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Datasheet for ABIN3086451 PPP1R3B Protein (AA 1-285) (Strep Tag)





Overview

Quantity:	1 mg
Target:	PPP1R3B
Protein Characteristics:	AA 1-285
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This PPP1R3B protein is labelled with Strep Tag.
Application:	SDS-PAGE (SDS), ELISA, Western Blotting (WB)

Product Details

Sequence:	MMAVDIEYRY NCMAPSLRQE RFAFKISPKP SKPLRPCIQL SSKNEASGMV APAVQEKKVK
	KRVSFADNQG LALTMVKVFS EFDDPLDMPF NITELLDNIV SLTTAESESF VLDFSQPSAD
	YLDFRNRLQA DHVCLENCVL KDKAIAGTVK VQNLAFEKTV KIRMTFDTWK SYTDFPCQYV
	KDTYAGSDRD TFSFDISLPE KIQSYERMEF AVYYECNGQT YWDSNRGKNY RIIRAELKST
	QGMTKPHSGP DLGISFDQFG SPRCSYGLFP EWPSYLGYEK LGPYY
	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
	system, a different complexity of the protein could make another tag necessary. In case you have a special request, please contact us.
Characteristics:	
Characteristics:	have a special request, please contact us.

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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®):
	 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.
Endotoxin Level:	Low Endotoxin less than 1 EU/mg (< 0.1 ng/mg)

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Product Details

Grade:

Crystallography grade

Target Details

Target:	PPP1R3B
Alternative Name:	PPP1R3B (PPP1R3B Products)
Background:	Protein phosphatase 1 regulatory subunit 3B (Hepatic glycogen-targeting protein phosphatase
	1 regulatory subunit GL) (Protein phosphatase 1 regulatory subunit 4) (PP1 subunit R4) (Proteir
	phosphatase 1 subunit GL) (PTG),FUNCTION: Acts as a glycogen-targeting subunit for
	phosphatase PP1. Facilitates interaction of the PP1 with enzymes of the glycogen metabolism
	and regulates its activity. Suppresses the rate at which PP1 dephosphorylates (inactivates)
	glycogen phosphorylase and enhances the rate at which it activates glycogen synthase and
	therefore limits glycogen breakdown. Its activity is inhibited by PYGL, resulting in inhibition of
	the glycogen synthase and glycogen phosphorylase phosphatase activities of PP1.
	Dramatically increases basal and insulin-stimulated glycogen synthesis upon overexpression in
	hepatocytes (By similarity). {ECO:0000250}.
Molecular Weight:	32.7 kDa
UniProt:	Q86XI6
Pathways:	Cellular Glucan Metabolic Process, Regulation of Carbohydrate Metabolic Process
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!

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Application Details

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

Images

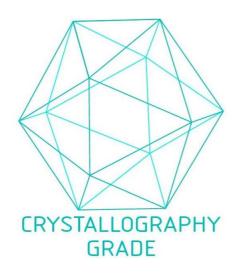


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process