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# **IL-4 ELISA Kit**

1	Image

**Publications** 



### Overview

Quantity:	96 tests
Target:	IL-4 (IL4)
Binding Specificity:	AA 21-140
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	7.8-500 pg/mL
Minimum Detection Limit:	7.8 pg/mL
Application:	ELISA

### **Product Details**

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Mouse IL-4
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: E.coli Immunogen sequence: H21-S140
Specificity:	Expression system for standard: E.coli Immunogen sequence: H21-S140
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.

## **Product Details**

Sensitivity:	<1pg/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	
Target:	IL-4 (IL4)
Alternative Name:	IL4 (IL4 Products)
Background:	Protein Function: Participates in at least several B-cell activation processes as well as of other cell types. It is a costimulator of DNA-synthesis. It induces the expression of class II MHC molecules on resting B-cells. It enhances both secretion and cell surface expression of IgE and IgG1. It also regulates the expression of the low affinity Fc receptor for IgE (CD23) on both lymphocytes and monocytes.  Background: Interleukin-4(IL-4), also knowns as a B-cell stimulatory factor1(BSF1), is an immunomodulatory cytokine, which can inhibit the growth of tumour cells. The human cDNA contains a single open reading frame encoding a protein of 153 amino acids, including a putative signal peptide. IL-4 may act as an autocrine growth factor in pancreatic cancer cells and also give rise to the possibility that cancer-derived IL-4 may suppress cancer-directed immunosurveillance in vivo in addition to its growth-promoting effects, thereby facilitating pancreatic tumor growth and metastasis.1 The mouse and human genes and their protein products show structural and functional similarities. The human IL-4 gene, which occurs as a single copy in the haploid genome, is mapped on chromosome 5.  Synonyms: Interleukin-4,IL-4,B-cell IgG differentiation factor,B-cell growth factor 1,B-cell stimulatory factor 1,BSF-1,IGG1 induction factor,Lymphocyte stimulatory factor 1,II4,II-4, Full Gene Name: Interleukin-4
Gene ID:	16189
UniProt:	P07750
Pathways:	JAK-STAT Signaling, Regulation of Leukocyte Mediated Immunity, Positive Regulation of Immune Effector Process, Production of Molecular Mediator of Immune Response, Proton Transport, Activated T Cell Proliferation

# **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.
Plate:	Pre-coated
Protocol:	mouse IL-4 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assay
	technology. A monoclonal antibody from rat specific for IL-4 has been precoated onto 96-well
	plates. Standards(E.coli, H21-S140) and test samples are added to the wells, a biotinylated
	detection polyclonal antibody from goat specific for IL-4 is added subsequently and then
	followed by washing with PBS or TBS buffer. Avidin-Biotin-Peroxidase Complex was added an
	unbound conjugates were washed away with PBS or TBS buffer. HRP substrate TMB was use
	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 IL-4 amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 500pg/mL, 250pg/mL, 125pg/mL, 62.5pg/mL, 31.3pg/mL,
,	15.6pg/mL, 7.8pg/mL mouse IL-4 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 or serum to each empty well. See
	"Sample Dilution Guideline" above for details. We recommend that each mouse IL-4 standard
	solution and each sample is measured in duplicate.
	Solution and each sample is measured in duplicate.
Assay Precision:	• Sample 1: n=16, Mean(pg/ml): 44, Standard deviation: 2.38, CV(%): 5.4
	• Sample 2: n=16, Mean(pg/ml): 168, Standard deviation: 7.9, CV(%): 4.7
	<ul> <li>Sample 3: n=16, Mean(pg/ml): 283, Standard deviation: 17.55, CV(%): 6.2,</li> <li>Sample 1: n=24, Mean(pg/ml): 57, Standard deviation: 3.88, CV(%): 6.8</li> </ul>
	• Sample 2: n=24, Mean(pg/ml): 185, Standard deviation: 10.36, CV(%): 5.6
	• Sample 3: n=24, Mean(pg/ml): 326, Standard deviation: 23.8, CV(%): 7.3
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

Product cited in:

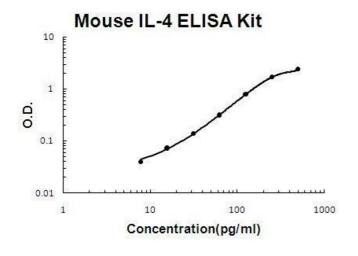
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There are more publications referencing this product on: Product page

### **Images**



### **ELISA**

Image 1. Mouse IL-4 PicoKine ELISA Kit standard curve