

## **Product Data Sheet**

### Anti-PPP2R1B Antibody

Catalog #	Source	Reactivity	Applications		
CPA1922	Rabbit	H, M, R, Mk, P	WB, IH		
Description	R	abbit polyclonal antibody to	PPP2R1B		
Immunogen	К	LH-conjugated synthetic pep	tide encompassing a sequence within the C-term		
	re	egion of human PPP2R1B. Th	e exact sequence is proprietary.		
Purification	Т	he antibody was purified by	mmunogen affinity chromatography.		
Specificity	R	ecognizes endogenous levels	of PPP2R1B protein.		
Clonality Polyclonal					
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	nd 0.01% sodium azide.			
Dilution	V	VB (1/500 - 1/1000), IH (1/50 -	1/100)		
Gene Symbol	Р	PP2R1B			
Alternative N	ames So	erine/threonine-protein pho	sphatase 2A 65 kDa regulatory subunit A beta isoform;		
	Р	P2A subunit A isoform PR65-	beta; PP2A subunit A isoform R1-beta		
Entrez Gene	5	519 (Human); 73699 (Mouse	); 315648 (Rat)		
SwissProt	P	30154 (Human); Q7TNP2 (M	ouse); Q4QQT4 (Rat)		
Storage/Stability		Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid			
	fr	reeze/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

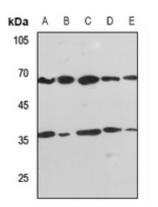
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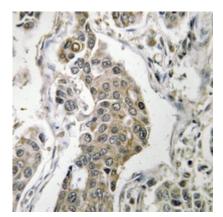




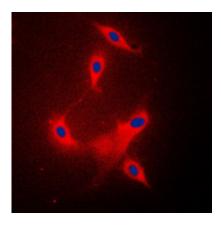
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Western blot analysis of PPP2R1B expression in HEK293T (A), Hela (B), HGC27 (C), mouse testis (D), rat testis (E) whole cell lysates. (Predicted band size: 66 kD; Observed band size: 66 kD)



Immunohistochemical analysis of PPP2R1B 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 PPP2R1B staining in Raw264.7 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark. DAPI was used to stain the cell nuclei (blue).

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