

# Recombinant Human PSMA Protein(Fc Tag)

Catalog Number: PDMH100295



**Note:** Centrifuge before opening to ensure complete recovery of vial contents.

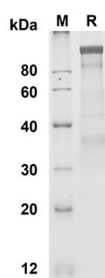
## Description

<b>Species</b>	Human
<b>Source</b>	Mammalian-derived Human PSMA protein Lys44-Ala750,with C-terminal Fc
<b>Mol_Mass</b>	102.6 kDa
<b>Accession</b>	Q04609
<b>Bio-activity</b>	Not validated for activity

## Properties

<b>Purity</b>	> 90% as determined by reducing SDS-PAGE.
<b>Endotoxin</b>	< 1.0 EU/mg of the protein as determined by the LAL method
<b>Storage</b>	Generally, lyophilized proteins are stable for up to 12 months when stored at -20 to -80 °C. Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.
<b>Shipping</b>	This product is provided as lyophilized powder which is shipped with ice packs.
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS with 5% Trehalose and 5% Mannitol.
<b>Reconstitution</b>	It is recommended that sterile water be added to the vial to prepare a stock solution of 0.5 mg/mL. Concentration is measured by UV-Vis.

## Data



SDS-PAGE analysis of Human PSMA proteins , 2µg/lane of Recombinant Human PSMAL proteins was resolved with SDS-PAGE under reducing conditions , showing bands at 100-110 KD

## Background

Glutamate carboxypeptidase 2 , also known as Glutamate carboxypeptidase II , Membrane glutamate carboxypeptidase , Prostate-specific membrane antigen , GCPII , PSMA , FOLH1 , and NAALAD1 , is a single-pass type II membrane protein which belongs to thepeptidase M28 family and M28B subfamily. FOLH1 is highly expressed in prostate epithelium. It is detected in urinary bladder , kidney , testis , ovary , fallopian tube , breast , adrenal gland , liver , esophagus , stomach , small intestine , colon , brain (at protein level) , and the capillary endothelium of a variety of tumors. FOLH1 has both folate hydrolase and N-acetylated alpha linked acidic dipeptidase (NAALADase) activity. It has a preference for tri-alpha a-glutamate peptides. Genetic variation in FOLH1 may be associated with low folate levels and consequent hyperhomocysteinemia. This condition can result in increased risk of cardiovascular disease , neural tube defects , and cognitive deficits. FOLH1 also shows a promising role in directed imaging and therapy of recurrent or metastatic disease.

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