

Product Data Sheet

Anti-c-FER (Phospho-Y402) Antibody

Catalog #	Source	e Reactivity	Applications			
CPA4641	Rabbit	-	WB, IH			
Description						
Immunogen		KLH-conjugated synthetic phosphopeptide corresponding to residues surrounding				
Y402 of human c-FER protein. The exact sequence is proprietary.						
Purification		The antibody was purified by immunogen affinity chromatography.				
Specificity		Recognizes endogenous levels of c-FER protein only when phosphorylated at Y402.				
Clonality Polyclonal						
, Conjugation		,				
Form		Liquid 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)			
Gene Symbol		FER				
Alternative Names		TYK3; Tyrosine-protein kinase Fer; Feline encephalitis virus-related kinase FER;				
		Fujinami poultry sarcoma/F	eline sarcoma-related protein Fer; Proto-oncogene c-Fer;			
		Tyrosine kinase 3; p94-Fer				
Entrez Gene		2241 (Human); 14158 (Mou	ise); 301737 (Rat)			
SwissProt		P16591 (Human); P70451 (I	Mouse); P09760 (Rat)			
Storage/Stabi	ity Shipped at 4°C. Upon delivery aliquot and store at -20°C for one year. Avoid					
		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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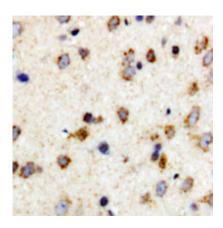
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Western blot analysis of c-FER (Phospho-Y402) expression in HeLa (A), Jurkat (B), NIH3T3 (C) whole cell lysates. (Predicted band size: 94 kD; Observed band size: 95 kD)



Immunohistochemical analysis of c-FER (Phospho-Y402) staining in human brain 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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