

## **FoxP3 Monoclonal Antibody**

YM0286 Catalog No:

Human; Mouse Reactivity:

**Applications:** WB;IHC;IF;ELISA

Target: FoxP3

Fields: >>Th17 cell differentiation;>>Inflammatory bowel disease

**Gene Name:** FOXP3

**Protein Name:** Forkhead box protein P3

Q9BZS1

Q99JB6

**Human Gene Id:** 50943

**Human Swiss Prot** 

No:

Mouse Gene Id: 20371

**Mouse Swiss Prot** 

No:

Purified recombinant fragment of human FoxP3 expressed in E. Coli. Immunogen:

FoxP3 Monoclonal Antibody detects endogenous levels of FoxP3 protein. **Specificity:** 

**Formulation:** Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.

Monoclonal, Mouse Source:

**Dilution:** WB 1:500 - 1:2000, IHC 1:200 - 1:1000, IF 1:200 - 1:1000, ELISA: 1:10000, Not

yet tested in other applications.

**Purification:** Affinity purification

**Concentration:** 1 mg/ml

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**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

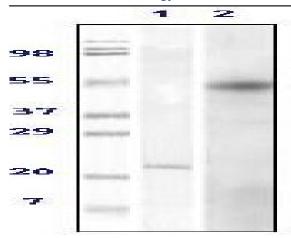
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**No3**: ab20034

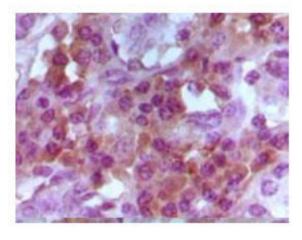
No4:

## **Products Images**

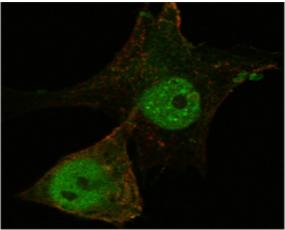
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Western Blot analysis using FoxP3 Monoclonal Antibody against truncated Foxp3 recombinant (1) and HEK293 cell lysate (2).



Immunohistochemistry analysis of paraffin-embedded human lymphocyto 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.