

Datasheet for ABIN3096105 TDP2 Protein (AA 1-362) (Strep Tag)



Overview

Quantity:	250 μg
Target:	TDP2
Protein Characteristics:	AA 1-362
Origin:	Human
Source:	Cell-free protein synthesis (CFPS)
Protein Type:	Recombinant
Purification tag / Conjugate:	This TDP2 protein is labelled with Strep Tag.
Application:	ELISA, SDS-PAGE (SDS), Western Blotting (WB)

Product Details

Brand:	AliCE®
Sequence:	MELGSCLEGG REAAEEEGEP EVKKRRLLCV EFASVASCDA AVAQCFLAEN DWEMERALNS
	YFEPPVEESA LERRPETISE PKTYVDLTNE ETTDSTTSKI SPSEDTQQEN GSMFSLITWN
	IDGLDLNNLS ERARGVCSYL ALYSPDVIFL QEVIPPYYSY LKKRSSNYEI ITGHEEGYFT
	AIMLKKSRVK LKSQEIIPFP STKMMRNLLC VHVNVSGNEL CLMTSHLEST RGHAAERMNQ
	LKMVLKKMQE APESATVIFA GDTNLRDREV TRCGGLPNNI VDVWEFLGKP KHCQYTWDTQ
	MNSNLGITAA CKLRFDRIFF RAAAEEGHII PRSLDLLGLE KLDCGRFPSD HWGLLCNLDI IL
	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:

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- 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:	TDP2

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Target Details	
Alternative Name:	TDP2 (TDP2 Products)
Background:	Tyrosyl-DNA phosphodiesterase 2 (Tyr-DNA phosphodiesterase 2) (hTDP2) (EC 3.1.4) (5'-
	tyrosyl-DNA phosphodiesterase) (5'-Tyr-DNA phosphodiesterase) (ETS1-associated protein 2)
	(ETS1-associated protein II) (EAPII) (TRAF and TNF receptor-associated protein) (Tyrosyl-RNA
	phosphodiesterase) (VPg unlinkase),FUNCTION: DNA repair enzyme that can remove a variety
	of covalent adducts from DNA through hydrolysis of a 5'-phosphodiester bond, giving rise to
	DNA with a free 5' phosphate. Catalyzes the hydrolysis of dead-end complexes between DNA
	and the topoisomerase 2 (TOP2) active site tyrosine residue. The 5'-tyrosyl DNA
	phosphodiesterase activity can enable the repair of TOP2-induced DNA double-strand
	breaks/DSBs without the need for nuclease activity, creating a 'clean' DSB with 5'-phosphate
	termini that are ready for ligation (PubMed:27099339, PubMed:27060144). Thereby, protects
	the transcription of many genes involved in neurological development and maintenance from
	the abortive activity of TOP2. Hydrolyzes 5'-phosphoglycolates on protruding 5' ends on DSBs
	due to DNA damage by radiation and free radicals. Has preference for single-stranded DNA or
	duplex DNA with a 4 base pair overhang as substrate. Acts as a regulator of ribosome
	biogenesis following stress. Has also 3'-tyrosyl DNA phosphodiesterase activity, but less
	efficiently and much slower than TDP1. Constitutes the major if not only 5'-tyrosyl-DNA
	phosphodiesterase in cells. Also acts as an adapter by participating in the specific activation o
	MAP3K7/TAK1 in response to TGF-beta: associates with components of the TGF-beta
	receptor-TRAF6-TAK1 signaling module and promotes their ubiquitination dependent complex
	formation. Involved in non-canonical TGF-beta induced signaling routes. May also act as a
	negative regulator of ETS1 and may inhibit NF-kappa-B activation.
	{ECO:0000269 PubMed:19794497, ECO:0000269 PubMed:21030584,
	ECO:0000269 PubMed:21921940, ECO:0000269 PubMed:21980489,
	EC0:0000269 PubMed:22405347, EC0:0000269 PubMed:22822062,
	ECO:0000269 PubMed:24658003, ECO:0000269 PubMed:27060144,
	ECO:0000269 PubMed:27099339}., FUNCTION: (Microbial infection) Also acts as a 5'-tyrosyl-
	RNA phosphodiesterase following picornavirus infection: its activity is hijacked by picornavirus
	and acts by specifically cleaving the protein-RNA covalent linkage generated during the viral
	genomic RNA replication steps of a picornavirus infection, without impairing the integrity of
	viral RNA. {ECO:0000269 PubMed:22908287}.
Molecular Weight:	40.9 kDa
UniProt:	095551

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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