

Product Data Sheet

Anti-PFKFB2 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4151	Rabbit	H, Mk	WB, IH, IF/IC		
Description	Rab	bit polyclonal antibody	/ to PFKFB2		
Immunogen	KLH	-conjugated synthetic	peptide encompassing a sequence within the N-term		
	regi	on of human PFKFB2.	The exact sequence is proprietary.		
Purification	The	antibody was purified	by immunogen affinity chromatography.		
Specificity	Rec	ognizes endogenous le	vels of PFKFB2 protein.		
Clonality	Poly	/clonal			
Form	Liqu	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	and	0.01% sodium azide.			
Dilution	WB	(1/500 - 1/1000), IH (1/	100 - 1⁄200), IF/IC (1⁄100 - 1⁄500)		
Gene Symbol	PFK	FB2			
Alternative Na	ames 6-pł	nosphofructo-2-kinase	/fructose-2,6-bisphosphatase 2; 6PF-2-K/Fru-2,6-P2ase 2;		
	PFK	/FBPase 2; 6PF-2-K/Fru	I-2,6-P2ase heart-type isozyme		
Entrez Gene	520	5208 (Human); 18640 (Mouse); 24640 (Rat)			
SwissProt	060)825 (Human); P70265	(Mouse); Q9JJH5 (Rat)		
Storage/Stabi	lity Ship	oped at 4°C. Upon deliv	very 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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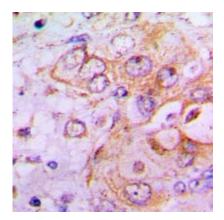
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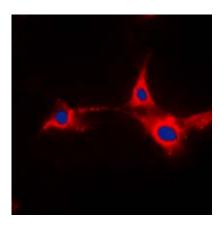
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Western blot analysis of PFKFB2 expression in HCT116 (A), NIH3T3 (B) whole cell lysates.



Immunohistochemical analysis of PFKFB2 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 conjugad compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.



Immunofluorescent analysis of PFKFB2 staining in HepG2 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. DAPI was used to stain the cell nuclei (blue).

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