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Datasheet for ABIN3113129 FUT3 Protein (AA 1-361) (Strep Tag)





Overview

Quantity:	1 mg
Target:	FUT3
Protein Characteristics:	AA 1-361
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This FUT3 protein is labelled with Strep Tag.
Application:	Western Blotting (WB), SDS-PAGE (SDS), ELISA

Product Details

Sequence:	MDPLGAAKPQ WPWRRCLAAL LFQLLVAVCF FSYLRVSRDD ATGSPRAPSG SSRQDTTPTR
	PTLLILLRTW PFHIPVALSR CSEMVPGTAD CHITADRKVY PQADMVIVHH WDIMSNPKSR
	LPPSPRPQGQ RWIWFNLEPP PNCQHLEALD RYFNLTMSYR SDSDIFTPYG WLEPWSGQPA
	HPPLNLSAKT ELVAWAVSNW KPDSARVRYY QSLQAHLKVD VYGRSHKPLP KGTMMETLSR
	YKFYLAFENS LHPDYITEKL WRNALEAWAV PVVLGPSRSN YERFLPPDAF IHVDDFQSPK
	DLARYLQELD KDHARYLSYF RWRETLRPRS FSWALDFCKA CWKLQQESRY QTVRSIAAWF T
	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®):
	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.
	 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 De	etails
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Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)
Grade:	Crystallography grade

Target Details

Target:	FUT3
Alternative Name:	FUT3 (FUT3 Products)
Background:	3-galactosyl-N-acetylglucosaminide 4-alpha-L-fucosyltransferase FUT3 (EC 2.4.1.65) (4-
	galactosyl-N-acetylglucosaminide 3-alpha-L-fucosyltransferase) (EC 2.4.1.152) (Alpha-3-
	fucosyltransferase FUT3) (EC 2.4.1) (Blood group Lewis alpha-4-fucosyltransferase) (Lewis
	FT) (Fucosyltransferase 3) (Fucosyltransferase III) (FucT-III),FUNCTION: Catalyzes the transfe
	of L-fucose, from a guanosine diphosphate-beta-L-fucose, to both the subterminal N-acetyl
	glucosamine (GlcNAc) of type 1 chain (beta-D-Gal-(1->3)-beta-D-GlcNAc) glycolipids and
	oligosaccharides via an alpha(1,4) linkage, and the subterminal glucose (Glc) or GlcNAc of typ
	2 chain (beta-D-Gal-(1->4)-beta-D-GlcNAc) oligosaccharides via an alpha(1,3) linkage,
	independently of the presence of terminal alpha-L-fucosyl-(1,2) moieties on the terminal
	galactose of these acceptors (PubMed:12668675, PubMed:1977660, PubMed:11058871).
	Through its catalytic activity, participates in the synthesis of antigens of the Lewis blood group
	system, i.e. Lewis a (Le(a)), lewis b (Le(b)), Lewis x/SSEA-1 (Le(x)) and lewis y (Le(y)) antigens
	(PubMed:12668675, PubMed:1977660, PubMed:11058871). Also catalyzes the transfer of L-
	fucose to subterminal GlcNAc of sialyl- and disialyl-lactotetraosylceramide to produce sialyl
	Lewis a (sLe(a)) and disialyl Lewis a via an alpha(1,4) linkage and therefore may regulate cell
	surface sLe(a) expression and consequently regulates adhesive properties to E-selectin, cell
	proliferation and migration (PubMed:12668675, PubMed:11058871, PubMed:27453266).
	Catalyzes the transfer of an L-fucose to 3'-sialyl-N-acetyllactosamine by an alpha(1,3) linkage,
	which allows the formation of sialyl-Lewis x structure and therefore may regulate the sialyl-
	Lewis x surface antigen expression and consequently adhesive properties to E-selectin
	(PubMed:11058871, PubMed:29593094). Prefers type 1 chain over type 2 acceptors
	(PubMed:7721776). Type 1 tetrasaccharide is a better acceptor than type 1 disaccharide
	suggesting that a beta anomeric configuration of GlcNAc in the substrate is preferred
	(PubMed:7721776). Lewis-positive (Le(+)) individuals have an active enzyme while Lewis-
	negative (Le(-)) individuals have an inactive enzyme (PubMed:1977660).
	{ECO:0000269 PubMed:11058871, ECO:0000269 PubMed:12668675,
	ECO:0000269 PubMed:1977660, ECO:0000269 PubMed:27453266,
	EC0:000269 PubMed:7721776}.

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Target Details	
Molecular Weight:	42.1 kDa
UniProt:	P21217
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 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

Expiry Date:

Storage Comment:

Store at -80°C.

Unlimited (if stored properly)

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

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