



Rabbit Anti-FoxP3 antibody

SL0269R

Product Name:	FoxP3
Chinese Name:	叉头蛋白P3抗体
Alias:	AIID; AIID; DIETER; DIETER; Forkhead box P3; Forkhead box protein P3; FOXP3_HUMAN; FOXP3delta7; Immune dysregulation polyendocrinopathy enteropathy X linked; Immunodeficiency polyendocrinopathy enteropathy X linked; IPEX; IPEX; JM2; JM2; MGC141961; MGC141963; OTTHUMP00000025832; OTTHUMP00000025833; OTTHUMP00000226737; PIDX; PIDX; SCURFIN; SCURFIN; XPID; XPID.
Organism Species:	Rabbit
Clonality:	Polyclonal
React Species:	Human,Mouse,Rat,
Applications:	ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800Flow-Cyt= 3µg/TestIF=1:100-500 (Paraffin sections need antigen repair) not yet tested in other applications. optimal dilutions/concentrations should be determined by the end user.
Molecular weight:	47kDa
Cellular localization:	The nucleus
Form:	Lyophilized or Liquid
Concentration:	1mg/ml
immunogen:	KLH conjugated synthetic peptide derived from human FoxP3:11-100/431
Lsotype:	IgG
Purification:	affinity purified by Protein A
Storage Buffer:	0.01M TBS(pH7.4) with 1% BSA, 0.03% Proclin300 and 50% Glycerol.
Storage:	Store at -20 °C for one year. Avoid repeated freeze/thaw cycles. The lyophilized antibody is stable at room temperature for at least one month and for greater than a year when kept at -20°C. When reconstituted in sterile pH 7.4 0.01M PBS or diluent of antibody the antibody is stable for at least two weeks at 2-4 °C.
PubMed:	PubMed
Product Detail:	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:

Probable transcription factor. Plays a critical role in the control of immune response.

Subunit:

Interacts with IKZF3.

Subcellular Location:

Nucleus (Potential).

Post-translational modifications:

Acetylation on lysine residues stabilizes FOXP3 and promotes differentiation of T-cells into induced regulatory T-cells (iTregs) associated with suppressive functions. Deacetylated by SIRT1.

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, thrombocytopenia, anemia and eczema. It is usually lethal in infancy.

Similarity:

Contains 1 C2H2-type zinc finger.
Contains 1 fork-head DNA-binding domain.

SWISS:

Q9BZS1

Gene ID:

50943

Database links:

[Entrez Gene: 50943](#)Human

[Entrez Gene: 20371](#)Mouse

[Entrez Gene: 317382](#)Rat

[Oimim: 300292](#)Human

[SwissProt: Q9BZS1](#)Human

[SwissProt: Q99JB6](#)Mouse

[SwissProt: D3ZKI1](#)Rat

[Unigene: 247700](#)Human

[Unigene: 182291](#)Mouse

Important Note:

This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

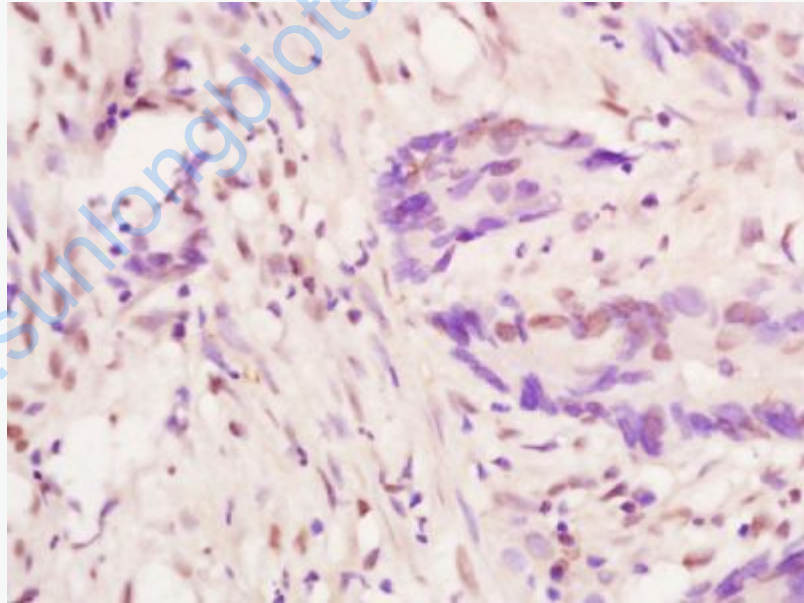
叉头蛋白3(FOXP3)是FOX蛋白家族成员之一,主要为T细胞转录蛋白,主要表达于T细胞+CD4+CD25,并调节该类T细胞的发育和功能.

FOXP3的表达受转化生长因子-

β 雌激素和糖皮质激素等调节,通过竞争性抑制活化TThe

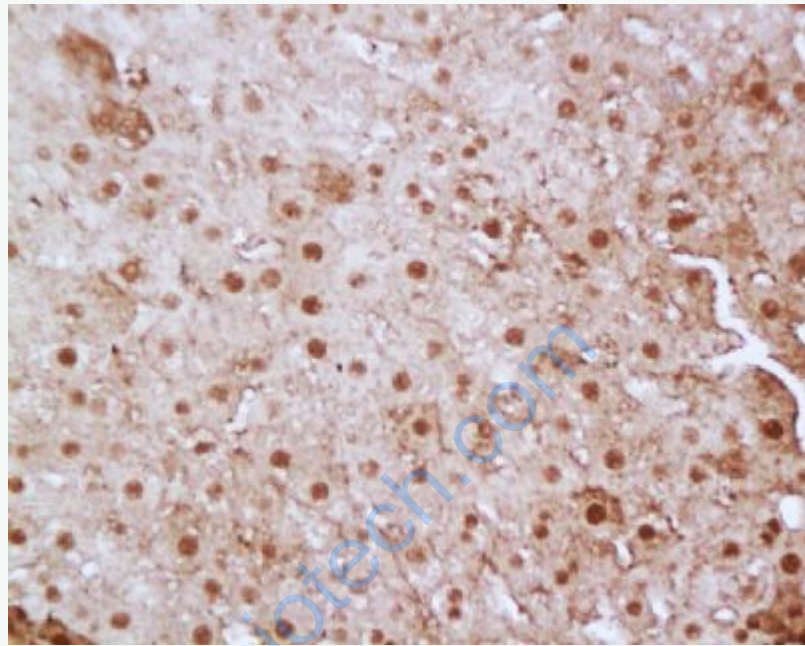
nucleus因子的转录活性而发挥作用.自身免疫性Diabetes患者体内CD4+CD25+T细胞减少,诱导FOXP3的表达或过继转移CD4+CD25+T细胞有可能预防自身免疫性Diabetes

Picture:

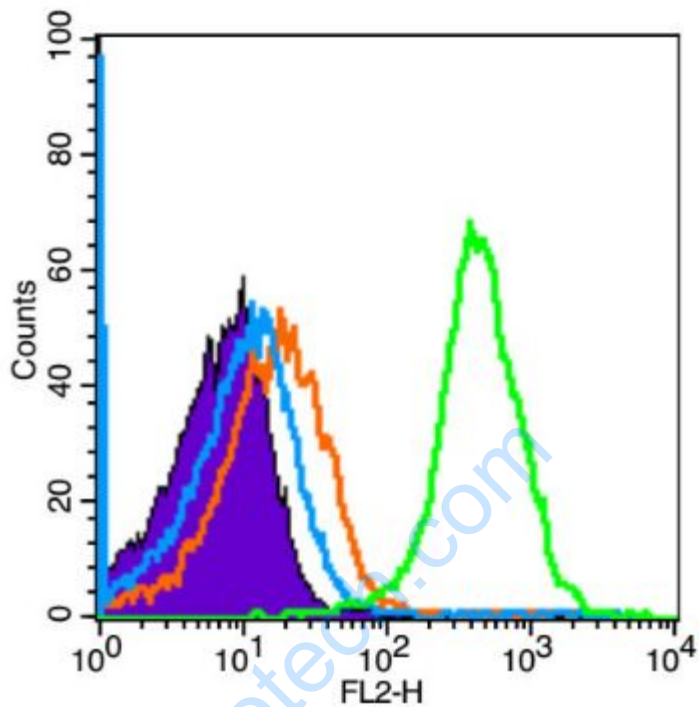


Paraformaldehyde-fixed, paraffin embedded (Human colon cancer); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (FoxP3) Polyclonal Antibody, Unconjugated (SL0269R) at 1:200 overnight at 4°C, followed by operating

according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Paraformaldehyde-fixed, paraffin embedded (Rat liver); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (FoxP3) Polyclonal Antibody, Unconjugated (SL0269R) at 1:200 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructions and DAB staining.



Blank control (Black line): Mouse spleen (Black).

Primary Antibody (green line): Rabbit Anti-FoxP3 antibody (SL0269R)

Dilution: $3\mu\text{g} / 10^6$ cells;

Isotype Control Antibody (orange line): Rabbit IgG .

Secondary Antibody (white blue line): Goat anti-rabbit IgG-PE

Dilution: $1\mu\text{g} / \text{test}$.

Protocol

The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at room temperature. The cells were then incubated in 5% BSA goat serum to block non-specific protein-protein interactions for 15 min at room temperature. Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min

	at room temperature. Acquisition of 20,000 events was performed.
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