ELISA Kit for Measuring Human FABP4/A-FABP

CircuLex Human FABP4/A-FABP ELISA Kit

Cat# CY-8122

Intended Use..... 1
Storage..... 1
Introduction.....2
Principle of the Assay..... 2-3
Materials Provided.....3
Materials Required but not Provided..... 4
Precautions and Recommendations..... 5
Sample Collection and Storage.....6
Detailed Protocol..... 7-8
Calculations.....9
Measurement Range..... 9
Troubleshooting..... 9
Reagent Stability.....9
Assay Characteristics..... 10-12
Example of Test Results..... 13
References.....14
Related Products..... 15

Intended Use

The MBL Research Product **CircuLex Human FABP4/A-FABP ELISA Kit** is used for the quantitative measurement of human FABP4/A-FABP in serum and cell culture supernatant.

Individual users should determine appropriate conditions when using other types of samples.

This assay kit is for research use only and not for use in diagnostic or therapeutic procedures.

Storage

- Upon receipt store all components at 4°C.
- Do not expose reagents to excessive light.

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Introduction

Adipocyte-specific fatty acid–binding protein (A-FABP), also designated aP2 and FABP4, belongs to the fatty acid-binding protein super family whose members have relative molecular masses of 15,000, and it is exclusively expressed in differentiated adipocytes (1, 2). FABP4 is a predominant cytosolic protein of mature adipocytes, accounting for 6% of total cellular proteins. This protein may be an important regulator of systemic insulin sensitivity and lipid and glucose metabolism (1). Mice deficient in FABP4/A-FABP are protected from development of hyperinsulinemia, hyperglycemia, and insulin resistance in the context of both dietary and genetic obesity (3, 4). Adipocytes obtained from FABP4/A-FABP-null mice had markedly reduced efficiency of lipolysis in vivo and in vitro (5, 6) and exhibited a 2- to 3-fold decrease in fatty acid release, suggesting that FABP4/A-FABP mediates efflux of fatty acids in normal physiology (7).

Although the physiological consequences of FABP4/A-FABP deficiency have been predominantly linked to changes in adipocytes, it has reported that the presence of FABP4/A-FABP in macrophages (2, 8) and has shown that FABP4/A-FABP expression can be induced by peroxisome proliferator activated receptor gamma (PPAR gamma) agonists (8), by toll-like receptor agonists (9), oxidized LDL (10), and the differentiation of monocytes to macrophages and can be suppressed by treatment with a cholesterol-lowering statin (11). In these cells, FABP4/A-FABP modulates inflammatory cytokine production and cholesterol ester accumulation (12).

In apolipoprotein E-deficient mice, ablation of the FABP4/A-FABP gene conferred remarkable protection against atherosclerosis, which commonly occurs in this mouse strain (13, 14). Taken together, these animal studies demonstrate that FABP4/A-FABP, by integrating metabolic and inflammatory pathways, provides a key link between various components of metabolic syndrome. Moreover, it was reported that an orally active small-molecule inhibitor of FABP4/A-FABP is an effective therapeutic agent against severe atherosclerosis and type 2 diabetes in mouse models (15).

Principle of the Assay

The MBL Research Product **CircuLex Human FABP4/A-FABP ELISA Kit** employs the quantitative sandwich enzyme immunoassay technique. An antibody specific for human FABP4/A-FABP is pre-coated onto a microplate. Standards and samples are pipetted into the wells and the immobilized antibody binds any FABP4/A-FABP present. After washing away any unbound substances, an HRP conjugated antibody specific for human FABP4/A-FABP is added to the wells. Following a wash to remove any unbound antibody HRP conjugate, the remaining conjugate is allowed to react with the substrate H_2O_2-tetramethylbenzidine. The reaction is stopped by addition of acidic solution and absorbance of the resulting yellow product is measured at 450 nm. The absorbance is proportional to the concentration of FABP4/A-FABP. A standard curve is constructed by plotting absorbance values versus FABP4/A-FABP concentrations of calibrators, and concentrations of unknown samples are determined using this standard curve.

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Summary of Procedure

Add 100 µL of diluted samples to the wells
↓ Incubate for 2 hours at room temp.
Wash the wells
↓
Add 100 µL of HRP conjugated anti-FABP4/A-FABP antibody
↓ Incubate for 1 hour at room temp.
Wash the wells
↓
Add 100 µL of Substrate Reagent
↓ Incubate for 10-20 minutes at room temp.
Add 100 µL of Stop Solution
↓
Measure absorbance at 450 nm

Materials Provided

All samples and standards should be assayed in duplicate. The following components are supplied and are sufficient for the one 96-well microplate kit.

Microplate: One microplate supplied ready to use, with 96 wells (12 strips of 8-wells) in a foil, zip-lock bag with a desiccant pack. Wells are coated with anti-human FABP4/A-FABP monoclonal antibody as a capture antibody.

10X Wash Buffer: One 100 mL bottle of 10X buffer containing Tween®-20

Dilution Buffer: One bottle containing 50 mL of 1X buffer; use for sample dilution. Ready to use.

Human FABP4 Standard: Two vials, each vial containing X* ng of lyophilized recombinant human FABP4/A-FABP
***The amount is changed depending on lot. See the real "User's Manual" included in the kit box.**

HRP conjugated Detection Antibody: One bottle containing 12 mL of HRP (horseradish peroxidase) conjugated anti-human FABP4/A-FABP monoclonal antibody. Ready to use.

Substrate Reagent: One bottle containing 20 mL of the chromogenic substrate, tetra-methylbenzidine (TMB). Ready to use.

Stop Solution: One bottle containing 20 mL of 1 N H_2SO_4. Ready to use.

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Materials Required but not Provided

- **Pipettors:** 2-20 µL, 20-200 µL and 200-1,000 µL precision pipettors with disposable tips.
- **Precision repeating pipettor**
- **Orbital microplate shaker**
- **Microcentrifuge and tubes** for sample preparation.
- **Vortex mixer**
- **(Optional) Microplate washer:** Manual washing is possible but not preferable.
- **Plate reader** capable of measuring absorbance in 96-well plates at dual wavelengths of 450 nm/540 nm. Dual wavelengths of 450/550 or 450/595 nm can also be used. The plate can also be read at a single wavelength of 450 nm, which will give a somewhat higher reading.
- **(Optional) Software package facilitating data generation and analysis**
- **500 or 1,000 mL graduated cylinder**.
- **Reagent reservoirs**
- **Deionized water of the highest quality**
- **Disposable paper towels**

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Precautions and Recommendations

- **Although we suggest to conduct experiments as outlined below, the optimal experimental conditions will vary depending on the parameters being investigated, and must be determined by the individual user.**
- Allow all the components to come to room temperature before use.
- All microplate strips that are not immediately required should be returned to the zip-lock pouch, which must be carefully resealed to avoid moisture absorption.
- Do not use kit components beyond the indicated kit expiration date.
- Use only the microtiter wells provided with the kit.
- Rinse all detergent residue from glassware.
- Use deionized water of the highest quality.
- Do not mix reagents from different kits.
- The buffers and reagents in this kit may contain preservatives or other chemicals. Care should be taken to avoid direct contact with these reagents.
- Do not mouth pipette or ingest any of the reagents.
- Do not smoke, eat, or drink when performing the assay or in areas where samples or reagents are handled.
- Dispose of tetra-methylbenzidine (TMB) containing solutions in compliance with local regulations.
- Avoid contact with the acidic Stop Solution and Substrate Solution, which contains hydrogen peroxide.
- Wear gloves and eye protection when handling immunodiagnostic materials and samples of mouse origin, and these reagents. In case of contact with the Stop Solution and the Substrate Solution, wash skin thoroughly with water and seek medical attention, when necessary.
- **Biological samples may be contaminated with infectious agents. Do not ingest, expose to open wounds or breathe aerosols. Wear protective gloves and dispose of biological samples properly.**
- **CAUTION: Sulfuric Acid is a strong acid. Wear disposable gloves and eye protection when handling Stop Solution.**

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Sample Collection and Storage

Serum: Use a serum separator tube and allow samples to clot for 60 ± 30 minutes. Centrifuge the samples at 4°C for 10 minutes at 1,000 x *g*. Remove serum and assay immediately or store samples on ice for up to 6 hours before assaying. Aliquots of serum may also be stored at below -70°C for extended periods of time. Avoid repeated freeze-thaw cycles.

Cell culture supernatant: Remove any particulates by centrifugation and assay immediately or aliquot and store samples at below -70°C. Avoid repeated freeze-thaw cycles.

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Detailed Protocol

The MBL Research Product **CircuLex Human FABP4/A-FABP ELISA Kit** is provided with removable strips of wells so the assay can be carried out on separate occasions using only the number of strips required for the particular determination. Since experimental conditions may vary, an aliquot of Human FABP4 Standard within the kit, should be included in each assay as a calibrator. Disposable pipette tips and reagent troughs should be used for all liquid transfers to avoid cross-contamination of reagents or samples.

Preparation of Working Solutions

All reagents need to be brought to room temperature prior to the assay. Assay reagents are supplied ready-to-use, with the exception of **10X Wash Buffer** and **Human FABP4 Standard**.

1. Prepare a working solution of Wash Buffer by adding 100 mL of the **10X Wash Buffer** to 900 mL of deionized (distilled) water (**ddH$_2$O**). Mix well. Store at 4°C for two weeks or -20°C for long-term storage.

2. Reconstitute **Human FABP4 Standard** with **X* µL** of **ddH$_2$O** by gently mixing. After reconstitution, immediately dispense it in small aliquots (e.g. 130 µL) to plastic micro-centrifuge tubes and store below -70°C to avoid non-specific adsorption to glass surface and multiple freeze-thaw cycles. The concentration of the reconstituted Human FABP4 Standard should be **12 ng/mL**, which is referred to as the **Master Standard** of human FABP4/A-FABP.

***The volume is changed depending on lot. See the real "User's Manual" included in the kit box.**

Prepare Standard Solutions as follows:
Use **Master Standard** to produce a dilution series (below). Mix each tube thoroughly before the next transfer. **Std.1 (2,400 pg/mL)** serves as the highest standard. **Dilution Buffer** serves as the zero standard (Blank).

	Volume of Standard	Dilution Buffer	Concentration
Std.1	120 µL of Master Standard	480 µL	2,400 pg/mL
Std.2	300 µL of Std. 1 (2,400 pg/mL)	300 µL	1,200 pg/mL
Std.3	300 µL of Std. 2 (1,200 pg/mL)	300 µL	600 pg/mL
Std.4	300 µL of Std. 3 (600 pg/mL)	300 µL	300 pg/mL
Std.5	300 µL of Std. 4 (300 pg/mL)	300 µL	150 pg/mL
Std.6	300 µL of Std. 5 (150 pg/mL)	300 µL	75 pg/mL
Std.7	300 µL of Std. 6 (75 pg/mL)	300 µL	37.5 pg/mL
Blank	-	300 µL	0 pg/mL

Note: Do not use a Repeating pipette. Change tips for every dilution. Wet tip with **Dilution Buffer** before dispensing. Discard any unused Standard Solutions after use.

Sample Preparation

Dilute samples with **Dilution Buffer**.
- Serum samples may require a 30-fold dilution.
- Cell culture supernatants require appropriate dilution.

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Assay Procedure

1. Remove the appropriate number of microtiter wells from the foil pouch and place them into the well holder. Return any unused wells to the foil pouch, refold, seal with tape and store at 4°C.

2. Dilute samples with **Dilution Buffer**. (See "Sample Preparation" above.)

3. Pipette **100 µL** of **Standard Solutions (Std1-Std7, Blank)** and **diluted samples** in duplicates, into the appropriate wells.

4. Incubate the plate **at room temperature (ca.25°C) for 2 hours**, shaking at ca. 300 rpm on an orbital microplate shaker.

5. Wash 4-times by filling each well with Wash Buffer (350 µL) using a squirt bottle, multi-channel pipette, manifold dispenser or microplate washer.

6. Add **100 µL** of **HRP conjugated Detection Antibody** into each well.

7. Incubate the plate **at room temperature (ca.25°C) for 1 hour**, shaking at ca. 300 rpm on an orbital microplate shaker.

8. Wash 4-times by filling each well with Wash Buffer (350 µL) using a squirt bottle, multi-channel pipette, manifold dispenser or microplate washer.

9. Add **100 µL** of **Substrate Reagent**. Avoid exposing the microtiter plate to direct sunlight. Covering the plate with e.g. aluminum foil is recommended. Return Substrate Reagent to 4°C immediately after the necessary volume is removed

10. Incubate the plate **at room temperature (ca.25°C) for 10-20 minutes**, shaking at ca. 300 rpm on an orbital microplate shaker. The incubation time may be extended up to 30 minutes if the reaction temperature is below than 20°C.

11. Add **100 µL** of **Stop Solution** to each well in the same order as the previously added Substrate Reagent.

12. Measure absorbance in each well using a spectrophotometric microplate reader at dual wavelengths of 450/540 nm. Dual wavelengths of 450/550 or 450/595 nm can also be used. Read the microplate at 450 nm if only a single wavelength can be used. Wells must be read within 30 minutes of adding the Stop Solution.

Note-1: Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining Wash Buffer by aspirating or decanting. Invert the plate and blot it against clean paper towels.

Note-2: Reliable standard curves are obtained when either O.D. values do not exceed 0.2 units for the blank (zero concentration), or 3.0 units for the highest standard concentration.

Note-3: If the microplate reader is not capable of reading absorbance greater than the absorbance of the highest standard, perform a second reading at 405 nm. A new standard curve, constructed using the values measured at 405 nm, is used to determine the concentration of off-scale samples. The readings at 405 nm should not replace the on-scale readings at 450 nm.

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Calculations

Average the duplicate readings for each standard, control and sample, and subtract the optical density of the average zero standard. Plot the optical density versus the concentration of standards and draw the best curve. Most microtiter plate readers perform automatic calculations of analyte concentration. The standard curve fits best to a sigmoidal four-parameter logistic equation. The results of unknown samples can be calculated with any computer program having a four-parameter logistic function.

A standard curve is also to be constructed by plotting the absorbance (Y) versus log of the known concentration (X) of standards, using a cubic function. Alternatively, the logit log function can be used to linearize the standard curve (i.e. logit of optical density (Y) is plotted versus log of the known concentration (X) of standards). To determine the concentration of each sample, first find the optical density on the y-axis and extend a horizontal line to the standard curve. At the point of intersection, extend a vertical line to the x-axis and read the corresponding concentration.

If the samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor.

Measurement Range

The measurement range is 37.5 pg/mL to 2,400 pg/mL. Any sample reading higher than the highest standard should be diluted with Dilution Buffer in higher dilution and re-assayed. Dilution factors need to be taken into consideration in calculating the human FABP4/A-FABP concentration.

Troubleshooting

1. All samples and controls should be assayed in duplicate, using the protocol described in the **Detailed Protocol**. Incubation times or temperatures significantly different from those specified may give erroneous results.

2. Poor duplicates, accompanied by elevated values for wells containing no sample, indicate insufficient washing. If all instructions in the **Detailed Protocol** were followed accurately, such results indicate a need for washer maintenance.

3. Overall low signal may indicate that desiccation of the plate has occurred between the final wash and addition of Substrate Reagent. Do not allow the plate to dry out. Add Substrate Reagent immediately after wash.

Reagent Stability

All of the reagents included in the MBL Research Product **CircuLex Human FABP4/A-FABP ELISA Kit** have been tested for stability. Reagents should not be used beyond the stated expiration date.

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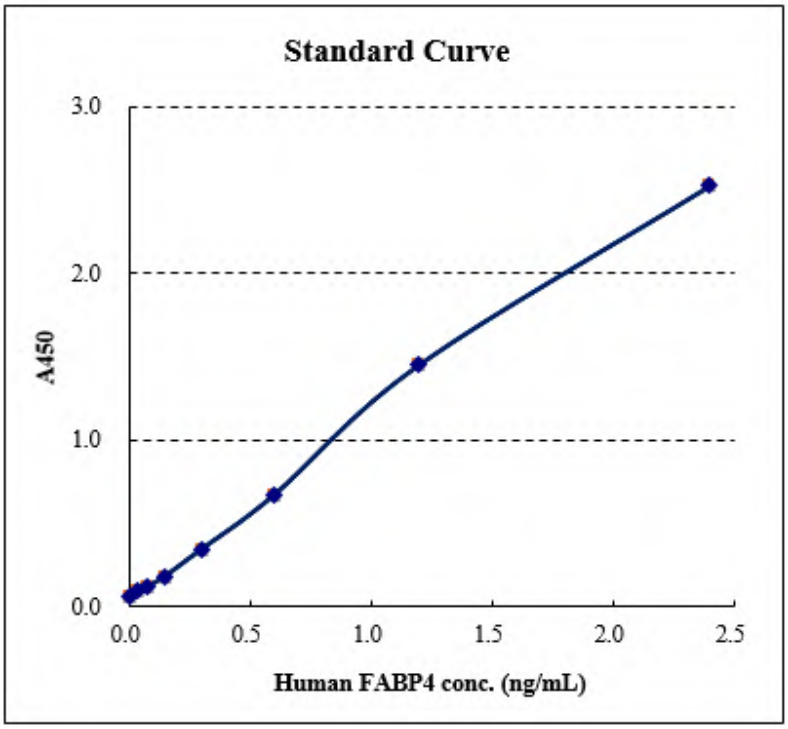
Assay Characteristics

1. Sensitivity

The limit of detection (defined as such a concentration of human FABP4/A-FABP giving absorbance higher than mean absorbance of blank* plus three standard deviations of the absorbance of blank: A blank + 3SD blank) is better than 27.8 pg/ml of sample.

* Dilution Buffer was pipetted into blank wells.

Typical Standard curve



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2. Precision

Intra-assay Precision (Precision within an assay)

Three samples of known concentration were tested sixteen times on one plate to assess intra-assay precision.

• Intra-assay (Within-Run, n=16)

	Human FABP4 conc. (ng/mL)		
	Serum 1	Serum 2	Serum 3
1	49.32	22.77	10.65
2	48.77	23.04	10.75
3	48.31	23.11	10.93
4	49.89	24.10	11.39
5	49.92	24.00	11.51
6	49.17	23.54	11.48
7	47.06	23.73	11.06
8	49.06	22.11	10.93
9	49.52	22.47	10.46
10	48.19	22.40	10.68
11	49.00	22.47	10.95
12	51.43	23.64	11.26
13	49.72	23.80	10.95
14	50.23	23.78	11.33
15	50.78	23.43	11.28
16	51.04	22.48	10.22
MAX.	**51.43**	**24.10**	**11.51**
MIN.	**47.06**	**22.11**	**10.22**
MEAN	**49.46**	**23.18**	**10.99**
S.D.	**1.12**	**0.65**	**0.37**
C.V.	**2.3%**	**2.8%**	**3.4%**

Inter-assay Precision (Precision between assays)

Three samples of known concentration were tested in five separate assays to assess inter-assay precision.

• Inter-assay (Run-to-Run, n=5)

	Human FABP4 conc. (ng/mL)		
	Serum 1	Serum 2	Serum 3
1	8.6	18.8	38.2
2	7.5	17.1	36.5
3	9.1	17.0	38.0
4	8.4	18.1	38.0
5	8.6	18.9	42.0
MAX.	**9.1**	**18.9**	**42.0**
MIN.	**7.5**	**17.0**	**36.5**
MEAN	**8.5**	**18.0**	**38.5**
S.D.	**0.59**	**0.91**	**2.08**
C.V.	**7.0%**	**5.1%**	**5.4%**

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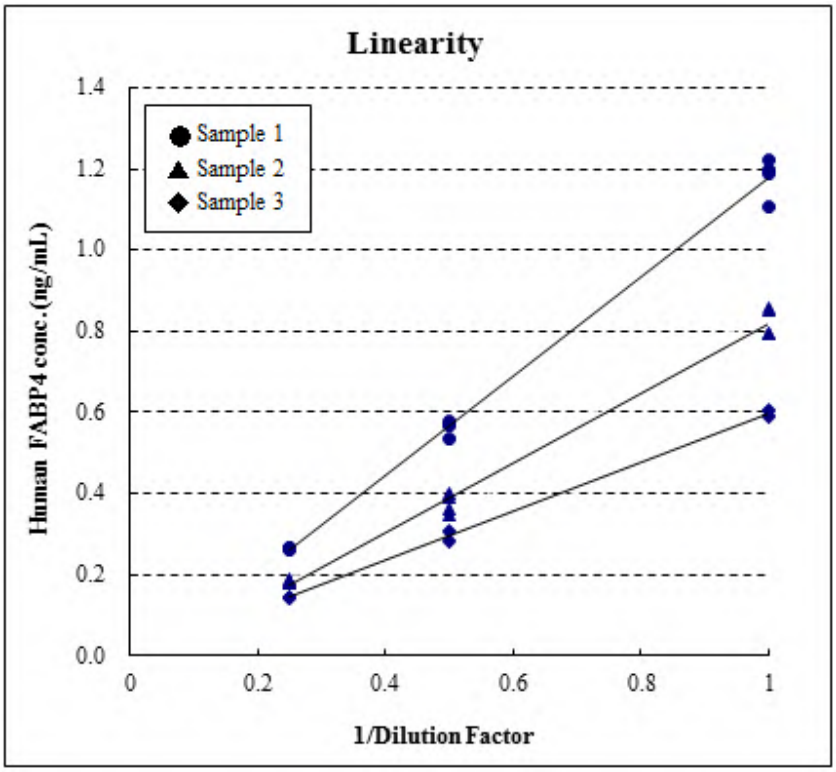
3. Linearity

To assess the linearity of the assay, three samples* containing and/or spiked with human FABP4/A-FABP were 30-fold diluted with Dilution Buffer and assayed.

* Sample: Serum



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Example of Test Results

Fig.1 Human FABP4/A-FABP concentration in sera from 38 healthy volunteers.

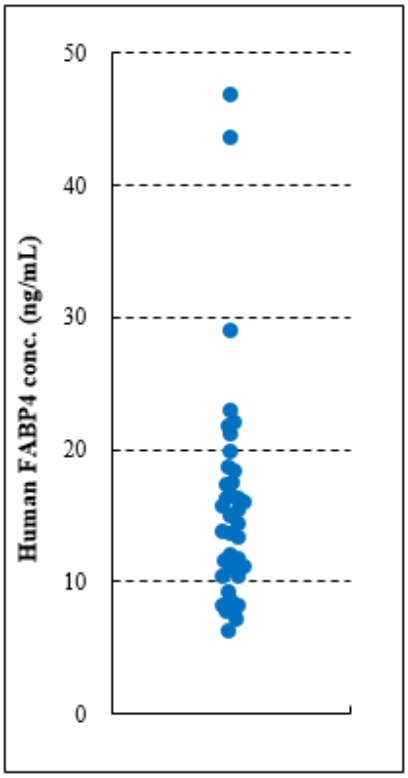
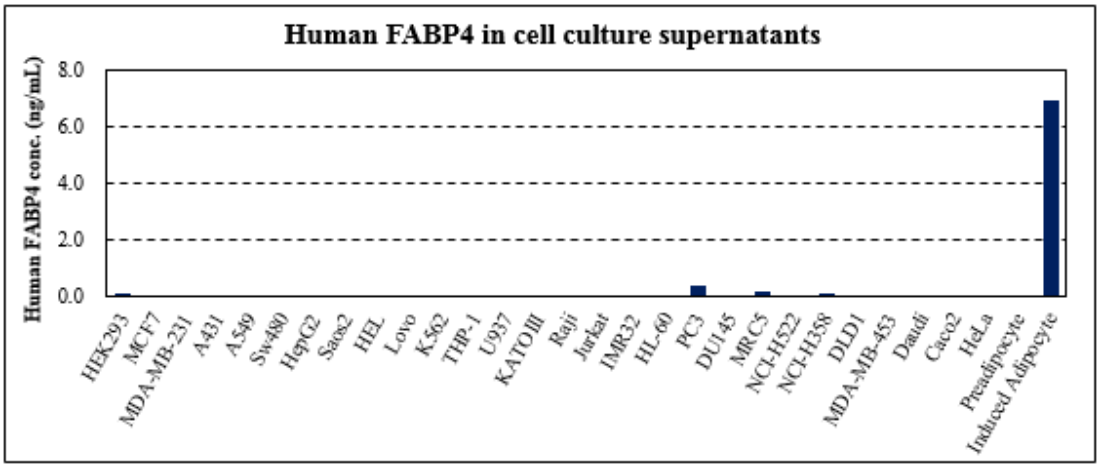


Fig.2 Human FABP4/A-FABP concentration in cell culture supernatants.



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Related Products

***CircuLex Mouse FABP1/L-FABP ELISA Kit: Cat# CY-8054**
***CircuLex Mouse FABP3/H-FABP ELISA Kit: Cat# CY-8055**
***CircuLex Mouse FABP5/E-FABP/mal1 ELISA Kit: Cat# CY-8056**
***CircuLex Mouse FABP4/A-FABP ELISA Kit: Cat# CY-8077**
***CircuLex Rat FABP4/A-FABP ELISA Kit: Cat# CY-8076**
***CircuLex Human FABP5/E-FABP ELISA Kit: Cat# CY-8116**
***CircuLex Mouse NGAL ELISA Kit: Cat# CY-8053**
***CircuLex Human NGAL ELISA Kit: Cat# CY-8070**
***CircuLex Rat Adiponectin ELISA Kit: Cat# CY-8049**
***CircuLex Mouse Adiponectin ELISA Kit: Cat# CY-8051**
***CircuLex Human Adiponectin ELISA Kit: Cat# CY-8050**
***CircuLex S100A12/EN-RAGE ELISA Kit: Cat# CY-8058**
***CircuLex S100A4 ELISA Kit: Cat# CY-8059**
***CircuLex S100P ELISA Kit: Cat# CY-8060**
***CircuLex S100A8 ELISA Kit: Cat# CY-8061**
***CircuLex S100A9 ELISA Kit: Cat# CY-8062**
***CircuLex S100A7 (psoriasin) ELISA Kit: Cat# CY-8073**
***CircuLex CML/Nε-(carboxymethyl)lysine ELISA Kit: Cat# CY-8066**
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