

# **Product Data Sheet**

### **Anti-PTRH2** Antibody

Catalog #	Source	Reactivity	Applications		
CQA1391	Rabbit	H, M, R	WB, IH, IF/IC		
Description	Ra	bbit polyclonal antibody	to PTRH2		
Immunogen	Re	combinant full length pro	tein of human PTRH2		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity		Recognizes endogenous levels of PTRH2 protein.			
Clonality	Ро	olyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/2000), IH (1/50	) - 1/200), IF/IC (1/50 - 1/200)		
Gene Symbol	РТ	RH2			
Alternative Na	ames Bl <sup>-</sup>	T1; PTH2; Peptidyl-tRNA ł	ydrolase 2 mitochondrial; PTH 2; Bcl-2 inhibitor of		
	tra	anscription 1			
Entrez Gene		51651 (Human); 217057 (Mouse)			
SwissProt	Q	9Y3E5 (Human); Q8R2Y8 (	Mouse)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	ry aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

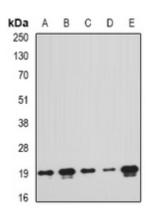
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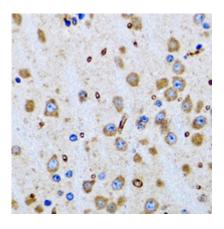


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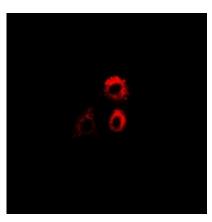
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Western blot analysis of PTRH2 expression in MCF7 (A), A549 (B), mouse liver (C), mouse spleen (D), rat brain (E) whole cell lysates. (Predicted band size: 19 kD; Observed band size: 19 kD)



Immunohistochemical analysis of PTRH2 staining in rat brain 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 PTRH2 staining in MCF7 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 DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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