

Rabbit Anti-TGF Beta 1+2+3 antibody

SL4538R

| Product Name: | TGF Beta 1+2+3 |
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| Chinese Name: | 转化生长因子β1抗体 |
| Alias: | TGF Beta 1/2/3; TGFβ1; TGFβ2; TGFβ3; CED; Diaphyseal dysplasia 1 progressive; DPD 1; DPD1; TGF beta 1; TGF beta; TGF beta1; TGF beta2; TGF beta3; TGF-beta 1; TGF-beta 2; TGF-beta 3; TGFB 1; TGFB; TGFB1; TGFB2; TGFB3; Transforming growth factor; Transforming growth factor beta 1; Transforming growth factor beta 2; Transforming growth factor beta 3; Transforming Growth Factor beta. |
| Organism Species: | Rabbit |
| Clonality: | Polyclonal |
| React Species: | Human,Mouse,Rat,Dog,Pig,Cow,Rabbit,Sheep,Guinea Pig, |
| Applications: | WB=1:500-2000ELISA=1:500-1000IHC-P=1:400-800IHC-F=1:400-800IF=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: | 12.8/44kDa |
| Cellular localization: | Extracellular matrixSecretory protein |
| Form: | Lyophilized or Liquid |
| Concentration: | 1mg/ml |
| immunogen: | KLH conjugated synthetic peptide derived from human TGF Beta 1:301-350/390 |
| 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 TGFBeta superfamily is composed of numerous growth and differentiation factors, including transforming growth factors Beta(TGFBeta)1,2 and 3; placental transforming growth factor (PTGFBeta); growth /differentiation factors (GDFs); mullerian inhibiting |

substance (MIS);bone morphogenic proteins (BMPs); glial cell line-derived neurotrophic factor (GDNF);inhibins or activins (Alpha,Beta-A, Beta-B and Beta-C),Lefty and Nodal. Members of the TGF superfamily are involved in embryonic development and adult tissue homeostasis.

Function:

Multifunctional protein that controls proliferation, differentiation and other functions in many cell types. Many cells synthesize TGFB1 and have specific receptors for it. It positively and negatively regulates many other growth factors. It plays an important role in bone remodeling as it is a potent stimulator of osteoblastic bone formation, causing chemotaxis, proliferation and differentiation in committed osteoblasts.

Subunit:

Homodimer; disulfide-linked, or heterodimer with TGFB2. Secreted and stored as a biologically inactive form in the extracellular matrix in a 290 kDa complex (large latent TGF-beta1 complex) containing the TGFB1 homodimer, the latency-associated peptide (LAP), and the latent TGFB1 binding protein-1 (LTBP1). The complex without LTBP1 is known as the'small latent TGF-beta1 complex'. Dissociation of the TGFB1 from LAP is required for growth factor activation and biological activity. Release of the large latent TGF-beta1 complex from the extracellular matrix is carried out by the matrix metalloproteinase MMP3. May interact with THSD4; this interaction may lead to sequestration by FBN1 microfibril assembly and attenuation of TGFB signaling. Interacts with the serine proteases, HTRA1 and HTRA3: the interaction with either inhibits TGFB1-mediated signaling. The HTRA protease activity is required for this inhibition. Interacts with CD109, DPT and ASPN.

Subcellular Location:

Secreted, extracellular space, extracellular matrix.

Tissue Specificity:

Highly expressed in bone. Abundantly expressed in articular cartilage and chondrocytes and is increased in osteoarthritis (OA). Co-localizes with ASPN in chondrocytes within OA lesions of articular cartilage.

Post-translational modifications:

Glycosylated.

The precursor is cleaved into mature TGF-beta-1 and LAP, which remains noncovalently linked to mature TGF-beta-1 rendering it inactive.

DISEASE:

Defects in TGFB1 are the cause of Camurati-Engelmann disease (CE) [MIM:131300]; also known as progressive diaphyseal dysplasia 1 (DPD1). CE is an autosomal dominant disorder characterized by hyperostosis and sclerosis of the diaphyses of long bones. The disease typically presents in early childhood with pain, muscular weakness and waddling gait, and in some cases other features such as exophthalmos, facial paralysis, hearing difficulties and loss of vision.

Similarity: Belongs to the TGF-beta family.

SWISS: P01137

Gene ID: 7040

Database links:

Entrez Gene: 7040 Human

Entrez Gene: 21803 Mouse

Entrez Gene: 59086 Rat

<u>Omim: 190180</u> Human

SwissProt: P01137 Human

SwissProt: P04202 Mouse

SwissProt: P17246 Rat

Unigene: 645227 Human

Unigene: 248380 Mouse

Unigene: 40136 Rat

Important Note:

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

Growth factors and hormones (Growth Factor and Hormones)

TGF是一种多效生长因子, 对epithelial

cells增值有潜在抑制作用, 可抑制Tumour生长用于许多恶性Tumour如:胃癌、肺癌、 膀胱癌、肾癌、前列腺癌、结肠癌等多种恶性Tumour的研究。

otech.com

TGFβ超级**家族由**为**数众多的生长的分化因子组成,包括转移生长因子**β1, 2和3(TGF β1, TGF β2, TGF β3);胎盘生长因子(PTGF-

β);生长/分化因子(GDFs);缪氏抑制物(MIS);骨形态形成蛋白(BMPs);交织细胞元 神经生长因子(GDNF);抑制素和活化素(α, β-A,和β-C), Lefty和Nodal。





