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## Datasheet for ABIN6974256 ADA ELISA Kit

1 Image

1 Publication



#### Overview

Quantity:	96 tests
Target:	ADA
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	1.56 mIU/mL - 100 mIU/mL
Minimum Detection Limit:	1.56 mIU/mL
Application:	ELISA

### Product Details

Purpose:	For the quantitative determination of human adenosine deaminase (ADA) concentrations in serum, urine, tissue homogenates.
Sample Type:	Serum, Tissue Homogenate, Urine
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of human ADA. No significant cross-reactivity or interference between human ADA and analogues was observed. Note: Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between human ADA and all the analogues, therefore, cross reaction may still exist.
Sensitivity:	0.39 mIU/mL

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#### Product Details

#### Components:

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

## Target Details

Target:	ADA
Alternative Name:	adenosine deaminase (ADA Products)
Background:	Abbreviation: ADA Alias: adenosine aminohydrolase
UniProt:	P00813
Pathways:	Regulation of G-Protein Coupled Receptor Protein Signaling, Ribonucleoside Biosynthetic Process

#### Application Details

Application Notes:	Optimal working dilution should be determined by the investigator.
Sample Volume:	100 μL
Assay Time:	1 - 4.5 h
Plate:	Pre-coated
Protocol:	1. Prepare reagents, samples and standards as instructed.
	2. Add 100 $\mu$ L standard or sample to each well. Incubate 2 hours at 37 °C.
	3. Remove the liquid of each well, don't wash.
	4. Add 100 $\mu$ L Biotin-antibody (1x) to each well. Incubate 1 hour at 37 °C.
	5. Aspirate and wash 3 times.
	6. Add 100 $\mu$ L HRP-avidin (1x) to each well. Incubate 1 hour at 37 °C
	7. Aspirate and wash 5 times.
	8. Add 90 $\mu$ L of TMB Substrate to each well. Incubate for 15-30 minutes at 37 °C. Protect from
	light.

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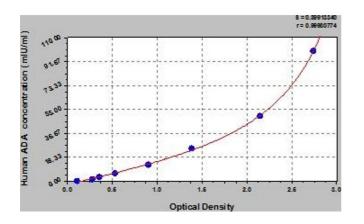
Solution to each well. Read at 450 nm within 5 minutes.
(1x) - Centrifuge the vial before opening. Biotin-antibody requires a 100-fold gested 100-fold dilution is 10 $\mu L$ of Biotin-antibody + 990 $\mu L$ of Biotin-antibody
) - Centrifuge the vial before opening. HRP-avidin requires a 100-fold dilution. A -fold dilution is 10 $\mu$ L of HRP-avidin + 990 $\mu$ L of HRP-avidin Diluent. x) - If crystals have formed in the concentrate, warm up to room temperature until the crystals have completely dissolved. Dilute 20 mL of Wash Buffer 5 x) into deionized or distilled water to prepare 500 mL of Wash Buffer (1 x). ifuge the standard vial at 6000-10000rpm for 30s. Reconstitute the Standard Sample Diluent. Do not substitute other diluents. This reconstitution produces n of 100 mIU/mL. Mix the standard to ensure complete reconstitution and lard to sit for a minimum of 15 minutes with gentle agitation prior to making te 250 $\mu$ L of Sample Diluent into each tube (S0-S6). Use the stock solution to d dilution series (below). Mix each tube thoroughly before the next transfer. Standard serves as the high standard (100 mIU/mL). Sample Diluent serves as
ard (0 mIU/mL).
luated containers to prepare the reagent. Please don't prepare the reagent biluent vials provided in the kit. Ints to room temperature (18-25 °C) before use for 30 min. Inta to room temperature (18-25 °C) before use for 30 min. Inta temperature (18-25 °C) before use for 30 min. Interview (18-25 °C) be
mall volumes and ensure that pipettors are calibrated. It is recommended to n 10 µL for once pipetting.
is recommended to be used to make the preparation for reagents. water or container for reagent preparation will influence the detection result.
ded to use fresh samples without long storage, otherwise protein degradation onmay occur in these samples, leading to false results. Samples should ored for a short periodat 2 - 8 °C or aliquoted at -20 °C (≤1 month) or -80 °C (≤ beated freeze-thawcycles should be avoided. Prior to assay, the frozen d be slowly thawed and centrifuged toremove precipitates. //pe is not specified in the instructions, a preliminary test is necessary to patibility with the kit. is used to prepare tissue homogenates or cell culture supernatant, there is a using a deviation due to the introduced chemical substance.The dilution factor is for reference only. e the concentration of the samples before performing the test. If the values

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Application Details	
	experiment has to be determined.Samples should then be diluted with PBS (pH =7.0-7.2).
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

Storage:	4 °C,-20 °C
Storage Comment:	Unopened kit Store at 2 - 8°C. Do not use the kit beyond the expiration date May be stored for
	up to 1 month at 2 - 8°C. Coated assay Try to keep it in a sealed aluminum foil bag, plate and
	avoid the damp. Standard May be stored for up to 1 month at 2 - 8° C. If Biotin-antibody don't
	make recent use, better keep it store at HRP-avidin -20°C. Biotin-antibody Diluent Opened kit
	HRP-avidin Diluent Sample May be stored for up to 1 month at 2 - 8°C. Diluent Wash Buffer
	TMB Substrate Stop Solution *Provided this is within the expiration date of the kit.
Expiry Date:	6 months
Publications	
Product cited in:	Liu, Asanoma, Takao, Tsukimori, Uchi, Furue, Kato, Wake: "Aryl hydrocarbon receptor SNP -130
	C/T associates with dioxins susceptibility through regulating its receptor activity and
	downstream effectors including interleukin 24." in: <b>Toxicology letters</b> , Vol. 232, Issue 2, pp. 384
	92, (2015) (PubMed).



**ELISA** 

Image 1. Typical Standard Curve

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