

FOXP3 Polyclonal Antibody, DyLight™ 550

Catalog NumberPA5-22776

Product data sheet

Details		Species Reactivity	
Size	100 µL	Species reactivity	Bovine, Human, Mouse, Rat
Host/Isotope	Rabbit / IgG	Tested Applications	Dilution *
Class	Polyclonal		
Type	Antibody		
Immunogen	A peptide corresponding to amino acids 43-100 of mouse FOXP3		
Conjugate	DyLight™ 550		
Form	Liquid	Flow Cytometry (Flow)	1:8,000
Concentration	0.84 mg/mL	Immunohistochemistry (Paraffin) (IHC (P))	Assay-Dependent
Purification	Protein G	Western Blot (WB)	1:2,000
Storage buffer	50mM sodium borate	Immunocytochemistry (ICC/IF)	Assay-Dependent
Contains	0.05% sodium azide	* Suggested working dilutions are given as a guide only. It is recommended that the user titrate the product for use in their own experiment using appropriate negative and positive controls.	
Storage Conditions	4° C, store in dark		

Product specific information

This antibody has shown reactivity against Ms (CD4+CD25+ T cells), does not react with Hu Jurkat, Ms CD4+CD25- T cell lysates.

Background/Target Information

FOXP3 (Forkhead box protein 3) is a member of the forkhead/winged-helix family of transcriptional regulators, highly conserved across mammals, and essential for normal immune homeostasis. FOXP3 is 381 amino acids long, stably and constitutively expressed at a high level in CD25 + CD4 positive regulatory T cells, a low level in CD4-positive/CD25-negative cells, and is absent in CD4-negative/CD8-positive T cells. FOXP3 may be a master regulatory gene, and a more specific marker of regulatory T cells. Defects in the gene encoding FOXP3 protein cause the scurfy phenotype in mice. In humans FOXP3 defects play a role in IPEX syndrome (immune dysfunction, polyendocrinopathy, enteropathy, X-linked syndrome), also known as X-linked autoimmunity-allergic dysregulation (XLAAD) syndrome. Transcript variants of FOXP3 encoding different isoforms have been identified. In human breast and colon cancer cells, expression of FOXP3 is regulated by p53 in response to the DNA damage.

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