antibodies -online.com





PSMA7 Protein (full length) (rho-1D4 tag)



Go to	Prod	luct	page

()	11/0	K\ /	iew
	\cup	'I V/I	$I \cap VV$

Overview	
Quantity:	0.5 mg
Target:	PSMA7
Protein Characteristics:	full length
Origin:	CHO cells
Source:	Insect Cells
Protein Type:	Recombinant
Purification tag / Conjugate:	This PSMA7 protein is labelled with rho-1D4 tag.
Application:	SDS-PAGE (SDS), ELISA, Western Blotting (WB), Crystallization (Crys), Functional Studies (Func)
Product Details	
Sequence:	VGVRGKDIVV LGVEKKSVAK LQDERTVRKI CALDDNVCMA FAGLTADARI VINRARVECQ
	SHRLTVEDPV TVEYITRYIA SLKQSNGRRP FGISALIVGF DFDGTPRLYQ TDPSGTYHAW
	KANAIGRGAK SVREFLEKNY TDDAIETDDL TIKLVIKALL EVVQSGGKNI ELAVMRRDQP
	LKILNPEEIE KYVAEIEKEK EENEKKKQKK AS
	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 Proteasome subunit alpha type-7 (Fragment) 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 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.

Purification:

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.
- 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:	>95 % as determined by SDS PAGE, Size Exclusion Chromatography and Western Blot.
Sterility:	0.22 μm filtered
Endotoxin Level:	Endotoxins have not been removed. Please contact us if you require an endotoxin-free version of this product.
Grade:	Crystallography grade

Biological Activity Comment:

Protein has not been tested for activity yet.

Target Details

Target:	PSMA7
Alternative Name: Proteasome subunit alpha type-7 (PSMA7 Products)	
UniProt:	G3GWR8
Pathways:	Mitotic G1-G1/S Phases, DNA Replication, Synthesis of DNA

Application Details

Application Notes: Optimal working dilution should be determined by the investigator.

Application Details

Restrictions:	For Research Use only
Handling	
Format:	Liquid
Buffer:	150 mM NaCL, 20 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)