

## PSP Protein, Human, Recombinant (His)

## General Information

|                       |  |
|-----------------------|--|
| Synonyms:             | PSPH;L-3-Phosphoserine Phosphatase;PSPase;Phosphoserine Phosphatase;O-Phosphoserine Phosphohydrolase;PSP |
| Protein Construction: | Met1-Glu225  |
| Species:              | Human  |
| Expression Host:      | E. coli  |
| Accession:            | P78330   |
| Molecular Weight:     | 25-30 KDa (reducing condition)   |
| AA Sequence:          | Met1-Glu225  |

## QC Testing

|                      |   |
|----------------------|---|
| Biological Activity: | Activity has not been tested. It is theoretically active, but we cannot guarantee it. If you require protein activity, we recommend choosing the eukaryotic expression version first. |
| Purity:              | Greater than 95% as determined by reducing SDS-PAGE. (QC verified)  |
| Endotoxin:           | < 0.1 ng/μg (1 EU/μg) as determined by LAL test.  |
| Formulation:         | Supplied as a 0.2 μm filtered solution of 20 mM Tris-HCL, 4M Urea, 5 mM EDTA, pH 8.0.   |

## Preparation and Storage

## Stability &amp; Storage:

Lyophilized powders can be stably stored for over 12 months, while liquid products can be stored for 6-12 months at -80°C. For reconstituted protein solutions, the solution can be stored at -20°C to -80°C for at least 3 months. Please avoid multiple freeze-thaw cycles and store products in aliquots.

Actual storage temperature shall be subject to the COA.

## Shipping:

Proteins are shipped with blue ice.

## Protein Background

Phosphoserine phosphatase (PSP) is an enzyme that belongs to the serB family. PSPH catalyzes magnesium-dependent hydrolysis of L-phosphoserine and is also involved in an exchange reaction between L-serine and L-phosphoserine. The reaction mechanism proceeds via the formation of a phosphoryl-enzyme intermediates. Deficiency of this protein is thought to be linked to Williams syndrome. A disorder that results in pre- and postnatal growth retardation, moderate psychomotor retardation and facial features suggestive of Williams syndrome.

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