

Product Data Sheet

Anti-NFAT4 (pS165) Antibody

Catalog #	Source	Reactivity	Applications	
CPA4119	Rabbit	H, M, Mk	WB, IH, IF/IC	
Description	ł	Rabbit polyclonal antibody t	o NFAT4 (pS165)	
Immunogen	ł	KLH-conjugated synthetic pe	ptide encompassing a sequence within the N-term	
	ı	region of human NFAT4 (pS1	65). The exact sequence is proprietary.	
Purification	-	The antibody was purified b	immunogen affinity chromatography.	
Specificity	I	Recognizes endogenous leve	ls of NFAT4 (pS165) protein.	
Clonality	I	Polyclonal		
Form	I	Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	ä	and 0.01% sodium azide.		
Dilution	١	WB (1/500 - 1/1000), IH (1/10	0 - 1/200), IF/IC (1/100 - 1/500)	
Gene Symbol	I	NFATC3		
Alternative N	ames I	NFAT4; Nuclear factor of act	vated T-cells, cytoplasmic 3; NF-ATc3; NFATc3; NFATx;	
	-	T-cell transcription factor NF	AT4; NF-AT4	
Entrez Gene	2	4775 (Human)		
SwissProt	(Q12968 (Human); P97305 (N	1ouse)	
Storage/Stabi	i lity S	Shipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid	
	f	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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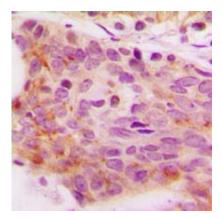
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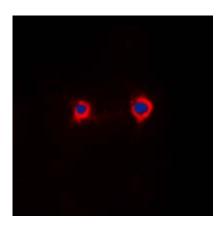
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Western blot analysis of NFAT4 (pS165) expression in Ramos (A), HeLa (B) whole cell lysates.



Immunohistochemical analysis of NFAT4 (pS165) staining in human breast 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 NFAT4 (pS165) 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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