APC909Mu01 100μg
Active Fibroblast Growth Factor 22 (FGF22)
Organism Species: Mus musculus (Mouse)
Instruction manual

FOR IN VITRO USE AND RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1th Edition (Apr, 2016)

[PROPERTIES]

Source: Prokaryotic expression.
Host: *E. coli*
Residues: His26~Ser162
Tags: N-terminal His-tag
Purity: >95%
Buffer Formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 0.05% sarcosyl and 5% trehalose.
Applications: Cell culture; Activity Assays.
(May be suitable for use in other assays to be determined by the end user.)
Predicted isoelectric point: 11.7
Predicted Molecular Mass: 19.9kDa
Accurate Molecular Mass: 20kDa as determined by SDS-PAGE reducing conditions.

[USAGE]

Reconstitute in 20mM Tris, 150mM NaCl (pH8.0) 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^{o}C for one month.
Aliquot and store at -80^{o}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 37ºC 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]

HLEGD VRWRRLFSST HFFLRVDLGG
RVQGTRWRHG QDSIVEIRSV RVGTVVIKAV YSGFYVAMNR RGRLYGSRVY
SVDCRFRERI EENGYNTYAS RRWRHRGRPM FLALDSQGIP RQGRRTRRHQ
LSTHFLPVLV SS

[ACTIVITY]

FGF22 (Fibroblast growth factor 22) is a member of the fibroblast growth factor (FGF) family. FGF family members possess broad mitogenic and cell survival activities and are involved in a variety of biological processes including embryonic development, cell growth, morphogenesis, tissue repair, tumor growth and invasion. A proliferation assay was conducted to detect the bioactivity of recombinant mouse FGF22 using 3T3 cells. Briefly, 3T3 cells were seeded into triplicate wells of 96-well plates at a density of 2,000 cells/well and allowed to attach overnight, then the medium was replaced with serum-free standard DMEM prior to the addition of various concentrations of FGF22. After incubated for 48h, cells were observed by inverted microscope and cell proliferation was measured by Cell Counting Kit-8 (CCK-8). Briefly, 10μL of CCK-8 solution was added to each well of the plate, then the absorbance at 450nm was measured using a microplate reader after incubating the plate for 1-4 hours at 37℃. Proliferation of 3T3 cells after incubation with FGF22 for 48h observed by inverted microscope was shown in Figure 1. Cell viability was assessed by CCK-8 (Cell Counting Kit-8) assay after incubation with recombinant FGF22 for 48h. The result was shown in Figure 2. It was obvious that FGF22 significantly increased cell viability of 3T3 cells.

A B

Figure 1. Cell proliferation of 3T3 cells after stimulated with FGF22.

(A) 3T3 cells cultured in DMEM, stimulated with 1000ng/mL FGF22 for 48h;

(B) Unstimulated 3T3 cells cultured in DMEM for 48h.

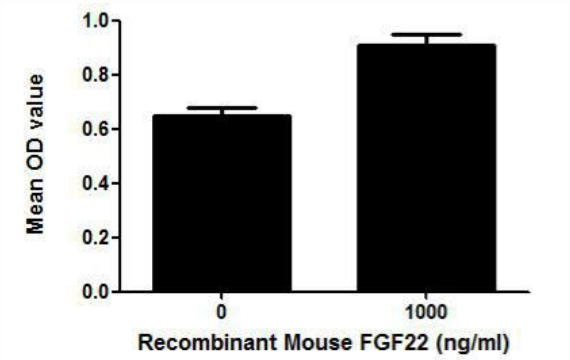

Figure 2. Cell proliferation of 3T3 cells after stimulated with FGF22.

[IDENTIFICATION]

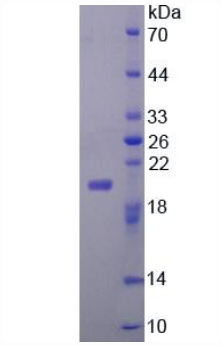

Figure 3. SDS-PAGE

Sample: Active recombinant FGF22, Mouse

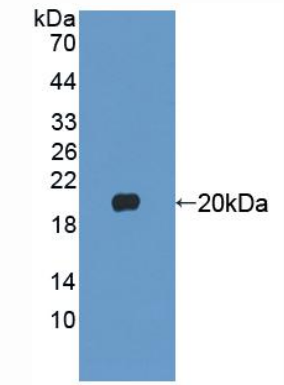

Figure 4. Western Blot

Sample: Recombinant FGF22, Mouse;

Antibody: Rabbit Anti-Mouse FGF22 Ab (PAC909Mu01)