FOXP3 Rabbit mAb

Catalog No	YP-Ab-17676
Isotype	IgG
Reactivity	Human
Applications	WB,FC
Gene Name	FOXP3
Alternative Names	FOXP3; IPEX; JM2; Forkhead box protein P3; Scurfin
Research Field	Defects in FOXP3 are the cause of immunodeficiency polyendocrinopathy, enteropathy, X-linked syndrome (IPEX) [MIM:304790]; also known as X-linked autoimmunity-immunodeficiency syndrome. IPEX is characterized by neonatal onset insulin-dependent diabetes mellitus, infections, secretory diarrhea, trombocytopenia, anemia and eczema. It is usually lethal in infancy.
Product Categories	Primary antibody
Host	Rabbit
Molecular Weight	Calculated MW: 47 kDa; Observed MW: 47 kDa
Clonality	Monoclonal Antibody
Clonality No.	R03-8G4
Dilution	WB: 1/500-1/1000 FC: 1/50-1/100
Immunogen	A synthesized peptide derived from human FOXP3 Purification Affinity Chromatography Conjugation Unconjugated Modification Unmodified Form Liquid
Buffer System	Rabbit IgG in phosphate buffered saline , pH 7.4, 150mM NaCl, 0.02% sodium azide and 50% glycerol.
Concentration	1 mg/ml
Purity	≥90%
Storage	Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.
Background	
matters needing attention	Avoid repeated freezing and thawing!
Usage suggestions	This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images