

## **Product Data Sheet**

## Anti-PAK1 (pT212) Antibody

Catalog #	Source	Reactivity	Applications
CPA4143	Rabbit	H, M, R	WB, IH, IF/IC
Description	Rabl	oit polyclonal antibody	to PAK1 (pT212)
Immunogen	KLH-	conjugated synthetic p	eptide encompassing a sequence within the center
	regio	on of human PAK1 (pT2	12). The exact sequence is proprietary.
Purification	The	antibody was purified	by immunogen affinity chromatography.
Specificity	Reco	ognizes endogenous lev	els of PAK1 (pT212) protein.
Clonality	Poly	clonal	
Form	Liqu	id 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 (1/1	00 - 1/200), IF/IC (1/100 - 1/500)
Gene Symbol	PAK	1	
Alternative Na	ames Serir	ne/threonine-protein k	inase PAK 1; Alpha-PAK; p21-activated kinase 1; PAK-1;
	p65-	РАК	
Entrez Gene	5058	3 (Human); 29431 (Rat)	
SwissProt	Q13	153 (Human); O88643	(Mouse); P35465 (Rat)
Storage/Stabi	lity Ship	ped at 4°C. Upon delive	ery 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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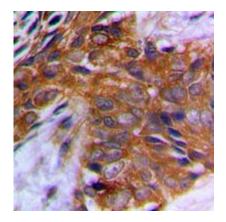
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# Coherion

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Western blot analysis of PAK1 (pT212) expression in HCT116 (A) whole cell lysates.



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Immunohistochemical analysis of PAK1 (pT212) staining in human prostate 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. w

Immunofluorescent analysis of PAK1 (pT212) 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 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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