Datasheet for ABIN1379961

E-cadherin ELISA Kit

1 Image

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Overview

Quantity:	96 tests
Target:	E-cadherin (CDH1)
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	188-12000 pg/mL
Minimum Detection Limit:	188 pg/mL
Application:	ELISA

Product Details

Purpose:	The OmniKine? Human E-Cadherin ELISA Kit contains the components necessary for quantitative determination of natural or recombinant Human E-Cadherin concentrations within any experimental sample including cell lysates, serum and plasma. This particular immunoassay utilizes the quantitative technique of a "Sandwich" Enzyme- Linked Immunosorbent Assay (ELISA) where the target protein (antigen) is bound in a "sandwich" format by the primary capture antibodies coated to each well-bottom and the secondary detection antibodies added subsequently by the investigator. The capture antibodies coated to the bottom of each well are specific for a particular epitope on Human E- Cadherin while the user-added detection antibodies bind to epitopes on the captured target protein. Amid each step of the procedure, a series of wash steps must be performed to ensure the elimination of non-specific binding between proteins to other proteins or to the solid phase. After incubation and "sandwiching" of the target antigen, a peroxidase enzyme is conjugated to the constant heavy chain of the secondary antibody (either covalently or via Avidin/Streptavidin-Biotin interactions), allowing for a colorimetric reaction to ensue upon substrate addition. When the

	substrate TMB (3, 3', 5, 5'-Tetramethylbenzidine) is added, the reaction catalyzed by peroxidase yields a blue color that is representative of the antigen concentration. Upon sufficient color development, the reaction can be terminated through addition of Stop Solution (2 N Sulfuric Acid) where the color of the solution will turn yellow. The absorbance of each well can then be read by a spectrophotometer, allowing for generation of a standard curve and subsequent determination of protein concentration.
Brand:	OmniKine™
Sample Type:	Cell Lysate, Serum, Plasma
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	The Human E-Cadherin ELISA Kit allows for the detection and quantification of endogenous levels of natural and/or recombinant Human E-Cadherin proteins.
Cross-Reactivity (Details):	The Human E-Cadherin ELISA is capable of recognizing both recombinant and naturally produced Human E-Cadherin proteins. The antigens listed below were tested at 60 ng/mL and exhibited less than 2% cross reactivity. Murine: E-Cadherin The antigens listed below were tested at 120 ng/mL and did not exhibit significant cross reactivity or interference. Human: N-Cadherin, P-Cadherin, VE-Cadherin Murine: P-Cadherin
Characteristics:	The Human E-Cadherin ELISA Kit allows for the detection and quantification of endogenous levels of natural and/or recombinant Human E-Cadherin proteins within the range of 188-12000 pg/mL.
Components:	• Microstrips Coated w / Capture Antibody: 12 x 8-Well Microstrips • Protein Standard: Lyophilized (100 ng), Red container • Biotinylated Detection Antibody: Lyophilized, Yellow container • 400x Streptavidin-HRP: 30 µL, Blue container • Wash Buffer (10x): 50 mL, Clear containter • Assay Diluent: 50 mL, Clear container • Ready-to-Use Substrate: 12 mL, Brown container • Stop Solution: 12 mL, Clear container • Adhesive Plate Sealers: 4 Sheets • Technical Manual 1 Manual
Material not included:	The following materials and/or equipment are NOT provided in this kit but are necessary to successfully conduct the experiment: Microplate reader able to measure absorbance at 450 nm (with correction wavelength set to 540 nm or 570 nm)

Product Details

	Micropipettes with capability of measuring volumes ranging from 1 µl to 1 mL Deionized or sterile water Squirt bottle, manifold dispenser, multichannel pipette reservoir or automated microplate washer Graph paper or computer software capable of generating or displaying logarithmic functions Absorbent paper or vacuum aspirator Test tubes or microfuge tubes capable of storing ≥1 mL Bench top centrifuge (optional) Bench top vortex (optional) Orbital shaker (optional)

Target Details

Target:	E-cadherin (CDH1)
Alternative Name:	E-Cadherin (CDH1 Products)
Background:	E-Cadherins are calcium-dependent cell adhesion proteins. They preferentially interact with themselves in a homophilic manner in connecting cells, thus may contribute to the sorting of heterogeneous cell types. CDH1 is involved in mechanisms regulating cell-cell adhesions, mobility and proliferation of epithelial cells. It has a potent invasive suppressor role and is a ligand for integrin alpha-E/beta-7. Fragment E- CAD/CTF2 promotes non-amyloidogenic degradation of Abeta precursors and as a strong inhibitory effect on APP C99 and C83 production. During apoptosis or with calcium influx, cleaved by a membrane-bound metalloproteinase (ADAM10), PS1/gamma-secretase and caspase-3 to produce fragments of about 38 kDa (E-CAD/CTF1), 33 kDa (E-CAD/CTF2) and 29 kDa (E-CAD/CTF3), respectively. In mechanism, E-Cadherin is induced by calcium influx, processed by the metalloproteinase, which causes disruption of cell-cell adhesion and the subsequent release of beta- catenin into the cytoplasm. The residual membrane-tethered cleavage product is rapidly degraded via an intracellular proteolytic pathway. Cleavage by caspase-3 releases the cytoplasmic tail resulting in disintegration of the actin microfilament system. The gamma-secretase- mediated cleavage promotes disassembly of adherens junctions. N- glycosylation at Asn-637 is essential for expression, folding and trafficking. Defects in CDH1 are the cause of hereditary diffuse gastric cancer (HDGC), an autosomal dominant cancer predisposition syndrome with increased susceptibility to diffuse gastric cancer. Diffuse gastric cancer is a malignant disease characterized by poorly differentiated infiltrating lesions resulting in thickening of the stomach.

Target Details

	Such malignant tumors start in the stomach, can spread to the esophagus or the small intestine, and can extend through the stomach wall to nearby lymph nodes and organs. It also can metastasize to other parts of the body. Note: Heterozygous germ line mutations CDH1 are responsible for familial cases of diffuse gastric cancer. Somatic mutations have also been found in patients with sporadic diffuse gastric cancer and lobular breast cancer. Source: Entrez Gene, Swiss-Prot
Pathways:	WNT Signaling, Sensory Perception of Sound, Cell-Cell Junction Organization, Tube Formation

Application Details

Plate:	Pre-coated
Protocol:	This particular immunoassay utilizes the quantitative technique of a Sandwich Enzyme-Linked Immunosorbent Assay (ELISA) where the target protein (antigen) is bound in a sandwich format by the primary capture antibodies coated to each well-bottom and the secondary detection antibodies added subsequently by the investigator. The capture antibodies coated to the bottom of each well are specific for a particular epitope on the Human E-Cadherin cytokine while the user-added detection antibodies bind to of epitopes on the captured target protein. Amid each step of the procedure, a series of wash steps must be performed to ensure the elimination of non-specific binding between proteins to other proteins or to the solid phase. After incubation and sandwiching of the target antigen, a peroxidase enzyme is conjugated to the constant heavy chain of the secondary antibody (either covalently or via Avidin/Streptavidin-Biotin interactions), allowing for a colorimetric reaction to ensue upon substrate addition. When the substrate TMB (3, 3', 5, 5'-Tetramethylbenzidine) is added, the reaction catalyzed by peroxidase yields a blue color that is representative of the antigen concentration. Upon sufficient color development, the reaction can be terminated through addition of Stop Solution (2 N Sulfuric Acid) where the color of the solution will turn yellow. The absorbance of each well can then be read by a spectrophotometer, allowing for generation of a standard curve and subsequent determination of protein concentration.
Sample Preparation:	If samples are to be used within 24 hours, aliquot and store at 4 °C. If samples are to be used over a long period of time, aliquot and store between -20 °C and -80 °C, depending on the duration of storage. Note: Samples containing a visible precipitate or pellet must be clarified prior to use in the assay. Caution: Avoid repeated freeze/thaw cycles to prevent loss of biological activity of proteins in experimental samples.

- Cell Lysate and Supernatants:
  Remove large cell components via centrifugation and perform the assay. Cell lysates and supernatants require a dilution using Assay Diluent. A serial dilution may be performed to determine a suitable dilution factor for the sample. For future use of the sample, follow the sample storage guidelines stated above.
- Serum:
  Allow samples to clot in a serum separator tube (SST) for 30 minutes. After sufficient clotting, centrifuge at 1000 x g for 15 minutes and remove serum from SST in preparation for the assay. Serum samples require at least a 1:50 dilution using Assay Diluent. For future use of the sample, follow the storage guidelines above.
- Plasma:
  Use heparin, citrate or EDTA as an anticoagulant to gather plasma from original biological sample. After collection of the plasma, centrifuge for 15 minutes at 1000 x g. This step must be performed within 30 minutes of plasma collection. Plasma samples require at least a 1:50 dilution using Assay Diluent. Afterwards, perform the assay or for future use of the sample, follow the storage guidelines stated above.

Assay Procedure:

Note: If possible, all incubation steps should be performed on an orbital shaker to equilibrate solutions when added to the microplate wells. Also, all provided solutions should be at ambient temperature prior to use.

Note: Avoid adding solutions into wells at an angle, always keep pipette tip perpendicular to plate bottom.

Reconstitution of Provided Materials:

1. Reconstitute the Biotin-Conjugated Detection Antibody in 67 µL of ddH□O for a concentration of 180 µg/ml.
2. Reconstitute the Protein Standard in 100 µL of ddH□O for a concentration of 340 ng/ml.
3. Dilute the 50 mL of 10x Wash Buffer in 450 mL of ddH2O for 500 mL of 1x Wash Buffer.

Addition of Known Standard and Unknown Sample to Immunoassay:

The OmniKine™ Human CD163 ELISA Kit allows for the detection and quantification of endogenous levels of natural and/or recombinant Human CD163 proteins

Calculation of Results:

Generation of Standard Curve and Interpretation of Data

1. Average the duplicate or triplicate readings for each standard, control and sample and subtract the average zero standard optical density.

2. Generate a standard curve by using Microsoft Excel or other computer software capable of establishing a 4- Parameter Logistic (4-PL) curve fit. If using Excel or an alternative graphing tool, plot the average optical density values in absorbance units (y-axis) against the known standard concentrations in pg/ml (x-axis). Note: Only use the values in which a noticeable gradient can be established. Afterwards, generate a best fit curve or trend-line through the

Application Details

	plotted points via regression analysis.
Restrictions:	For Research Use only

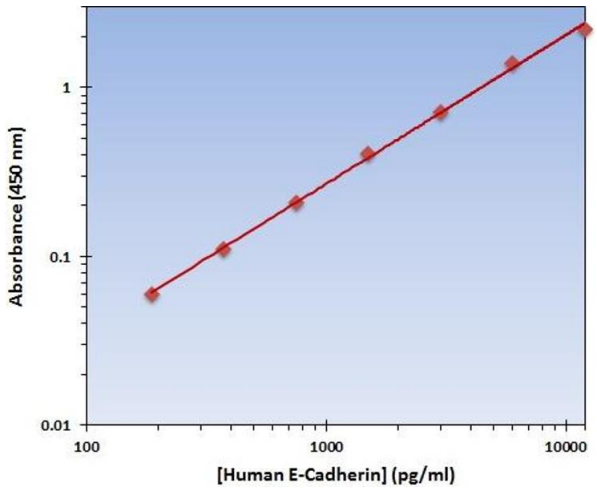
Handling

Precaution of Use:	Reagents provided in this kit may be harmful if ingested, inhaled or absorbed through the skin. Please carefully review the MSDS for each reagent before conducting the experiment. Stop Solution contains 2 N Sulfuric Acid (H2SO4) and is an extremely corrosive agent. Please wear proper eye, hand and face protection when handling this material. When the experiment is finished, be sure to rinse the plate with copious amounts of running water to dilute the Stop Solution prior to disposing the plate.
Handling Advice:	This ELISA kit is intended for research purposes only, NOT diagnostic or clinical procedures of any kind. Materials included in this kit should NOT be used past the expiration date on the kit label. Reagents or substrates included in this kit should NOT be mixed or substituted with reagents or substrates from any other kits. Variations in pipetting technique, washing technique, operator laboratory technique, kit age, incubation time or temperature may cause differences in binding affinity of the materials provided. The assay is designed to eliminate interference and background by other cellular macromolecules or factors present within any biological samples. However, the possibility of background noise cannot be fully excluded until all factors have been tested using the assay kit. Reagents provided in this kit may be harmful if ingested, inhaled or absorbed through the skin. Please carefully review the MSDS for each reagent before conducting the experiment. Stop Solution contains 2 N Sulfuric Acid (H2SO4) and is an extremely corrosive agent. Please wear proper eye, hand and face protection when handling this material. When the experiment is finished, be sure to rinse the plate with copious amounts of running water to dilute the Stop Solution prior to disposing the plate.
Storage:	4 °C
Storage Comment:	Note: If used frequently, reagents may be stored at 4 °C. • Unopened Kits: Store at 4 °C for 6 months. • Microstrips Coated w/ Capture Antibody, 400x Streptavidin-HRP Wash Buffer (10x), Assay Diluent Ready-to-Use Substrate, Stop Solution: 6 Months at 4 °C • Protein Standard, Biotinylated Detection Antibody: Lyophilized: 6 Months (if Reconstituted: 1

Handling

Month) at 4 °C

Images



ELISA

Image 1. This is an example of what a typical standard curve will look like. You must make your own standard curve. Do not use this example as your own standard curve.