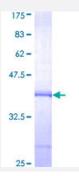


TPP1 (Human) Recombinant Protein (Q01)

Catalog # H00001200-Q01 Size 25 ug, 10 ug

Applications



Specification	
Product Description	Human TPP1 partial ORF (AAH14863, 195 a.a 304 a.a.) recombinant protein with GST-tag at N-t erminal.
Sequence	GLHLGVTPSVIRKRYNLTSQDVGSGTSNNSQACAQFLEQYFHDSDLAQFMRLFGGNFAHQASVA RVVGQQGRGRAGIEASLDVQYLMSAGANISTWVYSSPGRHEGQEPF
Host	Wheat Germ (in vitro)
Theoretical MW (kDa)	37.84
Interspecies Antigen Sequence	Mouse (87)
Preparation Method	in vitro wheat germ expression system
Purification	Glutathione Sepharose 4 Fast Flow
Quality Control Testing	12.5% SDS-PAGE Stained with Coomassie Blue.
Storage Buffer	50 mM Tris-HCI, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.
Storage Instruction	Store at -80°C. Aliquot to avoid repeated freezing and thawing.
Note	Best use within three months from the date of receipt of this protein.



Applications

- Enzyme-linked Immunoabsorbent Assay
- Western Blot (Recombinant protein)
- Antibody Production
- Protein Array

Gene Info — TPP1	
Entrez GenelD	1200
GeneBank Accession#	BC014863
Protein Accession#	<u>AAH14863</u>
Gene Name	TPP1
Gene Alias	CLN2, GIG1, LPIC, MGC21297
Gene Description	tripeptidyl peptidase I
Omim ID	<u>204500</u> <u>607998</u>
Gene Ontology	<u>Hyperlink</u>
Gene Summary	This gene encodes a member of the sedolisin family of serine proteases. The protease functions in the lysosome to cleave N-terminal tripeptides from substrates, and has weaker endopeptidase activity. It is synthesized as a catalytically-inactive enzyme which is activated and auto-proteolyzed upon acidification. Mutations in this gene result in late-infantile neuronal ceroid lipofuscinosis, which is associated with the failure to degrade specific neuropeptides and a subunit of ATP synthase in the lysosome. [provided by RefSeq
Other Designations	ceroid-lipofuscinosis, neuronal 2, late infantile (Jansky-Bielschowsky disease) growth-inhibiting protein 1 lysosomal pepstatin insensitive protease tripeptidyl aminopeptidase tripeptidyl-peptidase

Pathway

Lysosome



Disease

- Kidney Failure
- Neuronal Ceroid-Lipofuscinoses