antibodies

Datasheet for ABIN3103479 GDPD1 Protein (AA 1-314) (Strep Tag)



Overview

Image

Quantity:	1 mg
Target:	GDPD1
Protein Characteristics:	AA 1-314
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This GDPD1 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Sequence:	MSSTAAFYLL STLGGYLVTS FLLLKYPTLL HQRKKQRFLS KHISHRGGAG ENLENTMAAF
	QHAVKIGTDM LELDCHITKD EQVVVSHDEN LKRATGVNVN ISDLKYCELP PYLGKLDVSF
	QRACQCEGKD NRIPLLKEVF EAFPNTPINI DIKVNNNVLI KKVSELVKRY NREHLTVWGN
	ANYEIVEKCY KENSDIPILF SLQRVLLILG LFFTGLLPFV PIREQFFEIP MPSIILKLKE PHTMSRSQKF
	LIWLSDLLLM RKALFDHLTA RGIQVYIWVL NEEQEYKRAF DLGATGVMTD YPTKLRDFLH NFSA
	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 by multi-step, protein-specific process to ensure correct folding and modification.

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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 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 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:	GDPD1
Alternative Name:	GDPD1 (GDPD1 Products)
Background:	Lysophospholipase D GDPD1 (EC 3.1.4) (Glycerophosphodiester phosphodiesterase 4)
	(Glycerophosphodiester phosphodiesterase domain-containing protein 1),FUNCTION:
	Hydrolyzes lysoglycerophospholipids to produce lysophosphatidic acid (LPA) and the
	corresponding amines (PubMed:27637550, PubMed:25596343). Shows a preference for 1-0-
	alkyl-sn-glycero-3-phosphocholine (lyso-PAF), lysophosphatidylethanolamine (lyso-PE) and
	lysophosphatidylcholine (lyso-PC) (PubMed:27637550, PubMed:25596343). May be involved ir
	bioactive N-acylethanolamine biosynthesis from both N-acyl-lysoplasmenylethanolamin (N-
	acyl-lysoPlsEt) and N-acyl-lysophosphatidylethanolamin (N-acyl-lysoPE) (PubMed:27637550,
	PubMed:25596343). In addition, hydrolyzes glycerophospho-N-acylethanolamine to N-
	acylethanolamine (PubMed:27637550). Does not display glycerophosphodiester
	phosphodiesterase activity, since it cannot hydrolyze either glycerophosphoinositol or
	glycerophosphocholine (By similarity). {ECO:0000250 UniProtKB:Q9CRY7,
	ECO:0000269 PubMed:25596343, ECO:0000269 PubMed:27637550}.
Molecular Weight:	36.2 kDa
UniProt:	Q8N9F7
Application Details	
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Application Details	
	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. 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

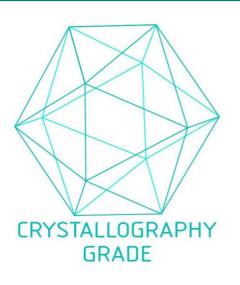


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process

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