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DACH2 Protein (AA 1-599) (Strep Tag)



Image



Overview

Quantity:	1 mg
Target:	DACH2
Protein Characteristics:	AA 1-599
Origin:	Human
Source:	Tobacco (Nicotiana tabacum)
Protein Type:	Recombinant
Purification tag / Conjugate:	This DACH2 protein is labelled with Strep Tag.
Application:	ELISA, SDS-PAGE (SDS), Western Blotting (WB)

Product Details

Sequence:

MAVSASPVIS ATSSGAGVPG GLFRAEPLYS TPREPPRLTP NMINSFVVNN HSNSAGGGGR GNTNTNECRM VDMHGMKVAS FLMDGQELIC LPQVFDLFLK HLVGGLHTVY TKLKRLDISP VVCTVEQVRI LRGLGAIQPG VNRCKLITRK DFETLFTDCT NARRKRQMTR KQAVNSSRPG RPPKRSLGVL QENARLLTHA VPGLLSPGLI TPTGITAAAM AEAMKLQKMK LMAMNTLQGN GSQNGTESEP DDLNSNTGGS ESSWDKDKMQ SPFAAPGPQH GIAHAALAGQ PGIGGAPTLN PLQQNHLLTN RLDLPFMMMP HPLLPVSLPP ASVAMAMNQM NHLNTIANMA AAAQIHSPLS RAGTSVIKER IPESPSPAPS LEENHRPGSQ TSSHTSSSVS SSPSQMDHHL ERMEEVPVQI PIMKSPLDKI QLTPGQALPA GFPGPFIFAD SLSSVETLLT NIQGLLKVAL DNARIQEKQI QQEKKELRLE LYREREIREN LERQLAVELQ SRTTMQKRLK KEKKTKRKLQ EALEFESKRR EQVEQALKQA TTSDSGLRML KDTGIPDIEI ENNGTPHDSA AMQGGNYYCL EMAQQLYSA

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:	DACH2
Alternative Name:	DACH2 (DACH2 Products)
Background:	Dachshund homolog 2 (Dach2),FUNCTION: Transcription factor that is involved in regulation of organogenesis. Seems to be a regulator for SIX1 and SIX6. Seems to act as a corepressor of SIX6 in regulating proliferation by directly repressing cyclin-dependent kinase inhibitors, including the p27Kip1 promoter. Is recruited with SIX6 to the p27Kip1 promoter in embryonal retina. SIX6 corepression seems also to involve NCOR1, TBL1, HDAC1 and HDAC3. May be involved together with PAX3, SIX1, and EYA2 in regulation of myogenesis. In the developing somite, expression of DACH2 and PAX3 is regulated by the overlying ectoderm, and DACH2 and PAX3 positively regulate each other's expression (By similarity). Probably binds to DNA via its DACHbox-N domain. {ECO:0000250}.
Molecular Weight:	65.3 kDa
UniProt: Application Details	Q96NX9
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

Application Details

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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

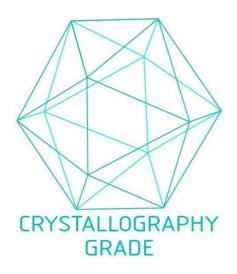


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process