

**FoxP3 Monoclonal Antibody**

<b>Catalog No :</b>	YM0286
<b>Reactivity :</b>	Human;Mouse
<b>Applications :</b>	WB;IHC;IF;ELISA
<b>Target :</b>	FoxP3
<b>Fields :</b>	>>Th17 cell differentiation;>>Inflammatory bowel disease
<b>Gene Name :</b>	FOXP3
<b>Protein Name :</b>	Forkhead box protein P3
<b>Human Gene Id :</b>	50943
<b>Human Swiss Prot No :</b>	Q9BZS1
<b>Mouse Gene Id :</b>	20371
<b>Mouse Swiss Prot No :</b>	Q99JB6
<b>Immunogen :</b>	Purified recombinant fragment of human FoxP3 expressed in E. Coli.
<b>Specificity :</b>	FoxP3 Monoclonal Antibody detects endogenous levels of FoxP3 protein.
<b>Formulation :</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
<b>Source :</b>	Monoclonal, Mouse
<b>Dilution :</b>	WB 1:500 - 1:2000. IHC 1:200 - 1:1000. IF 1:200 - 1:1000. ELISA: 1:10000. Not yet tested in other applications.
<b>Purification :</b>	Affinity purification
<b>Concentration :</b>	1 mg/ml

**Storage Stability :** -15°C to -25°C/1 year(Do not lower than -25°C)

**Molecularweight :** 47kD

**P References :** 1. Roncador G et al. Eur J Immunol. 2005. 35:1681-1691.  
2. Yisong YW. PNAS. 2005 102 (14): 5126-5131.

**Background :** The protein encoded by this gene is a member of the forkhead/winged-helix family of transcriptional regulators. Defects in this gene are the cause of immunodeficiency polyendocrinopathy, enteropathy, X-linked syndrome (IPEX), also known as X-linked autoimmunity-immunodeficiency syndrome. Alternatively spliced transcript variants encoding different isoforms have been identified. [provided by RefSeq, Jul 2008],

**Function :** disease: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.,function:Probable transcription factor. Plays a critical role in the control of immune response.,online information:FOXP3 entry,online information:FOXP3 mutation db,similarity:Contains 1 C2H2-type zinc finger.,similarity:Contains 1 fork-head DNA-binding domain.,

**Subcellular Location :** Nucleus . Cytoplasm . Predominantly expressed in the cytoplasm in activated conventional T-cells whereas predominantly expressed in the nucleus in regulatory T-cells (Treg). The 41 kDa form derived by proteolytic processing is found exclusively in the chromatin fraction of activated Treg cells (By similarity). .

**Tag :** orthogonal

**Sort :** 1

**No3 :** ab20034

**No4 :** 1

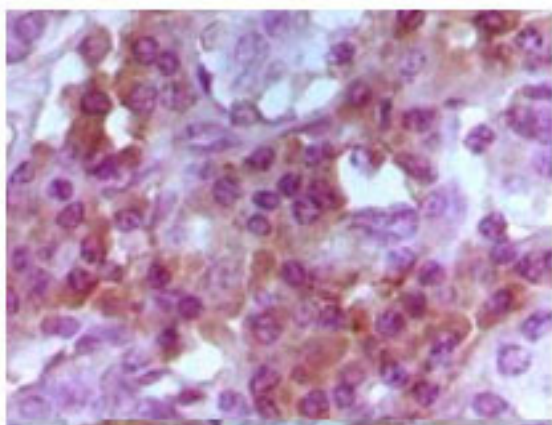
**Host :** Mouse

**Modifications :** Unmodified

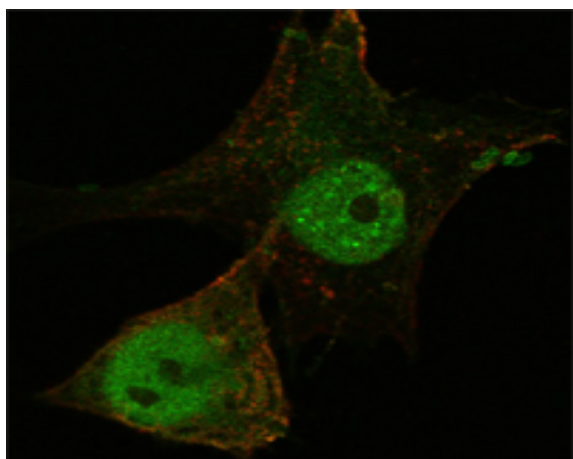
**Products Images**



Western Blot analysis using FoxP3 Monoclonal Antibody against truncated Foxp3 recombinant (1) and HEK293 cell lysate (2).



Immunohistochemistry analysis of paraffin-embedded human lymphocyte tissue, showing cytoplasmic and nuclear localization with DAB staining using FoxP3 Monoclonal Antibody.



Confocal immunofluorescence analysis of PANC-1 cells using FoxP3 Monoclonal Antibody (green). Red: Actin filaments have been labeled with Alexa Fluor-555 phalloidin.