

Datasheet for ABIN1571706 **PAFAH2 ELISA Kit**

Image



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Overview

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Quantity:	96 tests
Target:	PAFAH2
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	0.312 ng/mL - 20 ng/mL
Minimum Detection Limit:	0.312 ng/mL
Application:	ELISA

Product Details

Purpose:	The kit is a sandwich enzyme immunoassay for the in vitro quantitative measurement of PAFAH2 in human plasma.
Sample Type:	Plasma
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:	0.105 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 A
	• 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:	PAFAH2
Alternative Name:	Platelet Activating Factor Acetylhydrolase 2 (PAFAH2) (PAFAH2 Products)
Background:	Alternative name: SD-PLA2, Serine-dependent phospholipase A2, Platelet-activating factor acetylhydrolase 2, cytoplasmic
Gene ID:	5051
UniProt:	Q99487

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

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	5. Add 100µL prepared Detection Reagent B. Incubate 1 hour at 37°C
	6. Aspirate and wash 5 times
	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&lt10%Inter-assay: CV&lt12%
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,
	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.
Storage:	4 °C,-20 °C
Storage Comment:	The Assay Plate, Standard, Detection Reagent A and Detection Reagent B should be stored at -
	20°C upon being received. After receiving the kit , Substrate should be always stored at

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Expiry Date:

12 months

Images



