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 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.

Tested applications
Suitable for: WB, IHC-FrFl, IHC-P, IHC-Fr, ICC/IF

Species reactivity
Reacts with: Mouse, Rat, Human, Pig

Immunogen

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 TGFB1-1 and LAP, which remains non-covalently linked to mature TGFB1-1 rendering it inactive.

Cellular localization

Secreted > extracellular space > extracellular matrix.

### Application | Abreviews | Notes

<table>
<thead>
<tr>
<th>WB</th>
<th></th>
<th>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.</th>
</tr>
</thead>
<tbody>
<tr>
<td>IHC-FrFI</td>
<td></td>
<td>Use at an assay dependent concentration. PubMed: 24647450</td>
</tr>
<tr>
<td>IHC-P</td>
<td></td>
<td>Use a concentration of 10 - 20 µg/ml.</td>
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<tr>
<td>IHC-Fr</td>
<td></td>
<td>Use a concentration of 10 - 20 µg/ml.</td>
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<tr>
<td>ICC/IF</td>
<td></td>
<td>1/100.</td>
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</tbody>
</table>

### Target

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.

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Images
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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