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# Datasheet for ABIN6974404 AMBP ELISA Kit



Overview

Quantity:	96 tests
Target:	AMBP
Reactivity:	Mouse
Method Type:	Competition ELISA
Detection Range:	1.563 ng/mL - 100 ng/mL
Minimum Detection Limit:	1.563 ng/mL
Application:	ELISA

### Product Details

Purpose:	For the quantitative determination of mouse alpha-1-microglobulin/bikunin precursor (AMBP) concentrations in serum, plasma, tissue homogenates.
Sample Type:	Plasma, Serum, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of mouse AMBP. No significant cross-reactivity or interference between mouse AMBP and analogues was observed. Note: Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between mouse AMBP and all the analogues, therefore, cross reaction may still exist.
Sensitivity:	0.649 ng/mL

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#### Product Details

#### Components:

- Assay plate
- Standard
- HRP-conjugate (100 x concentrate)
- Sample Diluent
- HRP-conjugate Diluent
- Wash Buffer (25 x concentrate)
- TMB Substrate
- Stop Solution
- Adhesive Strip
- Stop Solution
- Adhesive Strip

#### Target Details

Target:	AMBP
Alternative Name:	alpha-1-microglobulin/bikunin precursor (AMBP) (AMBP Products)
Background:	Abbreviation: AMBP Alias: A1M, EDC1, HCP, HI30, IATIL, ITI, ITIL, ITILC, UTI, alpha-1- microglobulin/bikunin bikunin complex-forming glycoprotein heterogeneous in charge growth- inhibiting protein 19 inter-alpha-trypsin inhibi
UniProt:	Q07456

#### **Application Details**

Application Notes:	Optimal working dilution should be determined by the investigator.
Sample Volume:	50 µL
Assay Time:	1 - 4.5 h
Plate:	Pre-coated
Protocol:	1. Prepare reagents, samples and standards as instructed.
	2. Set a Blank well without any solution.
	3. Add 50 µL standard or sample to each well.
	4. Add 50 $\mu$ L HRP-conjugate (1x) to each well (Not to Blank well).
	5. Incubate 1 hour at 37 °C
	6. Aspirate and wash 5 times.
	7. Add 90 μL of TMB Substrate to each well. Incubate for 20 minutes at 37 °C. Protect from light.
	8. Add 50 $\mu$ L Stop Solution to each well. Read at 450 nm within 5 minutes.

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Reagent Preparation:	1. HRP-conjugate (1x) - Centrifuge the vial before opening. HRP-conjugate requires a 100-fold
	dilution. A suggested 100-fold dilution is 10 $\mu$ L of HRP-conjugate + 990 $\mu$ L of HRP-conjugate Diluent.
	<ol> <li>Wash Buffer(1x)- If crystals have formed in the concentrate, warm up to room temperature and mix gently until the crystals have completely dissolved. Dilute 20 mL of Wash Buffer Concentrate (25 x) into deionized or distilled water to prepare 500 mL of Wash Buffer (1 x).</li> <li>Standard Centrifuge the standard vial at 6000-10000rpm for 30s. Reconstitute the Standard with 1.0 mL of Sample Diluent. Do not substitute other diluents. This reconstitution produce a stock solution of 100 ng/mL. Mix the standard to ensure complete reconstitution and allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making</li> </ol>
	dilutions. Pipette 150 µL of Sample Diluent into each tube (S0-S6). Use the stock solution to produce a 2-fold dilution series (below). Mix each tube thoroughly before the next transfer. The undiluted Standard serves as the high standard (100 ng/mL). Sample Diluent serves as the zero standard (0 ng/mL).
	Note:
	<ul> <li>Kindly use graduated containers to prepare the reagent. Please don't prepare the reagent directly in the Diluent vials provided in the kit.</li> <li>Bring all reagents to room temperature (18-25 °C) before use for 30 min.</li> <li>Prepare fresh standard for each assay. Use within 4 hours and discard after use.</li> <li>Making serial dilution in the wells directly is not permitted.</li> </ul>
	<ul> <li>Please carefully reconstitute Standards according to the instruction, and avoid foaming and mix gently until the crystals have 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 µL for once pipetting.</li> </ul>
	• Distilled water is recommended to be used to make the preparation for reagents or samples Contaminated water or container for reagent preparation will influence the detection result.
Sample Preparation:	<ul> <li>It is recommended to use fresh samples without long storage, otherwise protein degradatio 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> </ul>
	<ul> <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> </ul>
	<ul> <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>

Recommend to dilute the serum or plasma samples with Sample Diluent(1:500) before test.

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	The suggested 500-fold dilution can be achieved by adding 5 $\mu L$ sample to 95 $\mu L$ of Sample
	Diluent first, then complete the 500-fold dilution by adding 10 $\mu L$ of this solution to 240 $\mu L$ of
	Sample Diluent. The recommended dilution factor is for reference only. The optimal dilution
	factor should be determined by users according to their particular experiments. 6
Assay Precision:	Intra-assay Precision (Precision within an assay): CV%<8% Three samples of known
	concentration were tested twenty times on one plate to assess.
	Inter-assay Precision (Precision between assays): CV%<10% Three samples of known
	concentration were tested in twenty assays to assess.
Restrictions:	For Research Use only
Handling	
Handling Storage:	4 °C,-20 °C
	4 °C,-20 °C Unopened kit Store at 2 - 8°C. Do not use the kit beyond the expiration date. May be stored for
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