For Research Use Only

FOXP1 Polyclonal antibody

Catalog Number:22051-1-AP 2 Publications

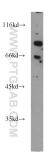


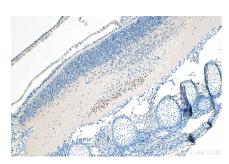
Basic Information	Catalog Number: 22051-1-AP	GenBank Accession Number: BC131720	Purification Method: Antigen affinity purification	
	Size:	GeneID (NCBI):	Recommended Dilutions:	
	150ul , Concentration: 800 ug/ml by Nanodrop; Source: Rabbit Isotype: IgG Immunogen Catalog Number: AG17045	27086	WB 1:500-1:2000	
		UNIPROT ID:	IHC 1:250-1:1000	
		Q9H334 Full Name: forkhead box P1		
				Calculated MW: 677 aa, 75 kDa
		Observed MW:		
		50 kDa, 60-65 kDa, 85 kDa		
		Applications	Tested Applications:	Positive Controls:
Cited Applications: MDA-MB-453s cells, mouse heart tissue			Irkat cells, A549 cells, LNCaP cells, MCF-7 cell	
			1B-453s cells, mouse heart tissue, mouse lung mouse testis tissue, PC-3 cells, Raji cells, Y79	
Species Specificity:	cells			
human, mouse, rat	IHC : m		nouse embryo tissue,	
Cited Species: human, mouse				
Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0				
	retrieval may be performed w	vely, antigen		
Background Information	retrieval may be performed w buffer pH 6.0 FOXP1, also known as Mac-1-regulat heterodimers with FOXP2 and FOXP4 important function in neuronal devel result in the development of autism as motor development delay. FOXP1	vely, antigen vith citrate ed forkhead, is a 677 amino acid (PubMed:25027557). Dimeriza opment.9 Mutations of its gene, spectrum disorder, intellectual of is also engaged in lung and esc ched role of FOXP1 in carcinoge	d protein, which forms homodimers and tion is required for DNA-binding. FOXP1 has a FOXP1, located on chromosome 3p14.1,7 car disability, speech and language deficits as we ophagus morphogenesis, as well as in B-cell enesis is of great importance, although still kDa, 65-67 kDa, 12 kDa.	
	retrieval may be performed w buffer pH 6.0 FOXP1, also known as Mac-1-regulat heterodimers with FOXP2 and FOXP4 important function in neuronal devel result in the development of autism as motor development delay. FOXP1 development.7,10 The widely resear unclear to some extent. FOXP1 existent	vely, antigen with citrate ed forkhead, is a 677 amino acid (PubMed:25027557). Dimeriza opment.9 Mutations of its gene, spectrum disorder, intellectual of is also engaged in lung and esc ched role of FOXP1 in carcinoge s some isoforms with MV 75-77	tion is required for DNA-binding. FOXP1 has a , FOXP1, located on chromosome 3p14.1,7 car disability, speech and language deficits as we ophagus morphogenesis, as well as in B-cell nesis is of great importance, although still kDa, 65-67 kDa, 12 kDa.	
Background Information Notable Publications	retrieval may be performed w buffer pH 6.0 FOXP1, also known as Mac-1-regulat heterodimers with FOXP2 and FOXP4 important function in neuronal devel result in the development of autism as motor development delay. FOXP1 development.7,10 The widely resear unclear to some extent. FOXP1 exists	vely, antigen vith citrate ed forkhead, is a 677 amino acid (PubMed:25027557). Dimeriza opment.9 Mutations of its gene, spectrum disorder, intellectual of is also engaged in lung and esc ched role of FOXP1 in carcinoge	tion is required for DNA-binding. FOXP1 has a , FOXP1, located on chromosome 3p14.1,7 car disability, speech and language deficits as we ophagus morphogenesis, as well as in B-cell enesis is of great importance, although still	
	retrieval may be performed w buffer pH 6.0 FOXP1, also known as Mac-1-regulat heterodimers with FOXP2 and FOXP4 important function in neuronal devel result in the development of autism as motor development delay. FOXP1 development.7,10 The widely resear unclear to some extent. FOXP1 exists Author Pub Liangyi Cui 333	vely, antigen with citrate ed forkhead, is a 677 amino acid (PubMed:25027557). Dimeriza: .opment.9 Mutations of its gene, spectrum disorder, intellectual of is also engaged in lung and eso ched role of FOXP1 in carcinoge s some isoforms with MV 75-77 med ID Journal	tion is required for DNA-binding. FOXP1 has a , FOXP1, located on chromosome 3p14.1,7 car disability, speech and language deficits as we ophagus morphogenesis, as well as in B-cell enesis is of great importance, although still kDa, 65-67 kDa, 12 kDa. Application WB	

For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll freeE: proteintech@ptglab.comin USA), or 1(312) 455-8498 (outside USA)W: ptglab.com

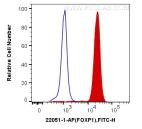
This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

Selected Validation Data





Jurkat cells were subjected to SDS PAGE followed by western blot with 22051-1-AP (FOXP1 antibody) at dilution of 1:1000 incubated at room temperature for 1.5 hours. Immunohistochemical analysis of paraffinembedded mouse embryo tissue slide using 22051-1-AP (FOXP1 antibody) at dilution of 1:500 (under 10x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0). Immunohistochemical analysis of paraffinembedded mouse embryo tissue slide using 22051-1-AP (FOXP1 antibody) at dilution of 1:500 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



1X10^6 Jurkat cells were intracellularly stained with 0.4 ug Anti-Human FOXP1 (22051-1-AP) and CoraLite®594-Conjugated Goat Anti-Rabbit IgG(H+L) at dilution 1:1000 (red), or 0.4 ug x. Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).