

## Product datasheet for **TP721098**

### **PPT1 (NM\_000310) Human Recombinant Protein**

#### **Product data:**

<b>Product Type:</b>	Recombinant Proteins
<b>Description:</b>	Purified recombinant protein of Human palmitoyl-protein thioesterase 1 (PPT1), transcript variant 1
<b>Species:</b>	Human
<b>Expression Host:</b>	HEK293
<b>Expression cDNA Clone or AA Sequence:</b>	Asp28-Gly306
<b>Tag:</b>	C-His
<b>Predicted MW:</b>	32.3 kDa
<b>Purity:</b>	>95% as determined by SDS-PAGE and Coomassie blue staining
<b>Buffer:</b>	Provided lyophilized from a 0.2 µm filtered solution of 20 mM Tris-HCl, 150 mM NaCl
<b>Endotoxin:</b>	Endotoxin level is < 0.1 ng/µg of protein (< 1 EU/µg)
<b>Storage:</b>	Store at -80°C.
<b>Stability:</b>	Stable for at least 3 months from date of receipt under proper storage and handling conditions.
<b>RefSeq:</b>	<a href="#">NP_000301</a>
<b>Locus ID:</b>	5538
<b>UniProt ID:</b>	<a href="#">P50897</a>
<b>RefSeq Size:</b>	2504
<b>Cytogenetics:</b>	1p34.2
<b>RefSeq ORF:</b>	918
<b>Synonyms:</b>	CLN1; INCL; PPT



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<b>Summary:</b>	The protein encoded by this gene is a small glycoprotein involved in the catabolism of lipid-modified proteins during lysosomal degradation. The encoded enzyme removes thioester-linked fatty acyl groups such as palmitate from cysteine residues. Defects in this gene are a cause of infantile neuronal ceroid lipofuscinosis 1 (CLN1, or INCL) and neuronal ceroid lipofuscinosis 4 (CLN4). Two transcript variants encoding different isoforms have been found for this gene.[provided by RefSeq, Dec 2008]
<b>Protein Families:</b>	Druggable Genome
<b>Protein Pathways:</b>	Fatty acid elongation in mitochondria, Lysosome, Metabolic pathways