

Product Data Sheet

Anti-FOXP1 Antibody

Catalog #	Source	Reactivity	Applications		
CPA4907	Rabbit	H, M, R, B	WB, IH		
Description	Ra	Rabbit polyclonal antibody to FOXP1			
Immunogen	KL	H-conjugated synthetic pe	ptide encompassing a sequence within the C-term		
	re	gion of human FOXP1. The	exact sequence is proprietary.		
Purification	Th	e antibody was purified b	y immunogen affinity chromatography.		
Specificity	Re	cognizes endogenous leve	ls of FOXP1 protein.		
Clonality	Ро	lyclonal			
Conjugation					
Form	Liq	ุนid in 0.42% Potassium p	nosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol,		
	an	nd 0.01% sodium azide.			
Dilution	W	B (1/500 - 1/1000), IH (1/50	- 1/100)		
Gene Symbol	FO)XP1			
Alternative N	ames Fo	rkhead box protein P1			
Entrez Gene 27086 (H		086 (Human); 108655 (M	Human); 108655 (Mouse); 297480 (Rat)		
SwissProt	QS	9H334 (Human); P58462 (I	Mouse); Q498D1 (Rat)		
Storage/Stabi	lity Sh	ipped at 4°C. Upon delive	y aliquot and store at -20°C for one year. Avoid		
	fre	eeze/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

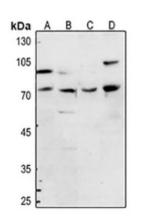
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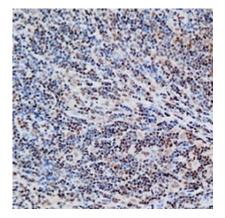


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Western blot analysis of FOXP1 expression in HEK293T (A), HGC27 (B), mouse lung (C), rat lung (D) whole cell lysates. (Predicted band size: 75 kD; Observed band size: 75 kD)



Immunohistochemical analysis of FOXP1 staining in human lymphoma 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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