





# FOXP3 Monoclonal Antibody (3G3), Alexa Fluor™ 647

Catalog Number MA5-18160 Product data sheet

Details	
Size	100 μg
Host/Isotope	Mouse / IgG1, kappa
Class	Monoclonal
Туре	Antibody
Clone	3G3
Immunogen	Full-length His-tagged recombinant murine FoxP3
Conjugate	Alexa Fluor™ 647
Form	Liquid
Concentration	0.5 mg/mL
Purification	Size-exclusion chromatography
Storage buffer	PBS, pH 7.4
Contains	15mM sodium azide
Storage Conditions	4° C, store in dark, DO NOT FREEZE!

Species Reactivity	
Species reactivity	Human, Mouse
Published species	Mouse, Not Applicable
Total Androdica	D2 4
Tested Applications	Dilution *
Flow Cytometry (Flow)	1-5 μg/mL
Western Blot (WB)	Assay-dependent
Published Applications	
Flow Cytometry (Flow)	See 1 publications below

\* 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.

## 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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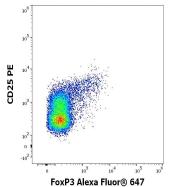
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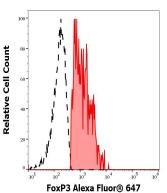


# Product Images For FOXP3 Monoclonal Antibody (3G3), Alexa Fluor™ 647



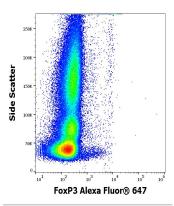
### FOXP3 Antibody (MA5-18160) in Flow

Flow cytometry multicolor surface staining of human CD4 positive lymphocytes using anti-human CD25 (MEM-181) PE antibody (20  $\mu$ L reagent / 100  $\mu$ L of peripheral whole blood) and intracellular staining of human CD4 positive lymphocytes using anti-human FoxP3 (3G3) Alexa Fluor® 647 Monoclonal antibody (Product # MA5-18160), concentration in sample 3  $\mu$ g/mL.



# FOXP3 Antibody (MA5-18160) in Flow

Separation of FoxP3 positive CD25 positive CD4 positive lymphocytes (red-filled) from FoxP3 negative CD25 negative CD4 positive lymphocytes (black-dashed) in flow cytometry analysis (intracellular staining) of human peripheral whole blood stained using anti-human FoxP3 (3G3) Alexa Fluor® 647 Monoclonal antibody (Product # MA5-18160), concentration in sample 3 µg/mL.



### FOXP3 Antibody (MA5-18160) in Flow

Flow cytometry intracellular staining pattern of human peripheral whole blood stained using anti-human FoxP3 (3G3) Alexa Fluor® 647Monoclonal antibody (Product # MA5-18160), concentration in sample 3 µg/mL.

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# PubMed References For FOXP3 Monoclonal Antibody (3G3), Alexa Fluor™ 647 1 Flow Cytometry References Species / Dilution MA5-18160 was used in Flow cytometry/Cell sorting to identify a small-molecule inhibitor that can be used to block type I IFN signaling during established infection and acts synergistically with conventional anti-parasitic drugs to improve parasite clearance and enhance anti-parasitic CD4+ T cell responses in mice and humans. Cell reports ( 2020; 30: 2512) "Type I Interferons Suppress Anti-parasitic Immunity and Can Be Targeted to Improve Treatment of Visceral Leishmaniasis." Author(s):Kumar R,Bunn PT,Singh SS,Ng SS,Montes de Oca M,De Labastida Rivera F,Chauhan SB,Singh N,Faleiro RJ, Edwards CL,Frame TCM,Sheel M,Austin RJ,Lane SW,Bald T,Smyth MJ,Hill GR,Best SE,Haque A,Corvino D,Waddell N, Koufariotis L,Mukhopadhay P,Rai M,Chakravarty J,Singh OP,Sacks D,Nylen S,Uzonna J,Sundar S,Engwerda CR PubMed Article URL:http://dx.doi.org/10.1016/j.celrep.2020.01.099

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