

Human FOXO1A Immunoassay

Catalog Number: SEKH-0541

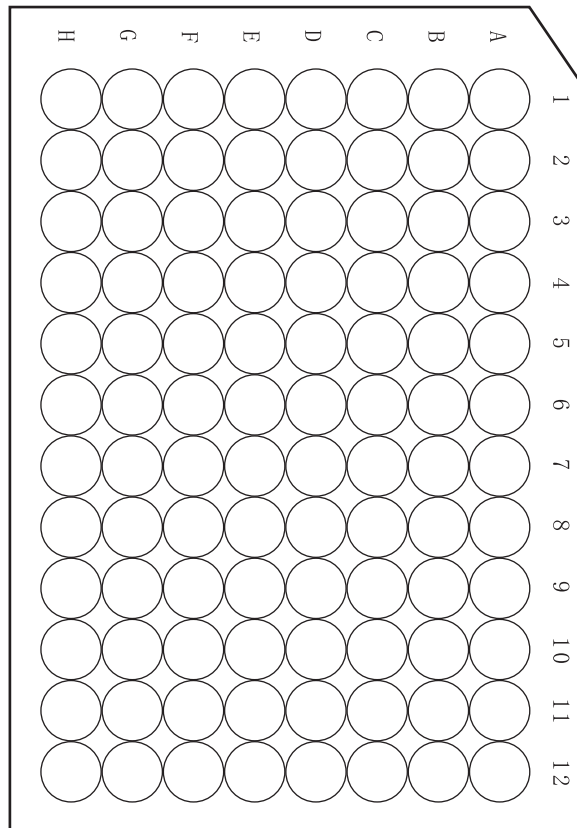
For the quantitative determination of human FOXO1A concentrations
in cell culture supernates, serum, and plasma.

For research use only. Not for use in diagnostic procedures.

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Performance Characteristics

SENSITIVITY: The minimum detectable dose was 0.8ng/mL.

RECOVERY: The coefficient of variation of both intra-assay and inter-assay were less than 10%.

RECOVERY: The recovery of FOXO1A spiked to three different levels in four samples throughout the range of the assay in various matrices was evaluated.

Recovery of FOXO1A in two matrices

Sample Type	Average % of Expected Range(%)	Range(%)
Citrate plasma	94	85-99
Cell culture supernatants	102	95-109

LINEARITY: To assess the linearity of the assay, three samples were spiked with high concentrations of FOXO1A in various matrices and diluted with the appropriate Sample Diluent to produce samples with values within the dynamic range of the assay.

Dilution ratio	Recovery(%)	Citrate plasma	Cell culture supernatants
1:2	Average% of Expected	89	102
	Range(%)	83-97	93-107
1:4	Average% of Expected	95	110
	Range(%)	90-102	104-119

BACKGROUND

FOXO1A (also known as Forkhead box protein O1 and forkhead in rhabdomyosarcoma) is an intracellular transcription factor. FOXO1A has widespread expression and functions in metabolic homeostasis and insulin signaling. Human FOXO1A has 92% protein sequence identity to mouse and rat FOXO1A.

PRINCIPLE OF THE ASSAY

This assay employs the quantitative sandwich enzyme immunoassay technique. A monoclonal antibody specific for FOXO1A has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and any FOXO1A present is captured by the coated antibody after incubation. Following extensive washing, a biotin-conjugate antibody specific for FOXO1A is added to detect the captured FOXO1A protein in sample. For signal development, horseradish peroxidase (HRP)-conjugated Streptavidin is added, followed by tetramethyl-benzidine (TMB) reagent. Following a wash to remove any unbound combination, and enzyme conjugate is added to the wells. Solution containing sulfuric acid is used to stop color development and the color intensity which is proportional to the quantity of bound protein is measurable at 450nm.

DESCRIPTION



TECHNICAL HINTS AND LIMITATIONS

1. This Solarbio ELISA should not be used beyond the expiration data on the kit label.
2. To avoid cross-contamination, use a fresh reagent reservoir and pipette tips for each step.
3. To ensure accurate results, some details, such as technique, plasticware and water sources should be emphasized.
4. A thorough and consistent wash technique is essential for proper assay performance.
5. A standard curve should be generated for each set of samples assayed.
6. It is recommended that all standards and samples be assayed in duplicate.
7. Avoid microbial contamination of reagents and buffers. Buffers containing protein should be made under aseptic conditions and be prepared fresh daily.
8. In order to ensure the accuracy of the results, the standard curve should be made every time.

PRECAUTIONS

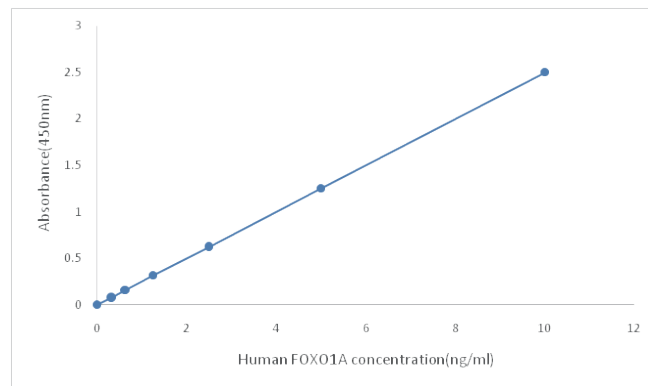
The Stop Solution suggested for use with this kit is an acid solution. Wear protective gloves, clothing, eye, and face protection. Wash hands thoroughly after handling.

DESCRIPTION

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. This standard curve is provided for demonstration only. A standard curve should be generated for each set of samples assayed.

Typical data using the FOXO1A ELISA

Standardized (ng/ml)	OD.	OD.	Average	Corrected
0	0.048	0.053	0.050	-
0.16	0.173	0.167	0.170	0.120
0.3	0.251	0.243	0.247	0.197
1	0.374	0.361	0.368	0.317
1.25	0.593	0.574	0.584	0.533
2.5	0.964	0.932	0.948	0.898
5	1.569	1.518	1.544	1.493
10	2.557	2.473	2.515	2.465



Representative standard curve for FOXO1A ELISA.

DESCRIPTION

ASSAY PROCEDURE

Prepare all reagents and standards as directed.



Add 100µl standard or samples to each well, shaking with Micro-oscillator(100r/min)incubate 120 minutes, at room temperature(25±2°C).



Aspirate and wash 4 times

Add 100µl working solution of Biotin-Conjugate anti-human FOXO1A antibody to each well, shaking with Micro-oscillator(100r/min) incubate 120 minutes,at room temperature.



Aspirate and wash 4 times

Add 100µl working solution of Streptavidin-HRP to each well, shaking with Micro-oscillator(100r/min)incubate 30 minutes,at room temperature



Aspirate and wash 5 times

Add 100µl Substrate solution to each well, incubate 5-20 minutes,at room temperature(25±2°C).Protect from light.



Add 50I Stop solution to each well. Read at 450nm within 30 minutes.

CALCULATION OF RESULTS

1. The standard curve is used to determine the amount of specimens.
2. First, average the duplicate readings for each standard, control, and sample. All O.D. values are subtracted by the mean value of blank control before result interpretation.
3. Construct 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 y-axis against the concentration on the x-axis and draw a best fit curve through the points on the graph.
4. The data may be linearized by plotting the log of the FOXO1A concentrations versus the log of the O.D. and the best fit line can be determined by

DESCRIPTION

KIT COMPONENTS& STORAGE CONDITIONS

PART	SIZE	STORAGE OF OPENED/ RECONSTITUTED MATERIAL
Microwell Plate - antibody coated 96-well Microplate (8 wells ×12 strips)	1 plate	Return unused wells to the foil pouch containing the desiccant pack. Reseal along entire edge of the zip-seal. May be stored for up to 1 month at 2 – 8 C**
Standard - lyophilized,10ng/ml upon reconstitution	2 vials	Store at 2-8°C ***for six months
Concentrated Biotin-Conjugated antibody(100X) - 120 ul/vial	1 vial	Store at 2-8°C ***for six months
Concentrated Streptavidin-HRP solution(100X) - 120 ul/vial	1 vial	Store at 2-8°C ***for six months
Standard /sample Diluent - 16 ml/vial	1 bottle	Store at 2-8°C ***for six months
Biotin-Conjugate antibody Diluent - 16 ml/vial	1 bottle	Store at 2-8°C ***for six months
Streptavidin-HRP Diluent - 16 ml/vial	1 bottle	Store at 2-8°C ***for six months
Wash Buffer Concentrate (20x) - 30 ml/vial	1 bottle	Store at 2-8°C ***for six months
Substrate Solution - 12 ml/vial	1 bottle	Store at 2-8°C ***for six months
Stop Solution - 12 ml/vial	1 bottle	Store at 2-8°C ***for six months
Plate Cover Seals	4 pieces	

**Provided this is within the expiration date of the kit.

OTHER SUPPLIES REQUIRED BUT NOT SUPPLIED

1. Microplate reader capable of measuring absorbance at 450 nm.
2. Pipettes and pipette tips.
3. Deionized or distilled water.
4. Squirrt bottle, manifold dispenser, or automated microplate washer.
5. 500 mL graduated cylinder.
6. Human FOXO1A controls (optional; available from Solarbio).

SPECIMEN COLLECTION & STORAGE

Cell Culture Supernates - Centrifuge cell culture media at 1000×g to remove debris. Assay immediately or aliquot and store samples at $\leq -20\text{ }^{\circ}\text{C}$. Avoid repeated freeze-thaw cycles.

Serum - Use a serum separator tube (SST) and allow samples to clot for 2 hours at room temperature or overnight at 2-8°C. Centrifuge at approximately for 15 minutes at 1000×g. Assay immediately or aliquot and store samples at $\leq -20\text{ }^{\circ}\text{C}$. Avoid repeated freeze-thaw cycles.

Plasma - Collect plasma using EDTA, heparin, or citrate as an anticoagulant. Centrifuge for 15 minutes at 1000×g within 30 minutes of collection. Assay immediately or aliquot and store samples at $\leq -20\text{ }^{\circ}\text{C}$. Avoid repeated freeze-thaw cycles.

Note: It is recommended to conduct a pre-test before the formal experiment to determine the dilution ratio.

REAGENTS PREPARATION

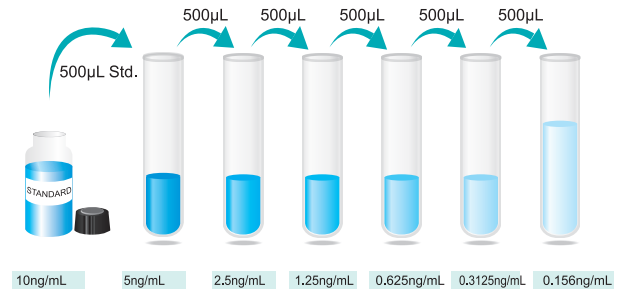
1. **Temperature returning** - Bring all kit components and specimen to room temperature (20-25°C) before use.
2. **Wash Buffer** - Dilute 30mL of Wash Buffer Concentrate with 570mL of deionized or distilled water to prepare 600mL of Wash Buffer. If crystals have formed in the concentrate Wash Buffer, warm to room temperature and mix gently until the crystals have completely dissolved.

3. **Standard\Sample** - Reconstitute the Standard with 1.0mL of deionized or distilled water. This reconstitution produces a stock solution of 10ng/mL. Allow the standard to sit for a minimum of 15 minutes with gentle agitation prior to making dilutions. Pipette 500μL of Standard/Specimen Diluent into 5ng/ml tube and the remaining tubes. Use the stock solution of 10ng/mL to produce a 2-fold dilution series (below). Mix each tube thoroughly and change pipette tips between each transfer. The 10ng/mL standard serves as the high standard. The Standard/specimen Diluent serves as the zero standard (0 pg/mL).

***If you do not run out of re-melting standard, store it at -20°C. Diluted standard shall not be reused.**

4. **Working solution of Biotin-Conjugate anti-human FOXO1A antibody:** Make a 1:100 dilution of the concentrated Biotin-Conjugate solution with the Biotin-Conjugate antibody Diluent in a clean plastic tube.
5. **Working solution of Streptavidin-HRP:** Make a 1:100 dilution of the concentrated Streptavidin-HRP solution with the Streptavidin-HRP Diluent in a clean plastic tube.

***The working solution should be used within one day after dilution.**



Preparation of FOXO1A standard dilutions