

## **Product Data Sheet**

## **Anti-TMPase Antibody**

Catalog #	Source	Reactivity	Applications
CQA2775	Rabbit	H, M, R	WB, IH
Description	Rab	bit polyclonal antibody	to TMPase
Immunogen	Reco	ombinant full length pro	otein of human TMPase
Purification	The	antibody was purified l	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of TMPase protein.
Clonality	Poly	clonal	
Conjugation			
Form	Liqu	id in 0.42% Potassium	phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,
	and	0.01% sodium azide.	
Dilution	WB	(1/500 - 1/2000), IH (1/5	0 - 1/200)
Gene Symbol	ACP	Р	
Alternative Na	ames Pros	static acid phosphatase	PAP; 5'-nucleotidase; 5'-NT; Ecto-5'-nucleotidase;
	Thia	mine monophosphatas	e; TMPase
Entrez Gene	55 (	Human); 56318 (Mouse	e); 56780 (Rat)
SwissProt	P15	309 (Human); Q8CE08 (	Mouse); P20646 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ery aliquot and store at -20°C for one year. Avoid
	free	ze/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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kDa A

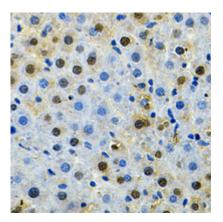
200

140

For research purposes only, not for human use

# **Product Data Sheet**

Western blot analysis of TMPase expression in MCF7 (A) whole cell lysates. (Predicted band size: 40; 44; 48 kD; Observed band size: 48 kD)



Immunohistochemical analysis of TMPase staining in human prostate 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.

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