For Research Use Only

CoraLite® Plus 488 Anti-Mouse Foxp3 (3G3)

Catalog Number: CL488-65089

1 Publications

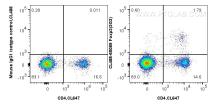


Basic Information	Catalog Number: CL488-65089	GenBank Accession Number: BC 132333	Purification Method: Affinity purification
	Size: 100ug , 500 µg/ml	GeneID (NCBI): 20371	CloneNo.: 3G3
	Source: Mouse	Full Name: forkhead box P3	Excitation/Emission maxima wavelengths:
	lsotype: IgG1, kappa		493 nm / 522 nm
Applications	Tested Applications: FC (Intra)		
	Cited Applications: FC (Intra)		
	Species Specificity: Mouse		
	Cited Species: mouse		
	Foxp3 (forkhead box P3), also named as IPEX, JM2 and scurfin, is a member of the forkhead transcription factor family and is essential for normal immune homeostasis (PMID: 11138001; 20429413). Foxp3 is specifically expressed in CD4+ CD25+ regulatory T cells and is crucial for their development and suppressive activity (PMID: 12612578; 12612581). In scurfy mouse, disruption of Foxp3 results in the fatal lymphoproliferative disorder attributed to the lack of an immune-suppressive component in the immune system (PMID: 25683611; 11138001). Similar to mice, genetic defects in human Foxp3 are the cause of immunodeficiency polyendocrinopathy, enteropathy, X-linked syndrome (IPEX) which also known as X-linked autoimmunity-immunodeficiency syndrom (PMID: 11137993).		
Background Information	expressed in CD4+ CD25+ re 12612578; 12612581). In scu attributed to the lack of an in Similar to mice, genetic def enteropathy, X-linked syndre	gulatory T cells and is crucial for their de rfy mouse, disruption of Foxp3 results in nmune-suppressive component in the im ects in human Foxp3 are the cause of imm	8001; 20429413). Foxp3 is specifically velopment and suppressive activity (PMID the fatal lymphoproliferative disorder mune system (PMID: 25683611; 11138001) nunodeficiency polyendocrinopathy,
	expressed in CD4+ CD25+ re 12612578; 12612581). In scu attributed to the lack of an in Similar to mice, genetic def enteropathy, X-linked syndre	gulatory T cells and is crucial for their de rfy mouse, disruption of Foxp3 results in nmune-suppressive component in the im ects in human Foxp3 are the cause of imm	8001; 20429413). Foxp3 is specifically velopment and suppressive activity (PMID: the fatal lymphoproliferative disorder mune system (PMID: 25683611; 11138001) nunodeficiency polyendocrinopathy,
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For technical support and original validation data for this product please contact:T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free
in USA), or 1(312) 455-8498 (outside USA)E: proteintech@ptglab.comW: ptglab.comW: ptglab.com

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

Selected Validation Data



1x10^{^6} mouse splenocytes were intracellularly stained with 0.06 ug CoraLite® Plus 488 Anti-Mouse Foxp3 (CL488-65089, Clone:3G3), and 0.06 ug CoraLite® Plus 647 Anti-Mouse CD4 (GK1.5) (CL647-65104, Clone: GK1.5), and 0.06 ug CoraLite® Plus 488 Mouse IgG1 Isotype Control (MOPC-21) (CL488-65124, Clone: MOPC-21). Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).