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Produktinformation



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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

SZABO-SCANDIC HandelsgmbH

Quellenstraße 110, A-1100 Wien

T. +43(0)1 489 3961-0

F. +43(0)1 489 3961-7

mail@szabo-scandic.com

www.szabo-scandic.com

[linkedin.com/company/szaboscandic](https://www.linkedin.com/company/szaboscandic) 

PRODUCT INFORMATION



PPT1 (human, recombinant)

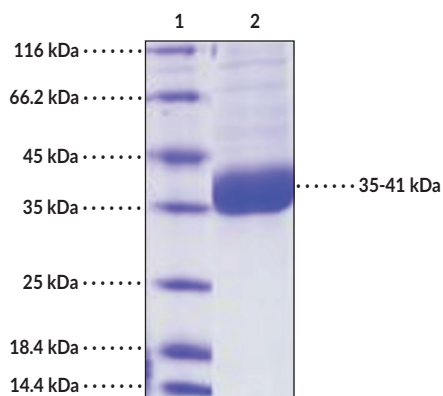
Item No. 38066

Overview and Properties

Synonyms: Palmitoyl-protein Hydrolase 1, Palmitoyl-protein Thioesterase 1, CLN1
Source: Recombinant human C-terminal His-tagged PPT1 expressed in HEK293 cells
Amino Acids: 28-306
Uniprot No.: P50897-1
Molecular Weight: 32.7 kDa
Storage: -80°C (as supplied)
Stability: ≥1 year
Purity: ≥90% estimated by SDS-PAGE
Supplied in: Lyophilized from sterile PBS, pH 7.4
Endotoxin Testing: <1.0 EU/μg, determined by the LAL endotoxin assay

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

Image



Lane 1: MW Markers
Lane 2: PPT1

SDS-PAGE Analysis of PPT1. This protein has a calculated molecular weight of 32.7 kDa. It has an apparent molecular weight of approximately 35-41 kDa by SDS-PAGE due to glycosylation.

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

SAFETY DATA
This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

WARRANTY AND LIMITATION OF REMEDY
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CAYMAN CHEMICAL

1180 EAST ELLSWORTH RD

ANN ARBOR, MI 48108 · USA

PHONE: [800] 364-9897

[734] 971-3335

FAX: [734] 971-3640

CUSTSERV@CAYMANCHEM.COM

WWW.CAYMANCHEM.COM

PRODUCT INFORMATION



Description

Palmitoyl-protein thioesterase 1 (PPT1) is a lysosomal hydrolase involved in removing palmitic acid (Item No. 10006627) from post-translationally modified proteins prior to their degradation.¹ It is mainly expressed in the brain but is also found in visceral macrophages from liver, lung, and bowel tissues.² PPT1 depalmitoylates proteins in cells undergoing autophagy, and PPT1 activity is important for axon outgrowth, neurite extension, and dendritic spine morphology.¹ Knockout of *PPT1* decreases tumor growth in a mouse xenograft model, and high expression of *PPT1* is associated with poor overall survival in several cancers.³ Mutations in *PPT1* cause infantile neuronal ceroid lipofuscinoses (INCL), an encephalopathy characterized by granular deposits in neurons, vision loss, seizures, mental deterioration, and brain death.^{4,5} Cayman's PPT1 (human, recombinant) protein consists of 290 amino acids, has a calculated molecular weight of 32.7 kDa, and a predicted N-terminus of Asp28 after signal peptide cleavage. By SDS-PAGE, under reducing conditions, the apparent molecular mass is 35-41 kDa due to glycosylation.

References

1. Koster, K.P. and Yoshii, A. Depalmitoylation by palmitoyl-protein thioesterase 1 in neuronal health and degeneration. *Front. Synaptic Neurosci.* **11**, 25 (2019).
2. Margraf, L.R., Boriack, R.L., Routheut, A.A., *et al.* Tissue expression and subcellular localization of CLN3, the Batten disease protein. *Mol. Genet. Metab.* **66(4)**, 283-289 (1999).
3. Rebecca, V.W., Nicastrì, M.C., Fennelly, C., *et al.* PPT1 promotes tumor growth and is the molecular target of chloroquine derivatives in cancer. *Cancer Discov.* **9(2)**, 220-229 (2019).
4. Vesa, J., Hellsten, E., Verkruyse, L.A., *et al.* Mutations in the palmitoyl protein thioesterase gene causing infantile neuronal ceroid lipofuscinosis. *Nature* **376(6541)**, 584-587 (1995).
5. Lu, J.-Y., Verkruyse, L.A., and Hofmann, S.L. Lipid thioesters derived from acylated proteins accumulate in infantile neuronal ceroid lipofuscinosis: Correction of the defect in lymphoblasts by recombinant palmitoyl-protein thioesterase. *Proc. Natl. Acad. Sci. USA* **93(19)**, 10046-10050 (1996).

CAYMAN CHEMICAL
1180 EAST ELLSWORTH RD
ANN ARBOR, MI 48108 · USA
PHONE: [800] 364-9897
[734] 971-3335
FAX: [734] 971-3640
CUSTSERV@CAYMANCHEM.COM
WWW.CAYMANCHEM.COM