

APA037Hu61 200µg

Active Fibronectin (FN)

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: Eukaryotic expression.

Host: 293F cell

Residues: Ser607~Pro1265

Tags: N-terminal His-tag

Purity: >95%

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

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

Original Concentration: 500µg/mL

Applications: Activity Assays.

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

Predicted isoelectric point: 4.5

Predicted Molecular Mass: 73.4kDa

Accurate Molecular Mass: 90kDa as determined by SDS-PAGE reducing conditions.

Phenomenon explanation:

The possible reasons that the actual band size differs from the predicted are as follows:

1. Splice variants: Alternative splicing may create different sized proteins from the same gene.
2. Relative charge: The composition of amino acids may affects the charge of the protein.
3. Post-translational modification: Phosphorylation, glycosylation, methylation etc.
4. Post-translation cleavage: Many proteins are synthesized as pro-proteins, and then cleaved to give the active form.
5. Polymerization of the target protein: Dimerization, multimerization etc.

[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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SSSGPVEVFITETPSQPNSHPIQWNAQPQSHISKYILRWRPKNSVGRWKEATIPGHLNSYTIKGLKPGVVYEGQLISIQYGHQEVTRF  
DFTTTSTSTPVTSTNTVTGETTTFSPVLATSESVTEITASSFVVSWSASDTVSGFRVEYELSEEGDEPQYLDLPSTATSVNIPDLLPGR  
KYIVNVYQISEDGEQSLILSTSTQTTAPDAPPDPTVDQVDDTSIVRWSRPQAPITGYRIVYSPSVEGSSTELNLPETANSVTLSDLQPG  
VQYNITIIYAVEENQESTPVVVIQETTGTPRSDTVPSRDLQFVEVTDVKVTIMWTPPEASVTCYRVDVIPVNLPGEHGQRLPISRNTFA  
EVTGLSPGVTTYFKVFAVSHGRESKPLTAQQTTKLDAPTNLQFVNETDSTVLVRWTPPPRAQITGYRLTVGLTRRGQPRQYNVGPVSKY  
PLRNLQPASEYTVSLVAIKGNQESPKATGVFTTLQPGSSIPPYNTEVTETTIVITWTWPAPRIGFKLGVRPSQGGGEAPREVTSDSGSIVV  
SGLTPGVEYVYTIQVLRDQGERDAPIVNKVVTPLSPTNLHLEANPDGTGLTVSWERSTTPDITGYRITTTPTNGQQGNLSLEEVVHADQ  
SSCTFDNLSPGLEYNVSVYTVKDDKESVPISDTIIP
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[ACTIVITY]

Fibronectin (FN) is a ubiquitous extracellular matrix glycoprotein encoded by the FN1 gene in mammals. It exists as soluble plasma fibronectin and insoluble cellular fibronectin assembled into fibrils. FN contains multiple functional domains that mediate interactions with cell-surface integrins, collagens and proteoglycans. It participates in cell adhesion, migration, proliferation and tissue remodeling, playing vital roles in wound healing, embryonic development, angiogenesis and tumor progression. Dysregulated FN expression and fibrillogenesis drive inflammation, fibrosis and cancer metastasis. Fibronectin can directly bind vitronectin (VTN), and this complex jointly regulates cell anchoring and extracellular matrix organization. Briefly, VTN was diluted serially in PBS with

0.01% BSA (pH 7.4). Duplicate samples of 100 μ L were then transferred to FN-coated microtiter wells and incubated for 1h at 37°C. Wells were washed with PBST and incubated for 1h with anti-VTN 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. Measured by its binding ability in a functional ELISA. When recombinant human FN is immobilized at 2 μ g/mL (100 μ L/well), the concentration of VTN that produces 50% optimal binding-response is found to be approximately 5.153 μ g/mL.

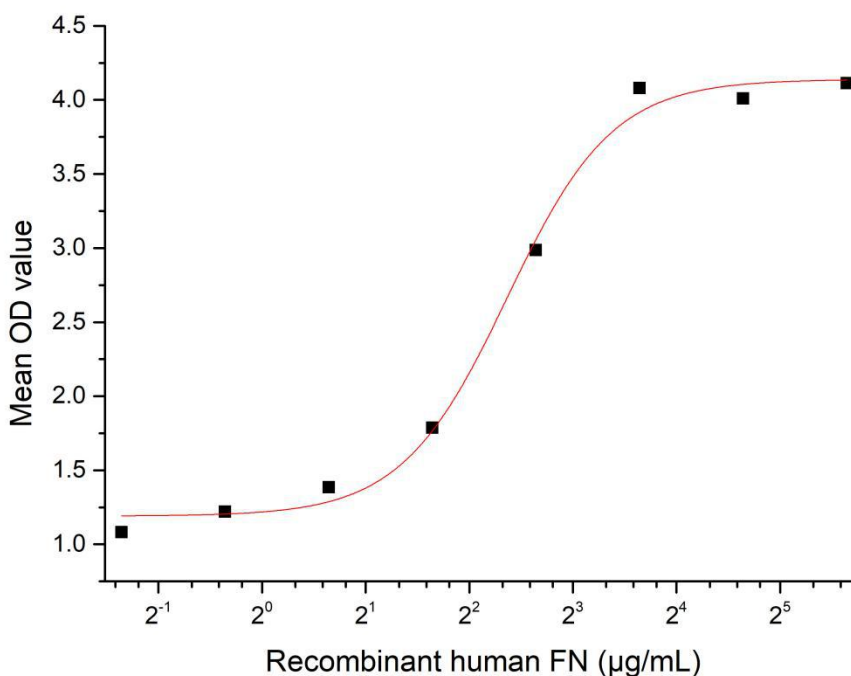


Figure 1. The binding activity of recombinant human FN and mouse VTN

SDS-PAGE gel showing a single band at approximately 90 kDa. Molecular weight markers are indicated on the left at 250, 150, 100, 70, and 50 kDa.

Sample: Active recombinant FN, Human

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.