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Datasheet for ABIN426442 COX1 ELISA Kit

Publication



Overview

Quantity:	96 tests
Target:	COX1
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	0.31 ng/mL - 20 ng/mL
Minimum Detection Limit:	0.31 ng/mL
Application:	ELISA

Product Details

Sample Type:	Plasma, Serum
Detection Method:	Colorimetric
Sensitivity:	0.113 ng/mL
Components:	Pre-coated, ready to use 96-well strip plate, flat buttom
	Plate sealer for 96 wells
	Reference Standard
	Standard Diluent
	Detection Reagent A
	Detection Reagent B
	Assay Diluent A
	Assay Diluent B
	Reagent Diluent (if Detection Reagent is lyophilized)
	TMB Substrate
	Stop Solution

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

Target Details

Target:	COX1
Abstract:	COX1 Products
Pathways:	Proton Transport

Application Details

Application Notes:	 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. 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. Kits from different batches may be a little different in detection range, sensitivity and color developing time. Do not mix or substitute reagents from one kit lot to another. Use only the reagents supplied
	 by manufacturer. 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.
	 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.
	 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 reader with a bandwidth of 10nm or less and an optical density range of 0-3 0.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.
	• 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.

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	 Furthermore, a preliminary experiment before assay for each batch is recommended. Each kit has been strictly passed Q.C test. However, results from end users might be inconsistent with our in-house data due to some unexpected transportation conditions or different lab equipments. Intra-assay variance among kits from different batches might arise from above factors, too. Kits from different manufacturers for the same item might produce different results, since we have not compared our products with other manufacturers.
Comment:	Information on standard material:
	The standard might be recombinant protein or natural protein, that will depend on the specific
	kit. Moreover, the expression system is E.coli or yeast or mammal cell. There is 0.05% proclin
	300 in the standard as preservative.
	Information on reagents:
	The stop solution used in the kit is sulfuric acid with concentration of 1 mol/L. And the wash
	solution is TBS. The standard diluent contains 0.02 % sodium azide, assay diluent A and assay
	diluent B contain 0.01% sodium azide. Some kits can contain is BSA in them.
	Information on antibodies:
	The provided antibodies and their host vary in different kits.
Sample Volume:	100 µL
Plate:	Pre-coated
Reagent Preparation:	 Bring all kit components and samples to room temperature (18-25 °C) before use. If the kit will not be used up in one time, please only take out strips and reagents for present experiment, and leave the remaining strips and reagents in required condition. Standard - Reconstitute the Standard with 1.0 mL of Standard Diluent, keep for 10 minutes al room temperature, shake gently (not to foam). The concentration of the standard in the stock solution is 20 ng/mL. Prepare 7 tubes containing 0.5 mL Standard Diluent and produce a double dilution series. Mix each tube thoroughly before the next transfer. Set up 7 points of diluted standard such as 20 ng/mL, 10 ng/mL, 5 ng/mL, 2.5 ng/mL, 1.25 ng/mL, 0.625 ng/mL, 0.312 ng/mL, and the last microcentrifuge tube with Standard Diluent is the blank as 0 ng/mL. Detection Reagent A and Detection Reagent B - If lyophilized reconstitute the Detection Reagent A with 150µL of Reagent Diluent, keep for 10 minutes at room temperature, shake gently (not to foam). Briefly spin or centrifuge the stock Detection A and Detection B before use. Dilute them to the working concentration 100-fold with Assay Diluent A and B, respectively. Wash Solution - Dilute 20 mL of Wash Solution concentrate (30x) with 580 mL of deionized or distilled water to prepare 600 mL of Wash Solution (1x).

	5. TMB substrate - Aspirate the needed dosage of the solution with sterilized tips and do not dump the residual solution into the vial again.
	Note:
	 Making serial dilution in the wells directly is not permitted. Prepare standards within 15 minutes before assay. Please do not dissolve the reagents at 37 °C directly.
	 3. Please carefully reconstitute Standards or working Detection Reagent A and B according to the instruction, and avoid foaming and mix gently until the crystals are completely dissolved. To minimize imprecision caused by pipetting, use small volumes and ensure that pipettors are calibrated. It is recommended to suck more than 10µL for one pipetting. 4. The reconstituted Standards, Detection Reagent A and Detection Reagent B can be used only once.
	 5. If crystals have formed in the Wash Solution concentrate (30x), warm to room temperature and mix gently until the crystals are completely dissolved. 6. Contaminated water or container for reagent preparation will influence the detection result.
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Sample Collection:	Serum: Allow samples to clot for two hours at room temperature or overnight at 4°C before
	centrifugation for 20 minutes at approximately 1000 × g. Assay immediately or store samples
	in aliquot at -20°C or -80°C. Avoid repeated freeze/thaw cycles.
	Plasma: Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge samples for 15
	minutes at 1000 $ imes$ g within 30 minutes of collection. Remove plasma and assay immediately or
	store samples in aliquot at -20°C or -80°C. Avoid repeated freeze/thaw cycles.
	Biological Fluids: Centrifuge samples for 20 minutes at 1000 × g. Remove particulates and
	assay immediately or store samples in aliquot at -20 °C or -80 °C for later use. Avoid repeated freeze/thaw cycles.
Calculation of Results:	Average the duplicate readings for each standard, control, and samples and subtract the
	average zero standard optical density. Construct a standard curve by plotting the mean O.D.
	and concentration for each standard and draw a best fit curve through the points on the graph
	or create a standard curve on log-log graph paper with the target concentration on the y-axis
	and absorbance on the x-axis. Draw the best fit straight line through the standard points and it
	can be determined by regression analysis.Using some plot software, for instance, curve expert
	1.30, is also recommended. If samples have been diluted, the concentration read from the
	standard curve must be multiplied by the dilution factor.
Assay Precision:	Intra-assay Precision (precision within an assay): Three samples with low, medium and high
	levels of the target antigen were tested twenty times on one plate, respectively.

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	Inter-assay Precision (precision between assays): Three samples with low, medium and high
	levels of the target antigen were tested on three different plates, eight replicates in each plate.
	CV (%) = SD/mean X 100
	 Intra-assay: CV less than 10 % Inter-assay: CV less than 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:	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/-20 °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.
	For opened kit: When the kit is opened, the remaining reagents still need to be stored
	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
Publications	
Product cited in:	Allocca, Corrigan, Mazumder, Fake, Samson: "Inflammation, necrosis, and the kinase RIP3 are
	key mediators of AAG-dependent alkylation-induced retinal degeneration." in: Science signaling,
	Vol. 12, Issue 568, (2020) (PubMed).