

Datasheet for ABIN1029412

ENO1 ELISA Kit

Image

Sensitivity:



Overview	
Quantity:	96 tests
Target:	ENO1
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 NNE
	in human serum, plasma and other biological fluids.
Sample Type:	Plasma, Serum
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.

exist.

0.126 ng/mL

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

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 \pm 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:	ENO1
Alternative Name:	Enolase, Non Neuronal (NNE) (ENO1 Products)
Background:	Alternative name: ENO1, ENO1L1, MPB1, PPH, Alpha Enolase, Enolase 1, Phosphopyruvate hydratase, Plasminogen-binding protein, 2-phospho-D-glycerate hydro-lyase, C-myc promoter-binding
Gene ID:	2023
UniProt:	P06733

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 100µL standard or sample to each well. Incubate 2 hours at 37°C
	3. Aspirate and add 100µL prepared Detection Reagent A. Incubate 1 hour at 37°C

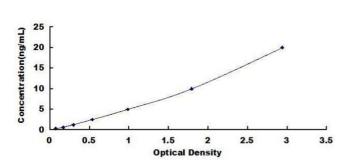
 4. Aspirate and wash 3 times 5. Add 100μL prepared Detection Reagent B. Incubate 1 hour at 37°C 6. Aspirate and wash 5 times 7. Add 90μL Substrate Solution. Incubate 15-25 minutes at 37°C 8. Add 50μL Stop Solution. Read at 450nm immediately. The microtiter plate provided in this kit has been pre-coated with an antibody specific to the
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. The microtiter plate provided in this kit has been pre-coated with an antibody specific to the
7. Add 90µL Substrate Solution. Incubate 15-25 minutes at 37°C 8. Add 50µL Stop Solution. Read at 450nm immediately. The microtiter plate provided in this kit has been pre-coated with an antibody specific to the
8. Add 50µL Stop Solution. Read at 450nm immediately. The microtiter plate provided in this kit has been pre-coated with an antibody specific to the
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 ± 10nm. The concentration of the index in
the samples is then determined by comparing the O.D. of the samples to the standard curve.
 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%
For Research Use only
The Stop Solution suggested for use with this kit is an acid solution. Wear eye, hand, face, and clothing protection when using this material.
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.
4 °C,-20 °C
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 4°C.Other reagents are kept according to the labels on vials. But for long term storage, please keep the whole kit at -20°C. The unused strips should be kept in a sealed bag with the desiccant provided to minimize exposure to damp air. The test kit may be used throughout the expiration date of the kit (six months from the date of manufacture). Opened test kits will remain stable until the expiring date shown, provided it is stored as prescribed above.

Expiry Date:

12 months

Images



ELISA

Image 1.