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NME7 Protein (full length) (rho-1D4 tag)



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Overview		
Quantity:	0.5 mg	
Target:	NME7	
Protein Characteristics:	full length	
Origin:	CHO cells	
Source:	Insect Cells	
Protein Type:	Recombinant	
Purification tag / Conjugate:	This NME7 protein is labelled with rho-1D4 tag.	
Application:	ELISA, Western Blotting (WB), Crystallization (Crys), SDS-PAGE (SDS), Functional Studies (Func)	
Product Details		
Sequence:	MANLERTFIA IKPDGVQRGL VGDIIKRFEQ KGFRLVAMKF LRASEEHLKQ HYIDLKDRPF	
	FPGLVKYMNS GPVVAMVWEG LNVVKTGRMM LGETNPADSK PGTIRGDFCI QVGRNIIHGS	
	DSVQSAEKEI SLWFKPEELI DYKPCAHDWV YE	
	Sequence without tag. The location of the tag depends on protein. You may also submit your	
	preference when ordering.	
Characteristics:	 Made in Germany - from design to production - by highly experienced protein experts. CHO Nucleoside diphosphate kinase 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 custom-made protein and will be made for the first time for your order. This	
	protein will be produced on the basis of on a Custom Service Project. We will make sure that	
	every step in the production is successful from the design of the expression plasmid to the	

Restrictions:

expression and purification of the final protein. Our experts in the lab will ensure that you receive a correctly folded protein.	
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.	
Three step purification of proteins expressed in baculovirus infected SF9 insect cells:	
1. Membrane proteins are fractioned by ultracentrifugation and subsequently solubilized with	
different detergents (detergent screen). Samples are analyzed by 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.	
0.22 μm filtered	
Endotoxins have not been removed. Please contact us if you require an endotoxin-free version	
of this product.	
Crystallography grade	
Protein has not been tested for activity yet.	
NME7	
Nucleoside diphosphate kinase (NME7 Products)	
G3HBD3	
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For Research Use only

Handling

Format:	Liquid
Buffer:	$150\mathrm{mM}$ NaCL, $20\mathrm{mM}$ NaH2PO4 pH 7.4, $10~\%$ glycerol. Note: Isoelectric point of protein taken into account regarding pH .
Handling Advice:	Avoid repeated freeze-thaw cycles.
Storage:	-80 °C
Storage Comment:	Store at -80°C.
Expiry Date:	Unlimited (if stored properly)