Datasheet for ABIN3136718

PARN Protein (AA 1-624) (His tag)

1 Image

Go to Product page

Overview

Quantity:	1 mg
Target:	PARN
Protein Characteristics:	AA 1-624
Origin:	Mouse
Source:	Insect Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This PARN protein is labelled with His tag.
Application:	SDS-PAGE (SDS), Western Blotting (WB), ELISA, Crystallization (Crys)

Product Details

Sequence:	MEIIRSNFKI NLHKVYQAIE EADFFAIDGE FSGISDGPSV TALTSGFDTP EERYQKLKKH SMDFLLFQFG LCAFKYDHTD SKHVTKSFNF YVFPKPFSRS SPDVKFVCQS SSIDFLASQG FDFNKVFCSG IPYLNQEEER QLREQFDEKR SQANGAGALA KCPVTIPEDQ KKFIDQVIEK IEDFLQSEEK RSLELDPCTG FQRKLIYQTL SWKYPKGIHV ETLETDKKER HIVISKVDEE ERKRREQEKY TKEQEELNDA VGFSRVIHAI ANSGKLVVGH NMLLDVMHTI HQFYCPLPAD LNEFKEMAIC VFPRLLDTKL MASTQPFKDI INNTSLAELE KRLKETPFDP PKVESAEGFP SYDTASEQLH EAGYDAYITG LCFISMANYL GSLLSPPKMC VSARSKLIEP FFNKLFLMRV MDIPYLNLEG PDLQPKRDHV LHVTFPKEWK TSDLYQLFSA FGNIQISWID DTSAFVSLSQ PEQVQIAVNT SKYAESYRIQ TYAEYVGKKQ EGKQVKRKWT EDSWKEVDRK RPHMQGPCYH SNSFTAAGVL GKRTLSPDPR EAALEDRESE EVSDSELEQT DSCTDPLPEG RKKSKKLKRM KKELSLAGSV SDSPAVLFEV PDTW
	Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a

	special request, please contact us.
Characteristics:	• Made in Germany - from design to production - by highly experienced protein experts. • Mouse Parn Protein (raised in Insect Cells) purified by multi-step, protein-specific process to ensure crystallization grade. • 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. In the unlikely event that the protein cannot be expressed or purified we do not charge anything (other companies might charge you for any performed steps in the expression process for custom-made proteins, e.g. fees might apply for the expression plasmid, the first expression experiments or purification optimization). When you order this made-to-order protein you will only pay upon receival of the correctly folded protein. With no financial risk on your end you can rest assured that our experienced protein experts will do everything to make sure that you receive the protein you ordered. 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. The concentration of the protein is calculated using its specific absorption coefficient. We use the Expasy's protparam tool to determine the absorption coefficient of each protein.
Purification:	Two step purification of proteins expressed in baculovirus infected SF9 insect cells: 1. In a first purification step, the protein is purified from the cleared cell lysate using three different His-tag capture materials: high yield, EDTA resistant, or DTT resistant. Eluate fractions are analyzed by SDS-PAGE. 2. 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:	>95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.
Sterility:	0.22 µm filtered
Endotoxin Level:	Protein is endotoxin free.
Grade:	Crystallography grade

Target Details

Target:	PARN
Alternative Name:	Parn (PARN Products)
Background:	3'-exoribonuclease that has a preference for poly(A) tails of mRNAs, thereby efficiently degrading poly(A) tails. Exonucleolytic degradation of the poly(A) tail is often the first step in the decay of eukaryotic mRNAs and is also used to silence certain maternal mRNAs translationally during oocyte maturation and early embryonic development. Interacts with both the 3'-end poly(A) tail and the 5'-end cap structure during degradation, the interaction with the cap structure being required for an efficient degradation of poly(A) tails. Involved in nonsense-mediated mRNA decay, a critical process of selective degradation of mRNAs that contain premature stop codons. Also involved in degradation of inherently unstable mRNAs that contain AU-rich elements (AREs) in their 3'-UTR, possibly via its interaction with KHSRP. Probably mediates the removal of poly(A) tails of AREs mRNAs, which constitutes the first step of destabilization (By similarity). {ECO:0000250}.
Molecular Weight:	72.5 kDa Including tag.
UniProt:	Q8VDG3

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 gurantee though.
Comment:	Protein has not been tested for activity yet. In cases in which it is highly likely that the recombinant protein with the default tag will be insoluble our protein lab may suggest a higher molecular weight tag (e.g. GST-tag) instead to increase solubility. We will discuss all possible options with you in detail to assure that you receive your protein of interest.
Restrictions:	For Research Use only

Handling

Format:	Liquid
Buffer:	100 mM NaCL, 20 mM Hepes, 10% glycerol. pH value is at the discretion of the manufacturer.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C

Handling

Storage Comment:	Store at -80°C.
Expiry Date:	Unlimited (if stored properly)

Images

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