Product datasheet

Anti-TGF beta 1 antibody ab92486

Overview

Product name: Anti-TGF beta 1 antibody
Description: Rabbit polyclonal to TGF beta 1
Host species: Rabbit
Specificity: Full length, inactive 44 kD TGFβ1 is cleaved into mature TGFβ1 (13 kD). TGFβ1 also homodimerizes and heterodimerizes with TGFβ2, so there is potential for multiple different band sizes in WB.

Tested applications: Suitable for: WB, IHC-FrFl, IHC-P, IHC-Fr, ICC/IF
Species reactivity: Reacts with: Mouse, Rat, Human, Pig
Positive control: WB: RAW 264.7 whole cell lysate (ab7187), Mouse 3T3 cell lysate. IHC-P: Mouse intestine tissue; human retina tissue.

General notes

Properties

Form: Liquid
Storage instructions: Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid repeated freeze / thaw cycles.
Storage buffer: pH: 7.20
Preservative: 0.01% Thimerosal (merthiolate)
 Constituents: 0.5% BSA, 30% Glycerol, 0.15% EDTA, PBS
Purity: Immunogen affinity purified
Clonality: Polyclonal
Isotype: IgG

Applications

Our Abpromise guarantee covers the use of ab92486 in the following tested applications.
The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.
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.

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.

Involvement in 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.

Sequence similarities
Belongs to the TGF-beta family.

Post-translational modifications
Glycosylated.
The precursor is cleaved into mature TGFB-beta-1 and LAP, which remains non-covalently linked to mature TGFB-beta-1 rendering it inactive.

Cellular localization
Secreted > extracellular space > extracellular matrix.

Notes
Use a concentration of 0.5 - 4 µg/ml. Predicted molecular weight: 44 kDa. Can be blocked with Human TGF beta 1 peptide (ab231650).
Full length, inactive 44 kD TGFB1 is cleaved into mature TGFB1 (13 kD). TGFB1 also homodimerizes and heterodimerizes with TGFB2, so there is potential for multiple different band sizes in WB.

Use at an assay dependent concentration. PubMed: 24647450
Use a concentration of 10 - 20 µg/ml.
Use a concentration of 10 - 20 µg/ml.
1/100.
Western blot of mouse 3T3 cell lysate with ab92486 at a concentration of 4 µg/ml.

ab92486 staining TGF beta 1 in Mouse intestine tissue sections by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections). Tissue was fixed with paraformaldehyde and blocked with 5% BSA for 30 minutes at 20°C; antigen retrieval was by heat mediation in a citrate buffer. Samples were incubated with primary antibody (1/100 in blocking buffer) for 24 hours at 4°C. An undiluted HRP-conjugated Human anti-rabbit polyclonal was used as the secondary antibody.
ab92486 staining TGF beta 1 in Human retina membrane by Immunohistochemistry (IHC-P - paraformaldehyde-fixed, paraffin-embedded sections).

Tissue was fixed with paraformaldehyde and blocked with 10% serum for 30 minutes at 20°C; antigen retrieval was by heat mediation. Samples were incubated with primary antibody (1/400 in PBS + 1% Triton X-100 + 2% goat serum) for 24 hours at 4°C. An HRP-conjugated Human polyclonal was used as the secondary antibody.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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