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Datasheet for ABIN3115084 KCNMB1 Protein (AA 2-191) (rho-1D4 tag)



Overview

000101000	
Quantity:	1 mg
Target:	KCNMB1
Protein Characteristics:	AA 2-191
Origin:	Human
Source:	Insect Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This KCNMB1 protein is labelled with rho-1D4 tag.
Application:	ELISA, Western Blotting (WB), Crystallization (Crys), SDS-PAGE (SDS)
Product Details	
Sequence:	VKKLVMAQKR GETRALCLGV TMVVCAVITY YILVTTVLPL YQKSVWTQES KCHLIETNIR
	DQEELKGKKV PQYPCLWVNV SAAGRWAVLY HTEDTRDQNQ QCSYIPGSVD NYQTARADVE
	KVRAKFQEQQ VFYCFSAPRG NETSVLFQRL YGPQALLFSL FWPTFLLTGG LLIIAMVKSN
	QYLSILAAQK
	Sequence without tag. Tag location is at the discretion of the manufacturer. If you have a
	special request, please contact us.
Characteristics:	Made in Germany - from design to production - by highly experienced protein experts.
	Human KCNMB1 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

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Durification	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:	Three step purification of membrane proteins expressed in baculovirus infected SF9 insect cells:
	 Membrane proteins are fractioned by ultracentrifugation and subsequently solubilized with different detergents (detergent screen). Samples are analyzed by Western blot.
	 The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot.
Purity:	 The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and
Purity: Sterility:	 The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot.
	 2. The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. 3. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot. >95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.
Sterility:	 2. The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. 3. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot. >95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot. 0.22 µm filtered
Sterility: Endotoxin Level:	 2. The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. 3. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot. >95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot. 0.22 µm filtered Protein is endotoxin-free.
Sterility: Endotoxin Level: Grade:	 2. The best performing detergent is used for solubilization and the proteins are purified via their rho1D4 tag via two rho1D4 antibody columns: one DTT resistant, the other one not. Eluate fractions are analyzed by Western blot. 3. Protein containing fractions of the best purification are subjected to second purification step through size exclusion chromatograph. Eluate fractions are analyzed by SDS-PAGE and Western blot. >95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot. 0.22 µm filtered Protein is endotoxin-free.

Background: Regulatory subunit of the calcium activated potassium KCNMA1 (maxiK) channel. Modulates

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Target Details

	the calcium sensitivity and gating kinetics of KCNMA1, thereby contributing to KCNMA1
	channel diversity. Increases the apparent Ca(2+)/voltage sensitivity of the KCNMA1 channel. It
	also modifies KCNMA1 channel kinetics and alters its pharmacological properties. It slows
	down the activation and the deactivation kinetics of the channel. Acts as a negative regulator of
	smooth muscle contraction by enhancing the calcium sensitivity to KCNMA1. Its presence is
	also a requirement for internal binding of the KCNMA1 channel opener dehydrosoyasaponin I
	(DHS-1) triterpene glycoside and for external binding of the agonist hormone 17-beta-estradiol
	(E2). Increases the binding activity of charybdotoxin (CTX) toxin to KCNMA1 peptide blocker by
	increasing the CTX association rate and decreasing the dissociation rate.
Molecular Weight:	22.8 kDa Including tag.
UniProt:	Q16558
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:	In cases in which it is highly likely that the 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
Buffer:	100 mM NaCL, 20 mM Hepes, 10% glycerol. pH value is at the discretion of the manufacturer.
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C
Storage Comment:	Store at -80°C.
Expiry Date:	Unlimited (if stored properly)

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