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Datasheet for ABIN1029145 LDL ELISA Kit

Image



#### Overview

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Quantity:	96 tests
Target:	LDL
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	15.625 ng/mL - 1000 ng/mL
Minimum Detection Limit:	15.625 ng/mL
Application:	ELISA

### Product Details

Purpose:	The kit is a sandwich enzyme immunoassay for the in vitro quantitative measurement of LDL in mouse 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. 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:	6.9 ng/mL

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## Product Details

Components:	<ul> <li>Pre-coated, ready to use 96-well strip plate</li> <li>Standard (freeze dried)</li> <li>Standard Diluent</li> <li>Detection Reagent A</li> <li>Detection Reagent B</li> <li>Assay Diluent A</li> <li>Assay Diluent B</li> <li>TMB</li> <li>Stop Solution</li> <li>Wash Buffer (30X)</li> <li>Plate sealer for 96 wells</li> <li>Instruction manual</li> </ul>
Material not included:	<ol> <li>Microplate reader with 450 ± 10nm filter.</li> <li>Precision single or multi-channel pipettes and disposable tips.</li> <li>Eppendorf Tubes for diluting samples.</li> <li>Deionized or distilled water.</li> <li>Absorbent paper for blotting the microtiter plate.</li> <li>Container for Wash Solution.</li> </ol>

# Target Details

Target:	LDL
Abstract:	LDL Products

# 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 $\mu L$ standard or sample to each well. Incubate 2 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 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.

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# Application Details

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.
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&It10%
Inter-assay: CV&It12%
For Research Use only
The Stop Solution suggested for use with this kit is an acid solution. Wear eye, hand, face, and
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### Handling

