**Product datasheet**

**Anti-TGF beta 1 antibody ab92486**

![Image](https://example.com)

**Overview**

<table>
<thead>
<tr>
<th>Product name</th>
<th>Anti-TGF beta 1 antibody</th>
</tr>
</thead>
<tbody>
<tr>
<td>Description</td>
<td>Rabbit polyclonal to TGF beta 1</td>
</tr>
<tr>
<td>Host species</td>
<td>Rabbit</td>
</tr>
<tr>
<td>Specificity</td>
<td>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.</td>
</tr>
<tr>
<td>Tested applications</td>
<td>Suitable for: WB, IHC-FrFl, IHC-P, IHC-Fr, ICC/IF</td>
</tr>
<tr>
<td>Species reactivity</td>
<td>Reacts with: Mouse, Rat, Human, Pig</td>
</tr>
<tr>
<td>Positive control</td>
<td>WB: Mouse 3T3 cell lysate.</td>
</tr>
</tbody>
</table>

**Properties**

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

**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.

<table>
<thead>
<tr>
<th>Application</th>
<th>Abreviews</th>
<th>Notes</th>
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</thead>
</table>

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**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.

**Cellular localization**
Secreted > extracellular space > extracellular matrix.

### Application | Notes |
---|---|
**WB** | 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. |
**IHC-FrFI** | Use at an assay dependent concentration. PubMed: 24647450 |
**IHC-P** | Use a concentration of 10 - 20 µg/ml. |
**IHC-Fr** | Use a concentration of 10 - 20 µg/ml. |
**ICC/IF** | 1/100. |

### Notes
- 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.
- Belongs to the TGF-beta family.
- Glycosylated.
- The precursor is cleaved into mature TGF-beta-1 and LAP, which remains non-covalently linked to mature TGF-beta-1 rendering it inactive.
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-mouse 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.

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