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Datasheet for ABIN6720507 CYP2E1 ELISA Kit

5 Images



#### Overview

Quantity:	96 tests		
Target:	CYP2E1		
Reactivity:	Mouse		
Method Type:	Sandwich ELISA		
Detection Range:	3.12 ng/mL - 200 ng/mL		
Minimum Detection Limit:	3.12 ng/mL		
Application:	ELISA		
Product Details			
Purpose:	The kit is a sandwich enzyme immunoassay for in vitro quantitative measurement of CYP2E1 in		
	mouse tissue homogenates, cell lysates, cell culture supernates.		
	We offer validation data (WB) for the kit components. So you can be sure to order a reliable		
	ELISA kit product composed of high quality reagents.		
Sample Type:	Cell Culture Supernatant, Cell Lysate, Tissue Homogenate		
Analytical Method:	Quantitative		
Detection Method:	Colorimetric		
Specificity:	This assay has high sensitivity and excellent specificity for detection of Cytochrome P450 2E1 (CYP2E1)		
Cross-Reactivity (Details):	No significant cross-reactivity or interference between Cytochrome P450 2E1 (CYP2E1) and analogues was observed.		

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#### Product Details

Sensitivity:	1.31 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		
	• Wash Buffer (30 x concentrate)		
	Instruction manual		

## Target Details

Target:	CYP2E1		
Alternative Name:	Cytochrome P450 2E1 (CYP2E1) (CYP2E1 Products)		
UniProt:	Q05421		

#### Application Details

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			
Assay Time:	3 h			

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## Application Details

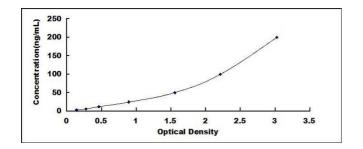
Plate:	Pre-coated
Protocol:	1. Prepare all reagents, samples and standards,
	2. Add 100 $\mu$ L standard or sample to each well. Incubate 1 hours at 37 °C,
	3. Aspirate and add 100 $\mu$ L prepared Detection Reagent A. Incubate 1 hour at 37 °C,
	4. Aspirate and wash 3 times,
	5. Add 100µL prepared Detection Reagent B. Incubate 30 minutes at 37 °C,
	6. Aspirate and wash 5 times,
	7. Add 90µL Substrate Solution. Incubate 10-20 minutes at 37 °C,
	8. Add 50µL Stop Solution. Read at 450nm immediately.
Reagent Preparation:	1. 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.
	2. Standard - Reconstitute the Standard with 1.0mL of Standard Diluent, kept for 10 minutes at
	room temperature, shake gently (not to foam). The concentration of the standard in the stock
	solution is 200ng/mL. Prepare 7 tubes containing 0.5mL 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 200ng/mL, 100ng/mL, 50ng/mL, 25ng/mL, 12.5ng/mL, 6.25ng/mL,
	3.12ng/mL, and the last tubes with Standard Diluent is the blank as 0ng/mL.
	3. 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.
	4. 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:
	1. Making serial dilution in the wells directly is not permitted.
	<ol> <li>Prepare standards within 15 minutes before assay. Please do not dissolve the reagents at 37 °C directly.</li> </ol>
	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 $10\mu$ L 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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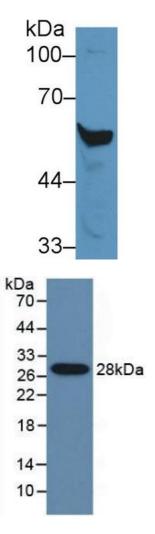
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Sample Preparation:	<ul> <li>It is recommended to use fresh samples without long storage, otherwise protein degradation and denaturationmay occur in these samples, leading to false results. Samples should therefore be stored for a short periodat 2 - 8 °C or aliquoted at -20 °C (≤1 month) or -80 °C (≤ 3 months). Repeated freeze-thawcycles should be avoided. Prior to assay, the frozen samples should be slowly thawed and centrifuged toremove precipitates.</li> <li>If the sample type is not specified in the instructions, a preliminary test is necessary to determinecompatibility with the kit.</li> <li>If a lysis buffer is used to prepare tissue homogenates or cell culture supernatant, there is a possibility of causing a deviation due to the introduced chemical substance. The recommended dilution factor is for reference only.</li> <li>Please estimate the concentration of the samples before performing the test. If the values are not in therange of the standard curve, the optimal sample dilution for the particular experiment has to be determined. Samples should then be diluted with PBS (pH =7.0-7.2).</li> </ul>
Assay Precision:	Intra-assay Precision (Precision within an assay): 3 samples with low, middle and high level of target were tested 20 times on one plate, respectively. Inter-assay Precision (Precision between assays): 3 samples with low, middle and high level of target 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 eve, hand, face, and

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.	
Storage:	4 °C/-20 °C	
Storage Comment:	<ol> <li>For unopened kit: All reagents should be stored according to the labels on the vials. The Standard, Detection Reagent A, Detection Reagent B, and 96-well Strip Plate should be stored at -20 °C upon receipt, while the other reagents should be stored at 4 °C.</li> <li>For opened kits: the remaining reagents must be stored according to the above storage conditions. In addition, please return the unused wells to the foil pouch containing the desiccant and seal the foil pouch with the zipper.</li> </ol>	
Expiry Date:	6 months	

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#### ELISA

Image 1. Typical standard curve

#### Western Blotting

**Image 2.** Rabbit Capture antibody from the kit in WB with Positive Control: Mouse liver lysate.

Western Blotting			
Image 3.	WB of	Protein	Standard: different control
antibodies	against	Highly	purified E. coli-expressed
recombinant mouse CYP2E1.			

Please check the product details page for more images. Overall 5 images are available for ABIN6720507.

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