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## Datasheet for ABIN4947739 IL1F10 ELISA Kit



Overview

Quantity:	96 tests
Target:	IL1F10
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Application:	ELISA

## Product Details

Purpose:	Mini Samples ELISA Kit for Interleukin 1 Theta (IL1q)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of Mini Samples Interleukin 1 Theta (IL1q). No significant cross-reactivity or interference between Mini Samples Interleukin 1 Theta (IL1q) and analogues was observed.
Components:	<ul> <li>Pre-coated, ready to use 96-well strip plate</li> <li>Plate sealer for 96 wells</li> <li>Standard Diluent</li> <li>Assay Diluent A</li> <li>Assay Diluent B</li> <li>Stop Solution</li> <li>Standard</li> <li>Detection Reagent A</li> <li>Detection Reagent B</li> </ul>

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	<ul> <li>TMB Substrate</li> <li>Wash Buffer (30 x concentrate)</li> <li>Instruction manual</li> </ul>
Material not included:	Microplate reader with 450 nm filter.
	<ul> <li>Precision single or multi-channel pipettes and disposable tips.</li> </ul>
	Eppendorf Tubes for diluting samples.
	Deionized or distilled water.
	Absorbent paper for blotting the microtiter plate.
	Container for Wash Solution

## Target Details

Target:	IL1F10
Alternative Name:	Interleukin 1 Theta (IL1F10 Products)
Application Details	
Application Notes:	<ul> <li>Limited by the current condition and scientific technology, we cannot completely conduct the comprehensive identification and analysis on the raw material provided by suppliers. So there might be some qualitative and technical risks to use the kit.</li> <li>The final experimental results will be closely related to validity of the products, operation skills of the end users and the experimental environments. Please make sure that sufficient samples are available.</li> <li>Kits from different batches may be a little different in detection range, sensitivity and color developing time.</li> <li>Do not mix or substitute reagents from one kit lot to another. Use only the reagents supplied by manufacturer.</li> <li>Protect all reagents from strong light during storage and incubation. All the bottle caps of reagents should be covered tightly to prevent the evaporation and contamination of microorganism.</li> <li>There may be some foggy substance in the wells when the plate is opened at the first time. It will not have any effect on the final assay results. Do not remove microtiter plate from the storage bag until needed.</li> <li>Wrong operations during the reagents preparation and loading, as well as incorrect parameter setting for the plate reader may lead to incorrect results. A microplate plate reade with a bandwidth of 10nm or less and an optical density range of 0-3 O.D. or greater at 450 ± 10nm wavelength is acceptable for use in absorbance measurement. Please read the instruction carefully and adjust the instrument prior to the experiment.</li> <li>Even the same operator might get different results in two separate experiments. In order to get better reproducible results, the operation of every step in the assay should be controlled. Furthermore, a preliminary experiment before assay for each batch is recommended.</li> </ul>

Plate:       Pre-coated         Protocol:       The test principle applied in this kit is Sandwich enzyme immunoassay. T         provided in this kit has been pre-coated with an antibody specific to Mini 1         Theta (IL1q). Standards or samples are then added to the appropriate mic         a biotin-conjugated antibody specific to Mini Samples Interleukin 1 Theta         conjugated to Horseradish Peroxidase (HRP) is added to each microplate         After TMB substrate solution is added, only those wells that contain Mini         Theta (IL1q), biotin-conjugated antibody and enzyme-conjugated Avidin w         color. The enzyme-substrate reaction is terminated by the addition of sulf         and the color change is measured spectrophotometrically at a wavelengt         The concentration of Mini Samples Interleukin 1 Theta (IL1q) in the samp         by comparing the O.D. of the samples to the standard curve.	
The standard might be recombinant protein or natural protein, that will de kit. Moreover, the expression system is E.coli or yeast or mammal cell. Th 300 in the standard as preservative.Information on reagents: The stop solution used in the kit is sulfuric acid with concentration of 1 m solution is TBS. The standard diluent contains 0.02 % sodium azide, assa diluent B contain 0.01% sodium azide. Some kits can contain is BSA in the Information on antibodies: The provided antibodies and their host vary in different kits.Sample Volume:25 µLAssay Time:3 hPlate:Pre-coatedProtocol:The test principle applied in this kit is Sandwich enzyme immunoassay. T provided in this kit has been pre-coated with an antibody specific to Mini 3 Theta (IL1q). Standards or samples are then added to the appropriate mic a biotin-conjugated antibody specific to Mini Samples Interleukin 1 Theta conjugated to Horseradish Peroxidase (HRP) is added to each microplate After TMB substrate solution is added, only those wells that contain Mini Theta (IL1q), biotin-conjugated antibody and enzyme-conjugated Avidin w color. The enzyme-substrate reaction is terminated by the addition of suli and the color change is measured spectrophotometrically at a wavelengt The concentration of Mini Samples Interleukin 1 Theta (IL1q) in the samples the concentration of Mini Samples to the standard curve.	ome unexpected transportation conditions or e among kits from different batches might arise ne item might produce different results, since
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and the color change is measured spectrophotometrically at a wavelengt The concentration of Mini Samples Interleukin 1 Theta (IL1q) in the samp by comparing the O.D. of the samples to the standard curve.	zyme-conjugated Avidin will exhibit a change in
The concentration of Mini Samples Interleukin 1 Theta (IL1q) in the samp by comparing the O.D. of the samples to the standard curve.	ated by the addition of sulphuric acid solution
by comparing the O.D. of the samples to the standard curve.	ometrically at a wavelength of 450nm ± 10nm.
	I Theta (IL1q) in the samples is then determined
	andard curve.
Sample Preparation:• We are only responsible for the kit itself, but not for the samples consuThe user should calculate the possible amount of the samples used in	not for the samples consumed during the assay. nt of the samples used in the whole test. Please

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	reserve sufficient samples in advance.
	<ul> <li>Please predict the concentration before assaying. If values for these are not within the range of the standard curve, users must determine the optimal sample dilutions for their particular experiments. Sample should be diluted by 0.01mol/L PBS(PH=7.0-7.2).</li> <li>If the samples are not indicated in the manual, a preliminary experiment to determine the validity of the kit is necessary.</li> <li>Tissue or cell extraction samples prepared by chemical lysis buffer may cause unexpected ELISA results due to the impacts from certain chemicals.</li> <li>Due to the possibility of mismatching between antigen from other origin and antibody used ir our kits (e.g.antibody targets conformational epitope rather than linear epitope), some native or recombinant proteins from other manufacturers may not be recognized by our products.</li> <li>Influenced by the factors including cell viability, cell number or sampling time, samples from cell culture supernatant may not be detected by the kit.</li> <li>Fresh samples without long time storage is recommended for the test. Otherwise, protein degradation and denaturalization may occur in those samples and finally lead to wrong results.</li> </ul>
Assay Procedure:	<ol> <li>Prepare all reagents, samples and standards,</li> <li>Add 100µL standard or sample to each well. Incubate 2 hours at 37 °C,</li> <li>Aspirate and add 100µL prepared Detection Reagent A. Incubate 1 hour at 37 °C,</li> <li>Aspirate and wash 3 times,</li> <li>Add 100µL prepared Detection Reagent B. Incubate 30 minutes at 37 °C,</li> <li>Aspirate and wash 5 times,</li> <li>Add 90µL Substrate Solution. Incubate 10-20 minutes at 37 °C,</li> <li>Add 50µL Stop Solution. Read at 450nm immediately.</li> </ol>
Assay Precision:	Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level Min Samples Interleukin 1 Theta (IL1q) were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level Mini Samples Interleukin 1 Theta (IL1q) 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.

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Handling Advice:	The stability of kit is determined by the loss rate of activity. The loss rate of this kit is less than
Handling Advice:	
	5 % within the expiration date under appropriate storage condition.
	To minimize extra influence on the performance, operation procedures and lab conditions,
	especially room temperature, air humidity, incubator temperature should be strictly controlled. It
	is also strongly suggested that the whole assay is performed by the same operator from the
	beginning to the end.
Storage:	4 °C
Storage Comment:	• For unopened kit: All the reagents should be kept according to the labels on vials. The
	Standard, Detection Reagent A, Detection Reagent B and the 96-well strip plate should be
	stored at -20°C upon receipt while the others should be at 4°C.
	<ul> <li>For opened kit: When the kit is opened, the remaining reagents still need to be stored</li> </ul>
	according to the above storage condition. Besides, please return the unused wells to the foil
	pouch containing the desiccant pack, and reseal along entire edge of zip-seal.
	Note: It is highly recommended to use the remaining reagents within 1 month provided this is
	within the expiration date of the kit.
	• For ELISA kit, 1 day storage at 37°C can be considered as 2 months at 4°C, which means 3
	days at 37°C equaling 6 months at 4°C.
Expiry Date:	6 months

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