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POLD4 Protein (AA 1-107) (Strep Tag)



Image



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Overview

Quantity:	1 mg
Target:	POLD4
Protein Characteristics:	AA 1-107
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This POLD4 protein is labelled with Strep Tag.
Application:	ELISA, Western Blotting (WB), SDS-PAGE (SDS)

Product Details

Sequence:	MGRKRLITDS YPVVKRREGP AGHSKGELAP ELGEEPQPRD EEEAELELLR QFDLAWQYGP
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CTGITRLQRW CRAKQMGLEP PPEVWQVLKT HPGDPRFQCS LWHLYPL

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.
- 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®):
	1. 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)
Grade:	Crystallography grade

Target Details

Target:	POLD4
Alternative Name:	POLD4 (POLD4 Products)
Background:	DNA polymerase delta subunit 4 (DNA polymerase delta subunit p12),FUNCTION: As a
	component of the tetrameric DNA polymerase delta complex (Pol-delta4), plays a role in high
	fidelity genome replication and repair. Within this complex, increases the rate of DNA synthesis
	and decreases fidelity by regulating POLD1 polymerase and proofreading 3' to 5' exonuclease
	activity (PubMed:16510448, PubMed:19074196, PubMed:20334433). Pol-delta4 participates in
	Okazaki fragment processing, through both the short flap pathway, as well as a nick translation
	system (PubMed:24035200). Under conditions of DNA replication stress, required for the repai
	of broken replication forks through break-induced replication (BIR), a mechanism that may
	induce segmental genomic duplications of up to 200 kb (PubMed:24310611). Involved in Pol-
	delta4 translesion synthesis (TLS) of templates carrying O6-methylguanine or abasic sites
	(PubMed:19074196). Its degradation in response to DNA damage is required for the inhibition
	of fork progression and cell survival (PubMed:24022480). {ECO:0000269 PubMed:16510448,
	ECO:0000269 PubMed:19074196, ECO:0000269 PubMed:20334433,
	ECO:0000269 PubMed:24022480, ECO:0000269 PubMed:24035200,
	ECO:0000269 PubMed:24310611}.
Molecular Weight:	12.4 kDa
UniProt:	Q9HCU8
Pathways:	Telomere Maintenance, DNA Damage Repair, DNA Replication, Synthesis of DNA
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.
	modifications. During lysate production, the cell wall and other cellular components that are not required for
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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

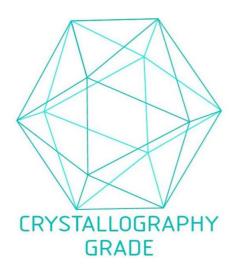


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process