

# **Product Data Sheet**

## **Anti-FUK Antibody**

Catalog #	Source	Reactivity	Applications		
CPA5393	Rabbit	H, M, R, Mk	WB, IH, IF/IC		
Description	F	Rabbit polyclonal antibody to	) FUK		
Immunogen	ķ	KLH-conjugated synthetic period	otide encompassing a sequence within the N-term		
	r	region of human FUK. The ex	act sequence is proprietary.		
Purification	T	The antibody was purified by	immunogen affinity chromatography.		
Specificity	F	Recognizes endogenous leve	s of FUK protein.		
Clonality	F	Polyclonal			
Conjugation					
Form		Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,			
	a	and 0.01% sodium azide.			
Dilution	١	WB (1/500 - 1/1000), IH (1/50	- 1/100), IF/IC (1/50 - 1/200)		
Gene Symbol	F	-UK			
Alternative Na	ames L	fucose kinase; Fucokinase			
Entrez Gene	1	197258 (Human)			
SwissProt	(	Q8N0W3 (Human)			
Storage/Stabi	lity S	Shipped at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	f	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

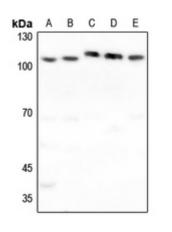
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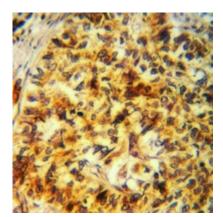


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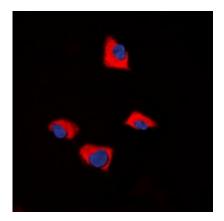
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Western blot analysis of FUK expression in Hela (A), A549 (B), mouse lung (C), mouse kidney (D), rat lung (E) whole cell lysates. (Predicted band size: 117 kD; Observed band size: 118 kD)



Immunohistochemical analysis of FUK staining in human stomach 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 FUK staining in NIH3T3 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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