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Datasheet for ABIN577819 AP36 ELISA Kit

Image



Overview

Quantity:	96 tests
Target:	AP36
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	22.86-400 pg/mL
Minimum Detection Limit:	22.86 pg/mL
Application:	ELISA

Product Details

Sample Type:	Serum, Plasma, Tissue Homogenate, Cell Lysate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Cross-Reactivity (Details):	Limited by current skills and knowledge, it is impossible for us to complete the cross-reactivity detection between the target antigen and all analogues for other species. Therefore, cross reaction may still exist.
Sensitivity:	14.29 pg/mL
Components:	 Assay plate (12 × 8 coated Microwells) Standard (freeze dried) Biotin-antibody (100 × concentrate) HRP-avidin (100 × concentrate) Biotin-antibody Diluent

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	 HRP-avidin Diluent Sample Diluent Wash Buffer (25 × concentrate) TMB Substrate Stop Solution Adhesive Strip (for 96 wells) Instruction manual
Material not included:	 Microplate reader capable of measuring absorbance at 450nm, with the correction wavelength set at 540nm or 570nm. An incubator which can provide stable incubation conditions up to 37°C ± 0.5°C. Squirt bottle, manifold dispenser or automated microplate washer. Absorbent paper for blotting the microtiter plate. 100mL and 500mL graduated cylinders. Deionized or distilled water. Pipettes and pipette tips. Test tubes for dilution.

Target Details

Target:	AP36
Abstract:	AP36 Products
Application Details	

Application Notes:	• The supplier is only responsible for the kit itself, but not for the samples consumed during the
	assay. The user should calculate the possible amount of the samples used in the whole test
	Please reserve sufficient samples in advance.
	 Samples to be used within 5 days may be stored at 2-8°C, otherwise samples must be store
	at -20°C (\leq 1 month) or -80°C (\leq 2 months) to avoid loss of bioactivity and contamination.
	 Grossly hemolyzed samples are not suitable for use in this assay.
	If the samples are not indicated in the manual, a preliminary experiment to determine the
	validity of the kit is necessary.
	Please predict the concentration before assaying. If values for these are not within the range
	of the standard curve, users must determine the optimal sample dilutions for their particular
	experiments.
	Tissue or cell extraction samples prepared by chemical lysis buffer may cause unexpected
	ELISA results due to the impacts of certain chemicals.
	Owing to the possibility of mismatching between antigens from another resource and
	antibodies used in this supplier's kits (e.g., antibody targets conformational epitope rather
	than linear epitope), some native or recombinant proteins from other manufacturers may no
	be recognized by this supplier's products.

	Influenced by factors including cell viability, cell number and cell sampling time, samples
	from cell culture supernatant may not be recognized by the kit.
	 Fresh samples without long time storage are recommended for the test. Otherwise, protein degradation and denaturalization may occur in those samples and finally lead to wrong results.
Comment:	Detection wavelength: 450 nm
	Information on standard material:
	Depending on the antigen to be detected, standards can be either native or recombinant
	protein. The recombinant proteins are being expressed in CHO cells in most cases. Please
	inquire for more information. The formulation of auxiliary material in the standard is considered
	proprietary information, however it does not contain any poisonous substance. Proclin 300
	(1:3000) is used as preservative.
	Information on reagents:
	In most cases the stop solution provided is 1 N H2SO4. The formulation of wash solution is
	proprietary information. None of the components contain (sodium) azide, thimerosal, 2-
	mercaptoethanol (2-ME) or any other poisonous materials. For the sandwich method kits, the
	sample diluent, antibody diluent, enzyme diluent and standard all contain BSA.
	Information on antibodies:
	The antibodies provided in different kits vary in regards to clonality and host. Some antibodies
	are affinity purified, some are Protein A
Sample Volume:	100 μL
Assay Time:	1 - 4.5 h
Plate:	Pre-coated
Sample Collection:	 Serum: Use a serum separator tube (SST) and allow samples to clot for two hours at room temperature or overnight at 4 °C before centrifugation for 15 minutes at 1000 × g. Remove serum and assay immediately or aliquot and store samples at -20 °C or -80 °C. Avoid repeated freeze-thaw cycles.
	 Plasma: Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge for 15 minutes at 1000 × g at 2-8 °C within 30 minutes of collection. Assay immediately or aliquot and store samples at -20 °C or -80 °C. Avoid repeated freeze-thaw cycles.

• **Tissue Homogenates**: Rinse 100 mg tissue with 1× PBS, homogenize in 1mL of 1× PBS and store overnight at -20 °C. After two freeze-thaw cycles to break the cell membranes, centrifuge the homogenates for 5 minutes at 5000 × g, 2-8 °C. Remove and assay the

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	supernate immediately. Alternatively, aliquot and store samples at -20 °C or -80 °C. Centrifuge the sample again after thawing before the assay. Avoid repeated freeze-thaw cycles.
Calculation of Results:	Average the duplicate readings for each standard and sample and subtract the average zero
	standard optical density.
	Create a standard curve by reducing the data using computer software capable of generating a
	four parameter logistic (4-PL) curve fit. As an alternative, construct a standard curve by plotting
	the mean absorbance for each standard on the x-axis against the concentration on the y-axis
	and draw a best fit curve through the points on the graph. The data may be linearized by
	plotting the log of the target antigen concentration versus the log of the O.D. and the best fit line
	can be determined by regression analysis. This procedure will produce an adequate but less precise fit of the data.
	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 of known concentration were
	tested twenty times on one plate to assess precision.
	Inter-assay precision (precision between assays): Three samples of known concentration were
	tested in twenty assays to assess precision.
	 Intra-assay: CV% less than 8%
	 Inter-assay: CV% less than 10%
Restrictions:	For Research Use only
Handling	
Precaution of Use:	The Stop Solution provided with this kit is an acid solution. Wear eye, hand, face and clothing
	protection when using this material.
Handling Advice:	The kit should not be used beyond the expiration date on the kit label.
	 Do not mix or substitute reagents with those from other lots or sources.
	If samples generate values higher than the highest standard, dilute the samples with Sample
	Diluent and repeat the assay.
	Any variation in Sample Diluent, operator, pipetting technique, washing technique, incubation
	time/temperature and kit age can cause variation in binding.
	 This assay is designed to eliminate interference by soluble receptors, binding proteins and other factors present in biological samples. Until all factors have been tested in the
	Immunoassay, the possibility of interference cannot be excluded.
Storage:	4 °C/-20 °C

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Handling

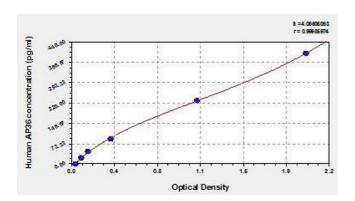
Storage Comment: For un

Expiry Date:

For unopened kit: All the reagents should be kept according to the labels on vials.

6 months

Images



ELISA

Image 1. Typical standard curve

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