

Product Data Sheet

Anti-STAT4 (Phospho-Y693) Antibody

Catalog #	Source	Reactivity	Applications		
CPA4838	Rabbit	H, M, Mk	WB, IH, IP		
Description	Rabb	Rabbit polyclonal antibody to STAT4 (Phospho-Y693)			
Immunogen	KLH-	conjugated synthetic ph	osphopeptide corresponding to residues surrounding		
	Y693	of human STAT4 protei	n. The exact sequence is proprietary.		
Purification	The	antibody was purified by	<i>immunogen affinity chromatography.</i>		
Specificity	Reco	gnizes endogenous leve	ls of STAT4 protein only when phosphorylated at Y693.		
Clonality	Poly	clonal			
Conjugation					
Form	Liqui	d in 0.42% Potassium pł	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	and	0.01% sodium azide.			
Dilution	WB (1/500 - 1/1000), IH (1/10	0 - 1⁄200), IP (1⁄10 - 1⁄100)		
Gene Symbol	STAT	4			
Alternative Na	ames Signa	al transducer and activat	tor of transcription 4		
Entrez Gene	6775	(Human); 20849 (Mous	e)		
SwissProt	Q147	765 (Human); P42228 (N	1ouse)		
Storage/Stabi	lity Ship	oed at 4°C. Upon deliver	y aliquot and store at -20°C for one year. Avoid		
	freez	e/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

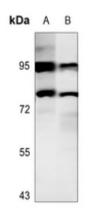
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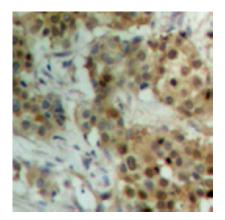
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For research purposes only, not for human use

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Western blot analysis of STAT4 (Phospho-Y693) expression in HEK293T (A), PC3 (B) whole cell lysates. (Predicted band size: 85 kD; Observed band size: 95 kD)



Immunohistochemical analysis of STAT4 (Phospho-Y693) 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 conjugated compact polymer system. DAB was used as the chromogen. The section was then counterstained with haematoxylin and mounted with DPX.

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