

Product Data Sheet

Anti-MYBPC2 Antibody

Catalog #	Source	e Reactivity	Applications			
CPA2986	Rabbit	H, M, R	WB, IH, IF/IC			
Description		Rabbit polyclonal antibody to MYBPC2				
Immunogen		KLH-conjugated synthetic peptide encompassing a sequence within the center				
		region of human MYBPC2. The exact sequence is proprietary.				
Purification		The antibody was purified by immunogen affinity chromatography.				
Specificity		Recognizes endogenous levels of MYBPC2 protein.				
Clonality		Polyclonal				
Conjugation						
Form	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycero					
		and 0.01% sodium azide.				
Dilution		WB (1/500 - 1/1000), IH (1/50 - 1/100), IF/IC (1/50 - 1/200)				
Gene Symbol		MYBPC2				
Alternative Na	ames	MYBPCF; Myosin-binding pr	otein C, fast-type; Fast MyBP-C; C-protein, skeletal			
		muscle fast isoform				
Entrez Gene		4606 (Human); 233199 (Mouse)				
SwissProt		Q14324 (Human); Q5XKE0 (Mouse)				
Storage/Stability Shi		Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid				
		freeze/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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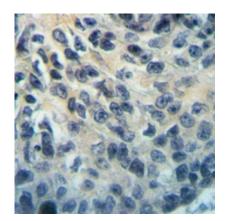
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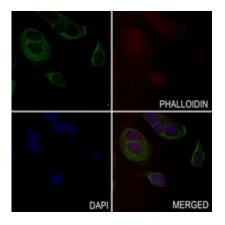
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Western blot analysis of MYBPC2 expression in HEK293T (A) whole cell lysates. (Predicted band size: 128 kD; Observed band size: 128 kD)



Immunohistochemical analysis of MYBPC2 staining in human lung 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 MYBPC2 staining in A549 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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