

Product Data Sheet

Anti-FOXO4 (Phospho-S197) Antibody

Catalog #	Source	Reactivity	Applications			
CPA1743	Rabbit	H, M, R, Mk, P	WB, IH, IF/IC			
Description	iption Rabbit polyclonal antibody to FOXO4 (Phospho-S197)					
Immunogen	KLH-	conjugated synthetic pho	osphopeptide corresponding to residues surrounding			
	S197	' of human FOXO4 protei	n. The exact sequence is proprietary.			
Purification	The	antibody was purified by	immunogen affinity chromatography.			
Specificity	Reco	ognizes endogenous level	s of FOXO4 protein only when phosphorylated at S197.			
Clonality	Poly	Polyclonal				
Conjugation						
Form	Liqu	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 (1/100) - 1/200), IF/IC (1/100 - 1/500)			
Gene Symbol	FOX	54				
Alternative Na	ames AFX;	AFX1; MLLT7; Forkhead	box protein O4; Fork head domain transcription factor			
	AFX	L				
Entrez Gene	4303	3 (Human); 54601 (Mouse	e)			
SwissProt	P98:	177 (Human); Q9WVH3 (I	Mouse)			
Storage/Stabi	lity Ship	ped at 4°C. Upon delivery	y 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

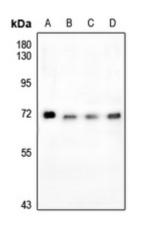
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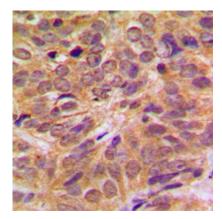


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Western blot analysis of FOXO4 (Phospho-S197) expression in LO2 (A), Panc1 (B), PC12 (C), AML12 (D) whole cell lysates. (Predicted band size: 53 kD; Observed band size: 70 kD)



Immunohistochemical analysis of FOXO4 (Phospho-S197) 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.

Immunofluorescent analysis of FOXO4 (Phospho-S197) 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 humidified chamber. Cells were washed with PBST and incubated with a DyLight 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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