

Datasheet for ABIN1569731 Fibulin 1 ELISA Kit

Image



Overview

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Quantity:	96 tests					
Target:	Fibulin 1 (FBLN1)					
Reactivity:	Human					
Method Type:	Sandwich ELISA					
Detection Range:	0.468 ng/mL - 30 ng/mL					
Minimum Detection Limit:	0.468 ng/mL					
Application:	ELISA					
Product Details						
Purpose:	The kit is a sandwich enzyme immunoassay for the in vitro quantitative measurement of FBLN1 in serum, plasma, tissue homogenates, cell lysates, cell culture supernates and other biological fluids.					
Sample Type:	Cell Culture Supernatant, Cell Lysate, Plasma, Serum, Tissue Homogenate					
Analytical Method:	Quantitative					
Detection Method:	Colorimetric					
Specificity:	This assay has high sensitivity and excellent specificity for detection of this index.					
Cross-Reactivity (Details):	No significant cross-reactivity or interference between this index and analogues was observed. Note: Limited by current skills and knowledge, it is impossible for us to complete the cross- reactivity detection between this index and all the analogues, therefore, cross reaction may still exist.					
Sensitivity:	0.15 ng/mL					

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Product Details

Components:	 Pre-coated, ready to use 96-well strip plate Standard (freeze dried) Standard Diluent Detection Reagent A Detection Reagent B Assay Diluent A Assay Diluent B TMB Stop Solution 				
	• Wash Buffer (30X)				
	Plate sealer for 96 wells				
	Instruction manual				
Material not included:	1. Microplate reader with 450 ± 10nm filter.				
	2. Precision single or multi-channel pipettes and disposable tips.				
	3. Eppendorf Tubes for diluting samples.				
	4. Deionized or distilled water.				
	5. Absorbent paper for blotting the microtiter plate.				
	6. Container for Wash Solution.				

Target Details

Target:	Fibulin 1 (FBLN1)		
Alternative Name:	FBLN1 (FBLN1 Products)		
Gene ID:	2192		
UniProt:	P23142		

Application Details

Sample Volume:	100 µL					
Assay Time:	1 - 4.5 h					
Plate:	Pre-coated					
Protocol:	1. Prepare all reagents, samples and standards					
	2. Add 100 μ L standard or sample to each well. Incubate 2 hours at 37°C					
	3. Aspirate and add 100 μ L prepared Detection Reagent A. Incubate 1 hour at 37°C					
	4. Aspirate and wash 3 times					
	5. Add 100µL prepared Detection Reagent B. Incubate 1 hour at 37°C					
	6. Aspirate and wash 5 times					

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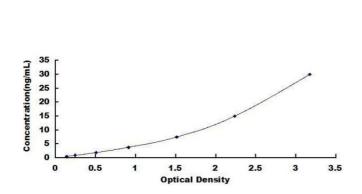
Application Details						
	7. Add 90µL Substrate Solution. Incubate 15-25 minutes at 37°C					
	8. Add 50µL Stop Solution. Read at 450nm immediately.					
Assay Procedure:	The microtiter plate provided in this kit has been pre-coated with an antibody specific to the					
	index. Standards or samples are then added to the appropriate microtiter plate wells with a					
	biotin-conjugated antibody preparation specific to the index. Next, Avidin conjugated to					
	Horseradish Peroxidase (HRP) is added to each microplate well and incubated. After TMB					
	substrate solution is added, only those wells that contain the index, biotin-conjugated antibody					
	and enzyme-conjugated Avidin will exhibit a change in color. The enzyme-substrate reaction is					
	terminated by the addition of sulphuric acid solution and the color change is measured					
	spectrophotometrically at a wavelength of 450nm \pm 10nm. The concentration of the index in					
	the samples is then determined by comparing the O.D. of the samples to the standard curve.					
Assay Precision:	• Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level the index were tested 20 times on one plate, respectively.					
	 Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level the index were tested on 3 different plates, 8 replicates in each plate. 					
	 CV(%) = SD/meanX100 					
	Intra-assay: CV<10%					
	Inter-assay: CV<12%					
Restrictions:	For Research Use only					
Handling						
Precaution of Use:	The Stop Solution suggested for use with this kit is an acid solution. Wear eye, hand, face, and clothing protection when using this material.					
Handling Advice:	The stability of ELISA kit is determined by the loss rate of activity. The loss rate of this kit is less					
	than 5 % within the expiration date under appropriate storage conditions. Note: To minimize					
	unnecessary influences on the performance, operation procedures and lab conditions,					
	unnecessary influences on the performance, operation procedures and lab conditions, especially room temperature, air humidity and incubator temperatures should be strictly					
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Storage:	especially room temperature, air humidity and incubator temperatures should be strictly regulated. It is also strongly suggested that the whole assay is performed by the same					
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-	 especially room temperature, air humidity and incubator temperatures should be strictly regulated. It is also strongly suggested that the whole assay is performed by the same experimenter from the beginning to the end. 4 °C,-20 °C The Assay Plate, Standard, Detection Reagent A and Detection Reagent B should be stored at - 					

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Expiry Date:

12 months

Images



ELISA			
Image 1.			

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