

Datasheet for ABIN3094920 RAG1 Protein (AA 1-1043) (Strep Tag)



Overview

Quantity:	250 µg
Target:	RAG1
Protein Characteristics:	AA 1-1043
Origin:	Human
Source:	Cell-free protein synthesis (CFPS)
Protein Type:	Recombinant
Purification tag / Conjugate:	This RAG1 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Brand:	AliCE®
Sequence:	MAASFPPTLG LSSAPDEIQH PHIKFSEWKF KLFRVRSFEK TPEEAQKEKK DSFEGKPSLE
	QSPAVLDKAD GQKPVPTQPL LKAHPKFSKK FHDNEKARGK AIHQANLRHL CRICGNSFRA
	DEHNRRYPVH GPVDGKTLGL LRKKEKRATS WPDLIAKVFR IDVKADVDSI HPTEFCHNCW
	SIMHRKFSSA PCEVYFPRNV TMEWHPHTPS CDICNTARRG LKRKSLQPNL QLSKKLKTVL
	DQARQARQHK RRAQARISSK DVMKKIANCS KIHLSTKLLA VDFPEHFVKS ISCQICEHIL
	ADPVETNCKH VFCRVCILRC LKVMGSYCPS CRYPCFPTDL ESPVKSFLSV LNSLMVKCPA
	KECNEEVSLE KYNHHISSHK ESKEIFVHIN KGGRPRQHLL SLTRRAQKHR LRELKLQVKA
	FADKEEGGDV KSVCMTLFLL ALRARNEHRQ ADELEAIMQG KGSGLQPAVC LAIRVNTFLS
	CSQYHKMYRT VKAITGRQIF QPLHALRNAE KVLLPGYHHF EWQPPLKNVS SSTDVGIIDG
	LSGLSSSVDD YPVDTIAKRF RYDSALVSAL MDMEEDILEG MRSQDLDDYL NGPFTVVVKE
	SCDGMGDVSE KHGSGPVVPE KAVRFSFTIM KITIAHSSQN VKVFEEAKPN SELCCKPLCL

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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.

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- 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:	RAG1
Alternative Name:	RAG1 (RAG1 Products)
Background:	V(D)J recombination-activating protein 1 (RAG-1) (RING finger protein 74) [Includes:
	Endonuclease RAG1 (EC 3.1), E3 ubiquitin-protein ligase RAG1 (EC 2.3.2.27) (RING-type E3
	ubiquitin transferase RAG1)],FUNCTION: Catalytic component of the RAG complex, a
	multiprotein complex that mediates the DNA cleavage phase during $V(D)J$ recombination.
	V(D)J recombination assembles a diverse repertoire of immunoglobulin and T-cell receptor
	genes in developing B and T-lymphocytes through rearrangement of different V (variable), in
	some cases D (diversity), and J (joining) gene segments. In the RAG complex, RAG1 mediates
	the DNA-binding to the conserved recombination signal sequences (RSS) and catalyzes the
	DNA cleavage activities by introducing a double-strand break between the RSS and the adjacent
	coding segment. RAG2 is not a catalytic component but is required for all known catalytic
	activities. DNA cleavage occurs in 2 steps: a first nick is introduced in the top strand
	immediately upstream of the heptamer, generating a 3'-hydroxyl group that can attack the
	phosphodiester bond on the opposite strand in a direct transesterification reaction, thereby
	creating 4 DNA ends: 2 hairpin coding ends and 2 blunt, 5'-phosphorylated ends. The chromatin
	structure plays an essential role in the $V(D)J$ recombination reactions and the presence of
	histone H3 trimethylated at 'Lys-4' (H3K4me3) stimulates both the nicking and haipinning
	steps. The RAG complex also plays a role in pre-B cell allelic exclusion, a process leading to
	expression of a single immunoglobulin heavy chain allele to enforce clonality and monospecific
	recognition by the B-cell antigen receptor (BCR) expressed on individual B-lymphocytes. The
	introduction of DNA breaks by the RAG complex on one immunoglobulin allele induces ATM-
	dependent repositioning of the other allele to pericentromeric heterochromatin, preventing
	accessibility to the RAG complex and recombination of the second allele. In addition to its

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	endonuclease activity, RAG1 also acts as an E3 ubiquitin-protein ligase that mediates monoubiquitination of histone H3. Histone H3 monoubiquitination is required for the joining step of V(D)J recombination. Mediates polyubiquitination of KPNA1 (By similarity). {ECO:0000250}.
Molecular Weight:	119.1 kDa
UniProt:	P15918
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

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