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Datasheet for ABIN411355

## TGFB1 ELISA Kit

1 Image

29 Publications

### Overview

Quantity:	96 tests
Target:	TGFB1
Binding Specificity:	AA 279-390
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	15.6-1000 pg/mL
Minimum Detection Limit:	15.6 pg/mL
Application:	ELISA

### Product Details

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of activated Mouse TGF beta 1
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (EDTA), Urine
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: CHO Immunogen sequence: A279-S390
Specificity:	Expression system for standard: CHO Immunogen sequence: A279-S390
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.

## Product Details

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Sensitivity: <2pg/mL

Material not included: Microplate reader in standard size. Automated plate washer. Adjustable pipettes and pipette tips. Multichannel pipettes are recommended in the condition of large amount of samples in the detection. Clean tubes and Eppendorf tubes. Washing buffer (neutral PBS or TBS). Preparation of 0.01M TBS: Add 1.2g Tris, 8.5g NaCl

## Target Details

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Target: TGFB1

Alternative Name: TGFB1 ([TGFB1 Products](#))

Background: Protein 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. Can promote either T- helper 17 cells (Th17) or regulatory T-cells (Treg) lineage differentiation in a concentration-dependent manner. At high concentrations, leads to FOXP3-mediated suppression of RORC and down-regulation of IL-17 expression, favoring Treg cell development. At low concentrations in concert with IL-6 and IL-21, leads to expression of the IL-17 and IL-23 receptors, favoring differentiation to Th17 cells (PubMed:18368049). .

Background: Transforming growth factor-beta1(TGF-beta1) is a multifunctional peptide that controls proliferation, differentiation, and other functions in many cell types. Many cells synthesize TGF-beta and essentially all of them have specific receptors for this peptide. TGF-beta regulates the actions of many other peptide growth factors and determines a positive or negative direction of their effects. TGFbeta1 is known for its potent and diverse biological effects, including immune regulation, and cell growth and differentiation. TGFbeta1 is also an important mediator of bone remodeling. TGFbeta1, a potent keratinocyte growth inhibitor, has been shown to be overexpressed in keratinocytes in certain inflammatory skin diseases and has been thought to counteract the effects of other growth factors at the site of inflammation. TGF-beta1, a multifunctional cytokine with fibrogenic properties, has been implicated in the pathogenesis of the vascular and target organ complications of hypertension. TGF-beta1 may also regulate blood pressure via stimulation of endothelin-1 and/or renin secretion. TGFbeta1 is secreted as a latent form, which consists of its mature form and a latency-associated peptide(beta1-LAP) in either the presence or the absence of additional latent TGF-beta1-binding protein. The standard product used in this kit is recombinant TGFbeta1 with the molecular

## Target Details

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mass of 25KDa.

Synonyms: Transforming growth factor beta-1, TGF-beta-1, Latency-associated peptide, LAP, Tgfb1,

Full Gene Name: Transforming growth factor beta-1

Cellular Localisation: Secreted, extracellular space, extracellular matrix.

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Gene ID: 21803

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UniProt: [P04202](#)

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Pathways: [EGFR Signaling Pathway](#), [Dopaminergic Neurogenesis](#), [Cellular Response to Molecule of Bacterial Origin](#), [Glycosaminoglycan Metabolic Process](#), [Regulation of Leukocyte Mediated Immunity](#), [Regulation of Muscle Cell Differentiation](#), [Positive Regulation of Immune Effector Process](#), [Cell-Cell Junction Organization](#), [Production of Molecular Mediator of Immune Response](#), [Ribonucleoside Biosynthetic Process](#), [Skeletal Muscle Fiber Development](#), [Regulation of Carbohydrate Metabolic Process](#), [Protein targeting to Nucleus](#), [Autophagy](#), [Cancer Immune Checkpoints](#)

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## Application Details

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Application Notes: Before using Kit, spin tubes and bring down all components to bottom of tube. Duplicate well assay was recommended for both standard and sample testing.

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Plate: Pre-coated

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Protocol: mouse TGF beta 1 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assay technology. A monoclonal antibody from rat specific for TGF beta 1 has been precoated onto 96-well plates. Standards(CHO, A279-S390) and test samples are added to the wells, a biotinylated detection polyclonal antibody from goat specific for TGF beta 1 is added subsequently and then followed by washing with PBS or TBS buffer. Avidin-Biotin-Peroxidase Complex was added and unbound conjugates were washed away with PBS or TBS buffer. HRP substrate TMB was used to visualize HRP enzymatic reaction. TMB was catalyzed by HRP to produce a blue color product that changed into yellow after adding acidic stop solution. The density of yellow is proportional to the mouse TGF beta1 amount of sample captured in plate.

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Assay Procedure: Aliquot 0.1 mL per well of the 1000pg/mL, 500pg/mL, 250pg/mL, 125pg/mL, 62.5pg/mL, 31.2pg/mL, 15.6pg/mL mouse TGF beta1 standard solutions into the precoated 96-well plate. Add 0.1 mL of the sample diluent buffer into the control well (Zero well). Add 0.1 mL of each properly diluted sample of mouse cell culture supernates, serum, plasma(EDTA) or urine to each empty well. See "Sample Dilution Guideline" above for details. It is recommended that

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## Application Details

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each mouse TGF beta 1 standard solution and each sample be measured in duplicate.

Assay Precision:

- Sample 1: n=16, Mean(pg/ml): 114, Standard deviation: 5.47, CV(%): 4.8
- Sample 2: n=16, Mean(pg/ml): 240, Standard deviation: 12, CV(%): 5
- Sample 3: n=16, Mean(pg/ml): 483, Standard deviation: 17.9, CV(%): 3.7,
- Sample 1: n=24, Mean(pg/ml): 125, Standard deviation: 6.75, CV(%): 5.4
- Sample 2: n=24, Mean(pg/ml): 248, Standard deviation: 15.4, CV(%): 6.2
- Sample 3: n=24, Mean(pg/ml): 515, Standard deviation: 30.4, CV(%): 5.9

Restrictions: For Research Use only

## Handling

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Handling Advice: Avoid multiple freeze-thaw cycles.

Storage: -20 °C, 4 °C

Storage Comment: Store at 4°C for 6 months, at -20°C for 12 months. Avoid multiple freeze-thaw cycles

Expiry Date: 12 months

## Publications

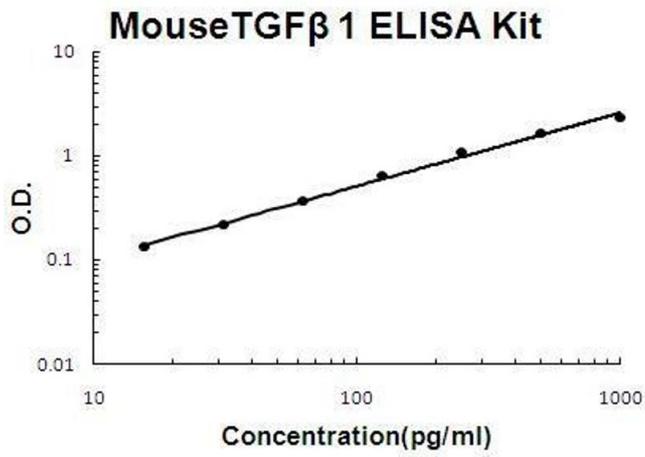
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Product cited in: Sai, Yao, Shen, Zheng, Sun, Wu, Wang, Yao: "Dynamic expression of hepatic GP73 mRNA and protein and circulating GP73 during hepatocytes malignant transformation." in: **Hepatobiliary & pancreatic diseases international : HBPD INT**, Vol. 19, Issue 5, pp. 449-454, (2020) ([PubMed](#)).

Dong, Chen, Li, Li, Wen, Lin, Ma, Wei, Chen, Ruan, Lin, Wang, Wu, Wu: "Serum Golgi protein 73 is a prognostic rather than diagnostic marker in hepatocellular carcinoma." in: **Oncology letters**, Vol. 14, Issue 5, pp. 6277-6284, (2017) ([PubMed](#)).

Kosanam, Prassas, Chrystoja, Soleas, Chan, Dimitromanolakis, Blasutig, Rückert, Gruetzmann, Pilarsky, Maekawa, Brand, Diamandis: "Laminin, gamma 2 (LAMC2): a promising new putative pancreatic cancer biomarker identified by proteomic analysis of pancreatic adenocarcinoma tissues." in: **Molecular & cellular proteomics : MCP**, Vol. 12, Issue 10, pp. 2820-32, (2013) ([PubMed](#)).

There are more publications referencing this product on: [Product page](#)



**ELISA**

**Image 1.** Mouse TGF beta 1 PicoKine ELISA Kit standard curve