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Datasheet for ABIN3135250 DYRK2 Protein (AA 1-599) (His tag)

Image



Overview

Quantity:	1 mg
Target:	DYRK2
Protein Characteristics:	AA 1-599
Origin:	Mouse
Source:	Insect Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This DYRK2 protein is labelled with His tag.
Application:	Western Blotting (WB), ELISA, SDS-PAGE (SDS), Crystallization (Crys)

Product Details

Sequence:	MLTRKPSAAA PAAYPTGRGG DTAVRQLQAS PGIGAGAPRS GVGTGPPSPI ALPPLRASNA
	TTTAHTIGGS KHTMNDHLHL NSHGQIQVQQ LFEDNSNKRT VLTTQPNGLT TVGKTGLPGV
	PERQLESIHR RQGSSTSLKS MEGMGKVKAS PMTPEQAMKQ YMQKLTAFEH HEIFSYPEIY
	FLGPNAKKRQ GMTGGPNNGG YDDDQGSYVQ VPHDHVAYRY EVLKVIGKGS FGQVVKAYDH
	KVHQHVALKM VRNEKRFHRQ AAEEIRILEH LRKQDKDNTM NVIHMLENFT FRNHICMTFE
	LLSMNLYELI KKNKFQGFSL PLVRKFAHSI LQCLDALHKN RIIHCDLKPE NILLKQQGRS
	SIKVIDFGSS CYEHQRVYTY IQSRFYRAPE VILGARYGMP IDMWSLGCIL AELLTGYPLL
	PGEDEGDQLA CMIELLGMPS QKLLDASKRA KNFVSSKGYP RYCTVTTLSD GSVVLNGGRS
	RRGKLRGPPE SREWGNALKG CDDPLFLDFL KQCLEWDPAV RMTPGQALRH PWLRRRLPKP
	PTGEKTAVKR VTESTGAITS ISKLPPPSSS ASKLRTNLAQ MTDANGNIQQ RTVLPKLVS
	Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a
	special request, please contact us.

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Product Details	
Characteristics:	 Made in Germany - from design to production - by highly experienced protein experts. Mouse Dyrk2 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:
	 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. 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

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Target Details	
Target:	DYRK2
Alternative Name:	Dyrk2 (DYRK2 Products)
Background:	Serine/threonine-protein kinase involved in the regulation of the mitotic cell cycle, cell
	proliferation, apoptosis, organization of the cytoskeleton and neurite outgrowth. Functions in
	part via its role in ubiquitin-dependent proteasomal protein degradation. Functions downstream
	of ATM and phosphorylates p53/TP53 at 'Ser-46', and thereby contributes to the induction of
	apoptosis in response to DNA damage. Phosphorylates NFATC1, and thereby inhibits its
	accumulation in the nucleus and its transcription factor activity. Phosphorylates EIF2B5 at 'Ser-
	544', enabling its subsequent phosphorylation and inhibition by GSK3B. Likewise,
	phosphorylation of NFATC1, CRMP2/DPYSL2 and CRMP4/DPYSL3 promotes their subsequent
	phosphorylation by GSK3B. May play a general role in the priming of GSK3 substrates.
	Inactivates GYS1 by phosphorylation at 'Ser-641', and potentially also a second phosphorylation
	site, thus regulating glycogen synthesis. Mediates EDVP E3 ligase complex formation and is
	required for the phosphorylation and subsequent degradation of KATNA1. Phosphorylates
	TERT at 'Ser-457', promoting TERT ubiquitination by the EDVP complex. Phosphorylates SIAH2,
	and thereby increases its ubiquitin ligase activity. Promotes the proteasomal degradation of
	MYC and JUN, and thereby regulates progress through the mitotic cell cycle and cell
	proliferation. Promotes proteasomal degradation of GLI2 and GLI3, and thereby plays a role in
	smoothened and sonic hedgehog signaling. Phosphorylates CRMP2/DPYSL2, CRMP4/DPYSL3,
	DCX, EIF2B5, EIF4EBP1, GLI2, GLI3, GYS1, JUN, MDM2, MYC, NFATC1, p53/TP53, TAU/MAPT
	and KATNA1. Can phosphorylate histone H1, histone H3 and histone H2B (in vitro). Can
	phosphorylate CARHSP1 (in vitro) (By similarity). Plays a role in cytoskeleton organization and
	neurite outgrowth via its phosphorylation of DCX. {ECO:0000250,
	EC0:0000269 PubMed:22359282}.
Molecular Weight:	67.5 kDa Including tag.
UniProt:	Q5U4C9
Pathways:	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 gurantee
	though.
Comment:	Protein has not been tested for activity yet. In cases in which it is highly likely that the

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

Images

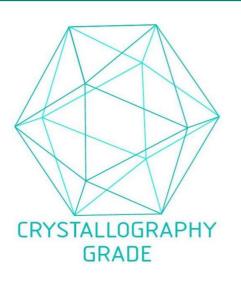


Image 1. "Crystallography Grade" protein due to multi-step, protein-specific purification process

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