

Product Data Sheet

Anti-PAK2 Antibody

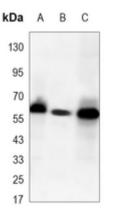
Catalog #	Source	Reactivity	Applications
CQA8390	Mouse	H, Mk	WB, IH, IF/IC
Description	Мо	use monoclonal antib	ody to PAK2
Immunogen	Pur	ified recombinant frag	ment of PAK2 expressed in E. Coli.
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Rec	ognizes endogenous l	evels of PAK2 protein.
Clonality	Мо	noclonal	
Conjugation			
Form	Liqu	uid in PBS containing 5	0% glycerol, 0.5% BSA and 0.02% sodium azide, pH 7.3.
Dilution	WB	(1/500 - 1/1000), IH (1,	/50 - 1/100), IF/IC (1/50 - 1/100)
Gene Symbol	PAk	(2	
Alternative Na	ames Ser	ine/threonine-protein	kinase PAK 2; Gamma-PAK; PAK65; S6/H4 kinase;
	p21	-activated kinase 2; P/	AK-2; p58
Entrez Gene	506	52 (Human)	
SwissProt	Q13	3177 (Human)	
Storage/Stabi	lity Shi	pped at 4°C. Upon deli	very aliquot and store at -20°C for one year. Avoid
	free	eze/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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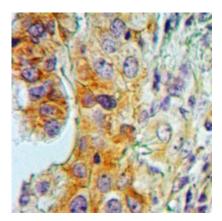




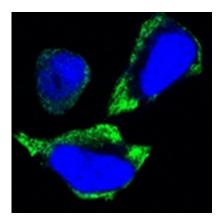
For research purposes only, not for human use

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Western blot analysis of PAK2 expression in Jurkat (A), MCF7 (B), Hela (C) whole cell lysates. (Predicted band size: 58 kD; Observed band size: 61 kD)



Immunohistochemical analysis of PAK2 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 PAK2 staining in HeLa 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.

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