Protein expressed with ALiCE® and purified by multi-step, protein-specific process to ensure correct folding and modification.

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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®):
	 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.
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)

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Product Details

Grade:

Crystallography grade

Target Details

Target:	SNAIL (SNAI1)
Alternative Name:	SNAI1 (SNAI1 Products)
Background:	Zinc finger protein SNAI1 (Protein snail homolog 1) (Protein sna),FUNCTION: Involved in
	induction of the epithelial to mesenchymal transition (EMT), formation and maintenance of
	embryonic mesoderm, growth arrest, survival and cell migration. Binds to 3 E-boxes of the E-
	cadherin/CDH1 gene promoter and to the promoters of CLDN7 and KRT8 and, in association
	with histone demethylase KDM1A which it recruits to the promoters, causes a decrease in
	dimethylated H3K4 levels and represses transcription (PubMed:20389281, PubMed:20562920
	The N-terminal SNAG domain competes with histone H3 for the same binding site on the
	histone demethylase complex formed by KDM1A and RCOR1, and thereby inhibits
	demethylation of histone H3 at 'Lys-4' (in vitro) (PubMed:20389281, PubMed:21300290,
	PubMed:23721412). During EMT, involved with LOXL2 in negatively regulating pericentromeric
	heterochromatin transcription (By similarity). SNAI1 recruits LOXL2 to pericentromeric regions
	to oxidize histone H3 and repress transcription which leads to release of heterochromatin
	component CBX5/HP1A, enabling chromatin reorganization and acquisition of mesenchymal
	traits (By similarity). Associates with EGR1 and SP1 to mediate tetradecanoyl phorbol acetate
	(TPA)-induced up-regulation of CDKN2B, possibly by binding to the CDKN2B promoter region $\$$
	TCACA-3. In addition, may also activate the CDKN2B promoter by itself.
	{ECO:0000250 UniProtKB:Q02085, ECO:0000269 PubMed:10655587,
	EC0:0000269 PubMed:15647282, EC0:0000269 PubMed:16096638,
	EC0:0000269 PubMed:20121949, EC0:0000269 PubMed:20389281,
	EC0:0000269 PubMed:20562920, EC0:0000269 PubMed:21300290,
	EC0:0000269 PubMed:21952048, EC0:0000269 PubMed:23721412}.
Molecular Weight:	29.1 kDa
UniProt:	095863
Pathways:	Negative Regulation of intrinsic apoptotic Signaling
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 Detail	S
	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

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