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## **TGFB3 ELISA Kit**





#### Overview

Quantity:	96 tests
Target:	TGFB3
Binding Specificity:	AA 301-412
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	31.2-2000 pg/mL
Minimum Detection Limit:	31.2 pg/mL
Application:	ELISA

# **Product Details**

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of activated Mouse TGF-beta 3
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (EDTA), Urine
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: sf21 Immunogen sequence: A301-S412
Specificity:	Expression system for standard: sf21 Immunogen sequence: A301-S412
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.

#### **Product Details**

Predicted Reactivity:

Sensitivity:	<10pg/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	

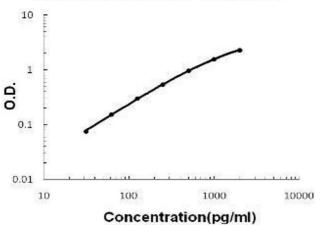
Bovine, Canine, Chicken, Horse, Monkey, Rabbit

Target Details	
Target:	TGFB3
Alternative Name:	TGFB3 (TGFB3 Products)
Background:	Protein Function: Involved in embryogenesis and cell differentiation.
	Background: Transforming growth factor beta 3(TGF-beta 3) is a type of protein, known as a
	cytokine, which is involved in cell differentiation, embryogenesis and development. It belongs to
	a large family of cytokines called the Transforming growth factor beta superfamily. TGF-beta 3
	is believed to regulate molecules involved in cellular adhesion and extracellular matrix(ECM)
	formation during the process of palate development. Without TGF-beta 3, mammals develop a
	deformity known as acleft palate. This is caused by failure of epithelial cells in both sides of the
	developing palate to fuse. TGF-beta 3 also plays an essential role in controlling the
	development of lungs in mammals, by also regulating cell adhesion and ECM formation in this
	tissue, and controls wound healing by regulating the movements of epidermal and dermal cells
	in injured skin. TGF-beta 3 activated Lef1 in the absence of beta-catenin(CTNNB1) via nuclear
	phospho-Smad2 and Smad4.
	Synonyms: Transforming growth factor beta-3,TGF-beta-3,Latency-associated
	peptide,LAP,Tgfb3,
	Full Gene Name: Transforming growth factor beta-3
	Cellular Localisation: Secreted.
Gene ID:	21809
UniProt:	P17125
Pathways:	Cell-Cell Junction Organization, Production of Molecular Mediator of Immune Response,
	Protein targeting to Nucleus

# **Application Details**

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.
Comment:	Tissue Specificity: Expressed in both testis and ovary.
Plate:	Pre-coated
Protocol:	mouse TGF-beta 3 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent
	assay technology. A monoclonal antibody from rat specific for TGF-beta 3 has been precoated
	onto 96-well plates. Standards(sf21, A301-S412) and test samples are added to the wells, a
	biotinylated detection polyclonal antibody from goat specific for TGF-beta 3 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-beta 3 amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 2000pg/mL,1000pg/mL, 500pg/mL, 250pg/mL, 125pg/mL,
	62.5pg/mL, 31.2pg/mL mouse TGF-beta 3 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 activated mouse cell culture supernates, serum, plasma(EDTA) or
	urine to each empty well. See "Sample Dilution Guideline" above for details. It is recommended
	that each mouse TGF-beta 3 standard solution and each sample be measured in duplicate.
Assay Precision:	Sample 1: n=16, Mean(pg/ml): 158, Standard deviation: 7.43, CV(%): 4.7
	<ul> <li>Sample 2: n=16, Mean(pg/ml): 476, Standard deviation: 20.5, CV(%): 4.3</li> </ul>
	• Sample 3: n=16, Mean(pg/ml): 935, Standard deviation: 50.5, CV(%): 5.4,
	<ul> <li>Sample 1: n=24, Mean(pg/ml): 182, Standard deviation: 10.56, CV(%): 5.8</li> <li>Sample 2: n=24, Mean(pg/ml): 563, Standard deviation: 32.1, CV(%): 5.7</li> </ul>
	Sample 3: n=24, Mean(pg/ml): 1138, Standard deviation: 74, CV(%): 6.5
Restrictions:	For Research Use only
Handling	
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

### Mouse TGF-beta 3 ELISA Kit



#### **ELISA**

**Image 1.** Mouse TGF-beta 3 PicoKine ELISA Kit standard curve