

Rat PSMA/FOLH1 Protein (active dimer)

Cat. No. PSM-RM110

Description

Source	Recombinant Rat PSMA/FOLH1 Protein (active dimer) is expressed from HEK293 with His tag at the N-terminus. It contains Lys45-Asp752.
Accession	P70627-1
Molecular Weight	The protein has a predicted MW of 80.62 kDa. Due to glycosylation, the protein migrates to 90-110 kDa based on Bis-Tris PAGE result.
Endotoxin	Less than 0.1 EU per μg by the LAL method.
Purity	> 90% as determined by Bis-Tris PAGE > 95% as determined by HPLC

Formulation and Storage

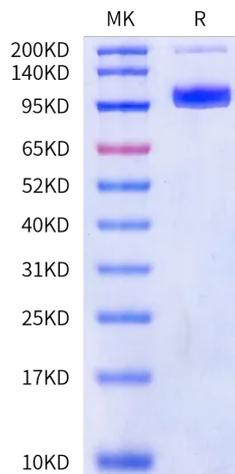
Formulation	Lyophilized from 0.22 μm filtered solution in 50mM MES, 150mM NaCl, 8% trehalose (pH 6.0).
Reconstitution	Dissolve the lyophilized protein in 50mM MES (pH 6.0). Please refer to the Certificate of Analysis for detailed instructions.
Storage	-20 to -80°C for 12 months as supplied from date of receipt. -80°C for 3 months after reconstitution. Recommend to aliquot the protein into smaller quantities for optimal storage. Please minimize freeze-thaw cycles.

Background

Prostate-specific membrane antigen (PSMA) is an enzyme that in humans is encoded by the FOLH1 (folate hydrolase 1) gene, also known as Glutamate carboxypeptidase II (GCPII). Human PSMA is highly expressed in the prostate, roughly a hundred times greater than in most other tissues. In some prostate cancers, PSMA is the second-most upregulated gene product, with an 8- to 12-fold increase over levels in noncancerous prostate cells.

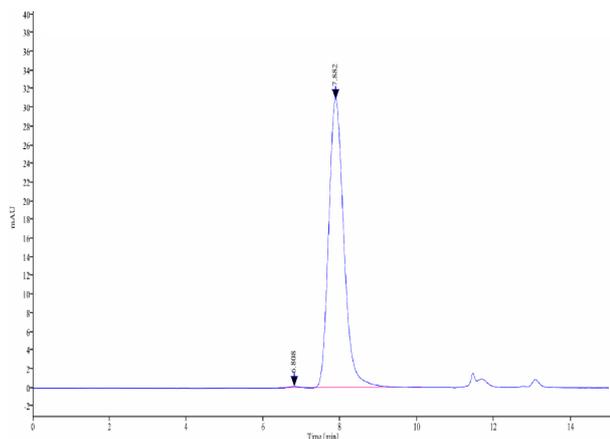
Assay Data

Bis-Tris PAGE



Rat PSMA on Bis-Tris PAGE under reduced condition. The purity is greater than 90%.

SEC-HPLC



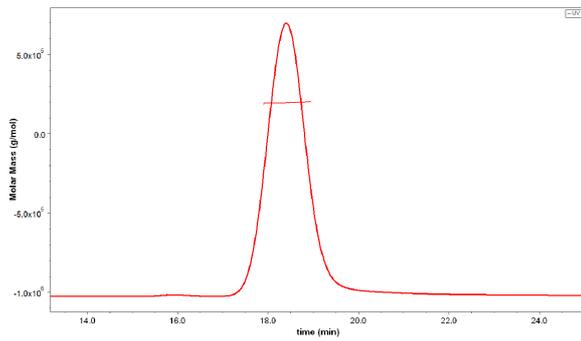
The purity of Rat PSMA is greater than 95% as determined by SEC-HPLC.

Rat PSMA/FOLH1 Protein (active dimer)

Cat. No. PSM-RM110

Assay Data

SEC-MALS



The purity of Rat PSMA is greater than 95% and the molecular weight of this protein is around 180-210 kDa as determined by SEC-MALS.

Bioactivity Data

Measured by its ability to hydrolyze the substrate N-acetyl-L-Asp-L-Glu into N-acetyl-L-Asp and L-Glu. The L-Glu product is measured by fluorescence after its derivatization by ortho-phthaldialdehyde. The specific activity is >500 pmol/min/ μ g.