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# Recombinant Human TPP1/CLN2 Protein (His Tag)

Catalog No. PKSH033493

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

## **Description**

Synonyms Tripeptidyl-Peptidase 1;TPP-1;Cell Growth-Inhibiting Gene 1 Protein;Lysosomal

Pepstatin-Insensitive Protease;LPIC;Tripeptidyl

Aminopeptidase;TPP1;CLN2;GIG1;LPIC;SCAR7;TPP-1

Species Human

Expression Host HEK293 Cells
Sequence Ser20-Pro563
Accession AAH14863.1
Calculated Molecular Weight 60.4 kDa
Observed molecular weight 74 kDa
Tag C-His

**Bioactivity** Not validated for activity

### **Properties**

**Purity** > 95 % as determined by reducing SDS-PAGE.

**Endotoxin**  $< 1.0 \text{ EU per } \mu \text{g of the protein as determined by the LAL method.}$ 

Storage Storage Store at < -20°C, stable for 6 months. Please minimize freeze-thaw cycles.

**Shipping** This product is provided as liquid. It is shipped at frozen temperature with blue

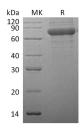
ice/gel packs. Upon receipt, store it immediately at < - 20°C.

Formulation Supplied as a 0.2 µm filtered solution of 20mM Tris-HCl, 150mM NaCl, 1mM

CaCl<sub>2</sub>, 10% Glycerol, pH 7.5.

**Reconstitution** Not Applicable

#### Data



> 95 % as determined by reducing SDS-PAGE.

## **Background**

Tripeptidyl-Peptidase 1 (TPP1) belongs to the peptidase S53 family. TPP1 is detected in all tissues examined with highest levels in heart and placenta and relatively similar levels in other tissues. TPP1 is lysosomal serine protease with tripeptidyl-peptidase I activity. TPP1 may act as a non-specific lysosomal peptidase which generates tripeptides from the breakdown

#### For Research Use Only

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products produced by lysosomal proteinases. TPP1 requires substrates with an unsubstituted N-terminus. TPP1 mutations have also been shown to cause neuronal ceroid lipofuscinosis type 2 (CLN2).

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