

APB985Hu01 100µg
Active Fatty Acid Binding Protein 5 (FABP5)
Organism Species: *Homo sapiens* (Human)
Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

13th Edition (Revised in Aug, 2023)

[PROPERTIES]

Source: Prokaryotic expression.

Host: *E. coli*

Residues: Ala2~Glu135

Tags: N-terminal His-tag

Purity: >90%

Endotoxin Level: <1.0EU per 1µg (determined by the LAL method).

Buffer Formulation: PBS, pH7.4, containing 0.01% Sarcosyl, 5%Trehalose .

Original Concentration: 200µg/mL

Applications: Activity Assays.

(May be suitable for use in other assays to be determined by the end user.)

Predicted isoelectric point: 7.0

Predicted Molecular Mass: 16.3kDa

Accurate Molecular Mass: 16kDa as determined by SDS-PAGE reducing conditions.

[USAGE]

Reconstitute in 10mM PBS (pH7.4) to a concentration of 0.1-1.0 mg/mL. Do not vortex.

[STORAGE AND STABILITY]

Storage: Avoid repeated freeze/thaw cycles.

Store at 2-8°C for one month.

Aliquot and store at -80°C for 12 months.

Stability Test: The thermal stability is described by the loss rate. The loss rate

was determined by accelerated thermal degradation test, that is, incubate the protein at 37°C for 48h, and no obvious degradation and precipitation were observed. The loss rate is less than 5% within the expiration date under appropriate storage condition.

[SEQUENCE]

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ATVQQLLEGR WRLVDSKGF D EYMKELGVGI ALRKMAMAK PDCIITCDGK  
NLTIKTESTL KTTQFSCTLG EKFEETTADG RKTQTVCNFT DGALVQHQE W  
DGKESTITRK LKDGKLVVEC VMNVTCTRI YEKVE
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[ACTIVITY]

Fatty Acid Binding Protein 5 (FABP5) , is a 15-kDa cytosolic protein abundant in epidermal keratinocytes, macrophages, and adipocytes. As a member of the FABP family, it selectively binds long-chain fatty acids and retinoids, shuttling ligands to nuclear receptors to regulate gene transcription. In the skin, FABP5 drives epidermal differentiation and inflammation by modulating lipid mediator biosynthesis . It also acts as a redox sensor, protecting against oxidative stress in psoriasis and wound healing. Furthermore, the interaction between FABP5 and S100A7 augments pro-inflammatory signaling and cancer cell migration by orchestrating lipid-mediated pathways and inflammatory cytokine production. Thus a functional ELISA assay was conducted to detect the interaction of recombinant human FABP5 and recombinant human S100A7. Briefly, FABP5 was diluted serially in PBS with 0.01% BSA (pH 7.4). Duplicate samples of 100 μ l were then transferred to S100A7-coated microtiter wells and incubated for 1h at 37°C . Wells were washed with PBST and incubated for 1h with anti-LOX pAb, then aspirated and washed 3 times. After incubation with HRP labelled secondary antibody for 1h at 37°C , wells were aspirated and washed 5 times. With the addition of substrate solution, wells were incubated 15-25 minutes at 37°C . Finally, add 50 μ L stop solution to the wells and read at 450/630nm immediately. The binding activity of recombinant human FABP5 and recombinant human S100A7 was shown in Figure 1, the EC50 for this effect is 0.298ug/mL.

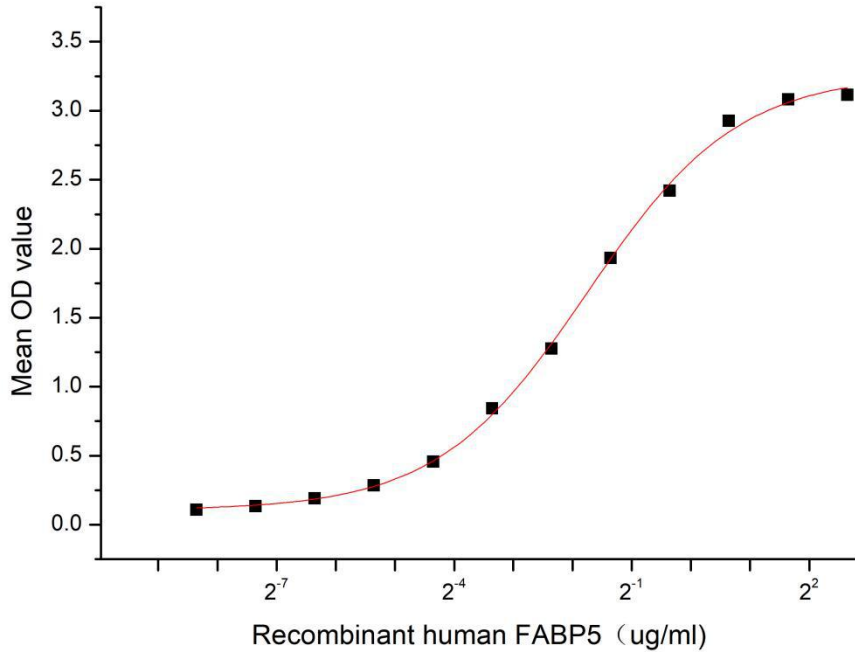


Figure 1. The binding activity of recombinant human FABP5 and recombinant human S100A7

[IDENTIFICATION]

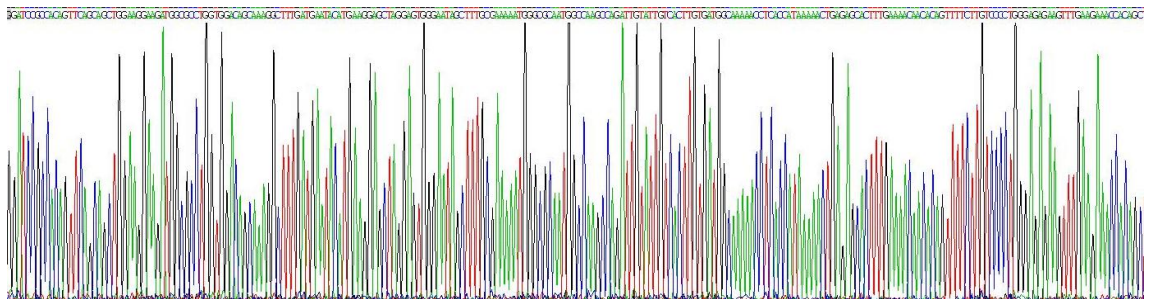


Figure 2. Gene Sequencing (extract)

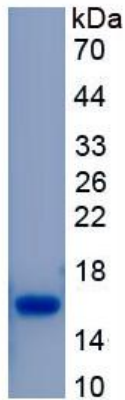


Figure 3. SDS-PAGE

Sample: Active recombinant FABP5, Human

[IMPORTANT NOTE]

The kit is designed for research use only, we will not be responsible for any issue if the kit was used in clinical diagnostic or any other procedures.