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Datasheet for ABIN813939 EDIL3 ELISA Kit

1 Image

1 Publication



Overview

Quantity:	96 tests
Target:	EDIL3 (DEL1)
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	125-8000 pg/mL
Minimum Detection Limit:	125 pg/mL
Application:	ELISA
Product Details	
Purpose:	For the quantitative determination of human EGF-like repeat and discoidin I-like domain- containing protein 3 (EDIL3) concentrations in serum, plasma, tissue homogenates.
Sample Type:	Serum, Plasma, Tissue Homogenate
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	This assay has high sensitivity and excellent specificity for detection of human EDIL3.
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:	31.25 pg/mL
Components:	Assay plate (12 × 8 coated Microwells)

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- Standard (freeze dried)
- Biotin-antibody (100 × concentrate)
- HRP-avidin (100 × concentrate)
- Biotin-antibody Diluent
- HRP-avidin Diluent
- Sample Diluent
- Wash Buffer (25 × concentrate)
- TMB Substrate
- Stop Solution
- Adhesive Strip (for 96 wells)
- Instruction manual

Target Details

-	
Target:	EDIL3 (DEL1)
Alternative Name:	EGF-Like Repeats and Discoidin I-Like Domains 3 (EDIL3) (DEL1 Products)
Background:	Synonyms: DEL1, MGC26287, EGF-like repeats and discoidin I-like domains-containing protein 3 developmental endothelial locus-1
HGNC:	3173
UniProt:	043854

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 stored
	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 not
	be recognized by this supplier's products.

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 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.
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
100 μL
1 - 4.5 h
Pre-coated
This assay employs the quantitative sandwich enzyme immunoassay technique. Antibody
specific for EDIL3 has been pre-coated onto a microplate. Standards and samples are pipetted
into the wells and any EDIL3 present is bound by the immobilized antibody. After removing any
unbound substances, a biotin-conjugated antibody specific for EDIL3 is added to the wells.
After washing, avidin conjugated Horseradish Peroxidase (HRP) is added to the wells. Following
After washing, avidin conjugated Horseradish Peroxidase (HRP) is added to the wells. Following a wash to remove any unbound avidin-enzyme reagent, a substrate solution is added to the

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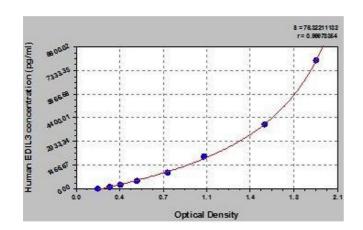
color development is stopped and the intensity of the color is measured.

Assay Precision:	contaminated water or container for reagent preparation will influence detection result.
	 Note: Kindly use graduated containers to prepare the reagent. Please don't prepare the reagent directly in the Diluent vials provided in the kit. Bring all reagents to room temperature (18-25°C) before use for 30 min. Prepare fresh standard for each assay. Use within 4 hours and discard after use. Making serial dilution in the wells directly is not permitted. Please carefully reconstitute Standards according to the instruction. 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 10µL when pipetting. It is recommended to use distilled water to prepare reagents and samples. Using
	 HRP-avidin (1×) - Centrifuge the vial before opening. HRP-avidin requires a 100-fold dilution. The suggested dilution is 10µL of HRP-avidin + 990µL of HRP-avidin Diluent. Wash Buffer (1×) - If crystals have formed in the concentrate, warm up to room temperature and mix gently until the crystals have completely dissolved. Dilute 20mL of Wash Buffer Concentrate (25×) into deionized or distilled water to prepare 500mL of Wash Buffer (1×). Standard - Centrifuge the standard vial at 6000-10000rpm for 30s. Reconstitute the Standard with 1ml of Sample Diluent. Do not substitute other diluents. This reconstitution produces a stock solution. 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 dilutions. Pipette 250µL of Sample Diluent into each tube. Use the stock solution to produce a 2-fold dilution series. Mix each tube thoroughly before the next transfer. The undiluted Standard serves as the high standard. Sample Diluent serves as the zero standard (0ng/mL).

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	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
Storage Comment:	For unopened kit: All the reagents should be kept according to the labels on vials.
Expiry Date:	6 months
Publications	
Product cited in:	Finn, Eapen, Manocha, Al Kassem, Lassegue, Ghasemzadeh, Quyyumi, Searles: "Coronary heart disease alters intercellular communication by modifying microparticle-mediated microRNA transport." in: FEBS letters , Vol. 587, Issue 21, pp. 3456-63, (2014) (PubMed).

Images



ELISA

Image 1. Typical standard curve