

Datasheet for ABIN3136242

TSEN34 Protein (AA 1-316) (Strep Tag)



_					
	W	0	rv	10	W

Quantity:	250 μg
Target:	TSEN34
Protein Characteristics:	AA 1-316
Origin:	Mouse
Source:	Cell-free protein synthesis (CFPS)
Protein Type:	Recombinant
Purification tag / Conjugate:	This TSEN34 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

T TOUUCE Details	
Brand:	AliCE®
Sequence:	MLVVEVANGR SLVWGAEAVQ ALRERLGVGG RTVGALPRGP RQNSRLGLPL LLLPEEARLL
	AEIGAVTLVS APRPDPRNHG LALASFKRQQ EQSFQDQNTL AAEARETRRQ ELLEKIVEGQ
	AAKKQKLEQD SGADEGGQEA GGSEATQGSE TSDDGQPSAE QEGAAPSLDS SSPQPGPSNG
	VTPLPRSALL IQLATARPRP VKAKPLDWRV QSKDWPHAGR PAHELRYSIY RDLWERGFFL
	SAAGKFGGDF LVYPGDPLRF HAHYIAQCWS AEDPIPLQDL VSAGRLGTSV RKTLLLCSPQ
	PDGKVVYTSL QWASLQ
	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 in one-step affinity chromatography
- 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 try to ensure that you receive soluble 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 posttranslational 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 against its specific reference buffer.
- · We use the Expasy's ProtParam tool to determine the absorption coefficient of each protein.

Purification:	One-step Strep-tag purification of proteins expressed in Almost Living Cell-Free Expression System (AliCE®).
Purity:	> 70-80 % as determined by SDS PAGE, Western Blot and analytical SEC (HPLC).
Grade:	custom-made
Target Details	
Target:	TSEN34

Target Details

Alternative Name:	Tsen34 (TSEN34 Products)
Background:	TRNA-splicing endonuclease subunit Sen34 (EC 4.6.1.16) (Leukocyte receptor cluster member
	5 homolog) (tRNA-intron endonuclease Sen34),FUNCTION: Constitutes one of the two catalytic
	subunit of the tRNA-splicing endonuclease complex, a complex responsible for identification
	and cleavage of the splice sites in pre-tRNA. It cleaves pre-tRNA at the 5'- and 3'-splice sites to
	release the intron. The products are an intron and two tRNA half-molecules bearing 2',3'-cyclic
	phosphate and 5'-OH termini. There are no conserved sequences at the splice sites, but the
	intron is invariably located at the same site in the gene, placing the splice sites an invariant
	distance from the constant structural features of the tRNA body. The tRNA splicing
	endonuclease is also involved in mRNA processing via its association with pre-mRNA 3'-end
	processing factors, establishing a link between pre-tRNA splicing and pre-mRNA 3'-end
	formation, suggesting that the endonuclease subunits function in multiple RNA-processing
	events (By similarity). {ECO:0000250}.
Molecular Weight:	34.2 kDa
UniProt:	Q8BMZ5
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. Standard Storage Buffer: PBS pH 7.4, 10 % Glycerol Might differ depending on protein.	
Handling Advice:	Avoid repeated freeze-thaw cycles.	
Storage:	-80 °C	
Storage Comment:	Store at -80°C.	
Expiry Date:	12 months	