

# **Product Data Sheet**

### **Anti-PPT Antibody**

Catalog #	Source	Reactivity	Applications		
CPA4493	Rabbit	H, M, R, B, Mk	WB, IH, IF/IC		
Description	Rabb	it polyclonal antibody to	) PPT		
Immunogen	KLH-	conjugated synthetic pe	ptide encompassing a sequence within the N-term		
	regio	region of human PPT. The exact sequence is proprietary.			
Purification	The	antibody was purified by	immunogen affinity chromatography.		
Specificity	Reco	gnizes endogenous leve	ls of PPT protein.		
Clonality	Poly	Polyclonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium ph	osphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB (	1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	PPT1				
Alternative Na	ames PPT;	Palmitoyl-protein thioes	terase 1; PPT-1; Palmitoyl-protein hydrolase 1		
Entrez Gene	5538	s (Human)			
SwissProt	P508	97 (Human)			
Storage/Stabi	lity Ship	oed at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	freez	e/thaw cycles.			

Application key: E- ELISA, WB- Western blot, IH- Immunohistochemistry, IF- Immunofluorescence, FC- Flow cytometry, IC-Immunocytochemistry, IP- Immunoprecipitation, ChIP- Chromatin Immunoprecipitation, EMSA- Electrophoretic Mobility Shift Assay, BL- Blocking, SE- Sandwich ELISA, CBE- Cell-based ELISA, RNAi- RNA interference Species reactivity key: H- Human, M- Mouse, R- Rat, B- Bovine, C- Chicken, D- Dog, G- Goat, Mk- Monkey, P- Pig, Rb-Rabbit, S- Sheep, Z- Zebrafish

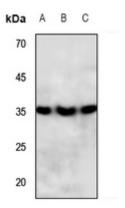
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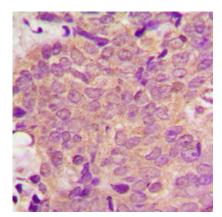


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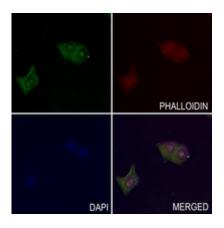
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Western blot analysis of PPT expression in mouse spleen (A), rat liver (B), rat spleen (C) whole cell lysates. (Predicted band size: 34 kD; Observed band size: 37 kD)



Immunohistochemical analysis of PPT staining in human breast cancer formalin fixed paraffin embedded tissue section. The section was pre-treated using heat mediated antigen retrieval with sodium citrate buffer (pH 6.0). The section was then incubated with the antibody at room temperature and detected using an HRP conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of PPT staining in LO2 cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4 °C in a hidified chamber. Cells were washed with PBST and incubated with a AF488-conjugated secondary antibody (green) in PBS at room temperature in the dark. Phalloidin -AF594 was used to stain Actin filaments (red). DAPI was used to stain the cell nuclei (blue).

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