

# Datasheet for ABIN411271 Fibronectin 1 ELISA Kit

1 Image

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Publications



### Overview

Quantity:	96 tests
Target:	Fibronectin 1 (FN1)
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	156-10000 pg/mL
Minimum Detection Limit:	156 pg/mL
Application:	ELISA

## Product Details

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Human Fibronectin
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin), Plasma (EDTA), Plasma (citrate)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	from plasma
Specificity:	Expression system for standard: from plasma
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.
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

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## Product Details

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:	Fibronectin 1 (FN1)
Alternative Name:	FN1 (FN1 Products)
Background:	Protein Function: Fibronectins bind cell surfaces and various compounds including collagen,
	fibrin, heparin, DNA, and actin. Fibronectins are involved in cell adhesion, cell motility,
	opsonization, wound healing, and maintenance of cell shape. Involved in osteoblast
	compaction through the fibronectin fibrillogenesis cell-mediated matrix assembly process,
	essential for osteoblast mineralization. Participates in the regulation of type I collagen
	deposition by osteoblasts.
	Background: Fibronectin(FN) also known as LETS, is identified on the surfFN of fibroblasts by
	labeling with radioactive compounds or specific antibodies. Fibronectin is a 430,000-dalton
	dimeric glycoprotein that exists in 2 forms, termed cellular and plasma fibronectin. Cellular and
	plasma fibronectins are heterodimers consisting of similar but not identical polypeptides.
	These two forms of FN differ in biologic activity. Fibronectins bind cell surfFNs and various
	compounds including collagen, fibrin, heparin, DNA, and actin. Because fibronectin stimulates
	endocytosis in several systems and promotes the clearance of particulate material from the
	circulation, it could function in the clearance of C1q-coated material such as immune
	complexes or cellular debris. Fibronectins are involved in cell adhesion, cell motility,
	opsonization, would healing, and maintenance of cell shape. LETS, encoded on chromosome &
	is responsible for the LETS protein expression in humans. Because LETS has been implicated i
	tumorigenicity and cellular transformation, it is of interest that rearrangement or modifications
	in the number of chromosome 8 have been associated with certain forms of cancer. The
	standard used in this kit is isolated from human plasma with the molecular mass of 200-
	250KDa.
	Synonyms: Fibronectin,FN,Cold-insoluble globulin,CIG,Anastellin,Ugl-Y1,Ugl-Y2,Ugl-Y3,FN1,FN,
	Full Gene Name: Fibronectin
	Cellular Localisation: Secreted, extracellular space, extracellular matrix.
Gene ID:	2335
UniProt:	P02751

Pathways:

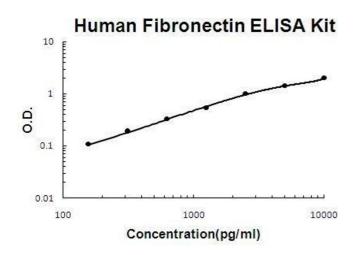
Cellular Response to Molecule of Bacterial Origin, Carbohydrate Homeostasis, Autophagy

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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.
Comment:	Sequence similarities: Contains 12 fibronectin type-I domains.
	Tissue Specificity: Plasma FN (soluble dimeric form) is secreted by hepatocytes. Cellular FN
	(dimeric or cross-linked multimeric forms), made by fibroblasts, epithelial and other cell types,
	is deposited as fibrils in the extracellular matrix. Ugl-Y1, Ugl-Y2 and Ugl-Y3 are found in urine
Plate:	Pre-coated
Protocol:	human Fibronectin ELISA Kit was based on standard sandwich enzyme-linked immune-sorben
	assay technology. A polyclonal antibody from rabbit specific for Fibronectin has been
	precoated onto 96-well plates. Standards(from plasma) and test samples are added to the
	wells, a biotinylated detection polyclonal antibody from goat specific for Fibronectin 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. HRF
	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 human Fibronectin amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 10000pg/mL, 5000pg/mL, 2500pg/mL, 1250pg/mL, 625pg/mL,
	312pg/mL, 156pg/mL human Fibronectin 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 human cell culture supernates, serum or plasma(heparin, EDTA,
	citrate) to each empty well. See "Sample Dilution Guideline" above for details. It is
	recommended that each human Fibronectin standard solution and each sample be measured
	in duplicate.
Assay Precision:	• Sample 1: n=16, Mean(ng/ml): 1.23, Standard deviation: 0.063, CV(%): 5.1
	<ul> <li>Sample 2: n=16, Mean(ng/ml): 2.35, Standard deviation: 0.099, CV(%): 4.2</li> <li>Sample 3: n=16, Mean(ng/ml): 5.53, Standard deviation: 0.265, CV(%): 4.8,</li> </ul>
	<ul> <li>Sample 3. n= 10, Mean(ng/ml): 3.33, Standard deviation: 0.203, CV(%): 4.8,</li> <li>Sample 1: n=24, Mean(ng/ml): 1.54, Standard deviation: 0.120, CV(%): 7.8</li> </ul>
	<ul> <li>Sample 2: n=24, Mean(ng/ml): 2.39, Standard deviation: 0.177, CV(%): 7.4</li> </ul>
	• Sample 3: n=24, Mean(ng/ml): 5.78, Standard deviation: 0.399, CV(%): 6.9
Restrictions:	For Research Use only
Handling	

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Handling	
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	
Product cited in:	Stevens, Scull, Ramanan, Fortin, Chaturvedi, Knouse, Xiao, Fung, Mirabella, Chen, McCue, Yang,
	Fleming, Chung, de Jong, Chen, Rice, Bhatia: "In situ expansion of engineered human liver tissue
	in a mouse model of chronic liver disease." in: <b>Science translational medicine</b> , Vol. 9, Issue 399, (2018) (PubMed).
	Banville, Burgess, Jaffar, Tjin, Richeldi, Cerri, Persiani, Black, Oliver: "A quantitative proteomic
	approach to identify significantly altered protein networks in the serum of patients with
	lymphangioleiomyomatosis (LAM)." in: <b>PLoS ONE</b> , Vol. 9, Issue 8, pp. e105365, (2014) (PubMed ).
	Lv, Wu, Zhou, Shao, Wang, Wang: "Alpha Lipoic Acid Modulated High Glucose-Induced Rat
	Mesangial Cell Dysfunction via mTOR/p70S6K/4E-BP1 Pathway." in: International journal of
	endocrinology, Vol. 2014, pp. 658589, (2014) (PubMed).
	Xu, Guan, Zheng, Gao, Zeng, Qin, Xue: "Exendin-4 alleviates high glucose-induced rat mesangial
	cell dysfunction through the AMPK pathway." in: Cellular physiology and biochemistry :
	international journal of experimental cellular physiology, biochemistry, and pharmacology,
	Vol. 33, Issue 2, pp. 423-32, (2014) (PubMed).
	Inoue, Sasaki, Katada, Taguchi: "Quantitative biocompatibility evaluation of nickel-free high-
	nitrogen stainless steel in vitro/in vivo." in: Journal of biomedical materials research. Part B,
	Applied biomaterials, Vol. 102, Issue 1, pp. 68-72, (2014) (PubMed).
	There are more publications referencing this product on: Product page



### ELISA

Image 1. Human Fibronectin PicoKine ELISA Kit standard curve

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