

EPA133Ra61 100µg
Eukaryotic Tumor Necrosis Factor Alpha (TNFα)
Organism Species: *Rattus norvegicus* (Rat)
Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

12th Edition (Revised in Aug, 2016)

[PROPERTIES]

Source: Eukaryotic expression

Host: CHO Cell

Residues: Leu80~Leu235

Tags: N-terminal His Tag and C-terminal Fc Region of Human IgG1

Subcellular Location: Secreted

Purity: > 97%

Traits: Freeze-dried powder

Buffer formulation: 20mM Tris, 150mM NaCl, pH8.0, containing 0.01% SKL, 5% Trehalose.

Original Concentration: 200µg/mL

Applications: Positive Control; Immunogen; SDS-PAGE; WB.

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

Predicted isoelectric point: 4.7

Predicted Molecular Mass: 47.8kDa

Accurate Molecular Mass: 25&18kDa 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 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°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**]

```

                                L R S S Q N S S D K P V A H V V A N H Q
A E E Q L E W L S Q R A N A L L A N G M D L K D N Q L V V P A D G L Y L I Y S Q V L F K G Q G C P D
Y V L L T H T V S R F A I S Y Q E K V S L L S A I K S P C P K D T P E G A E L K P W Y E P M Y L G G
V F Q L E K G D L L S A E V N L P K Y L D I T E S G Q V Y F G V I A L
  
```

[**IDENTIFICATION**]

T T C G A T A C T C T C A A C T C G G C A C C C G G C C T G T G C G A C C C C C C G G C G C T C G G C T C C G C G G C T C A A C C C C C T C T C T G A T G C A T C T A A G C A C T A C T C G G A C C G A G T C C G T G A C T T A C C C C G G T C T C T T A G G G C A A G C T C C C G A C T A G C T C C C C G G C G T T T
 L R S S Q N S S D K P V A H V V A N H Q A E E Q L E W L S Q R A N A L L A N G M D L K D N Q L V V P A D G L Y L I Y S Q V L F K G Q G C P D Y V L L T H T V S R F

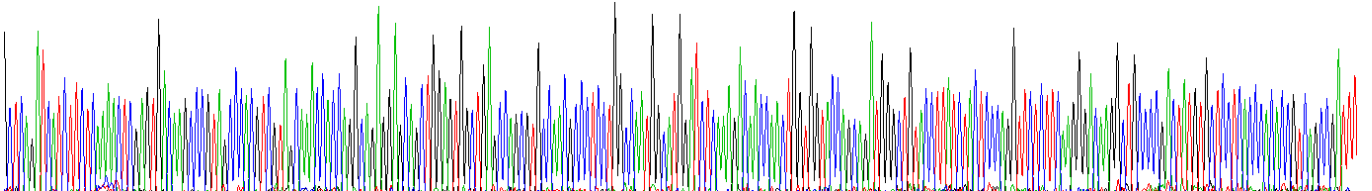


Figure . Gene Sequencing (extract)

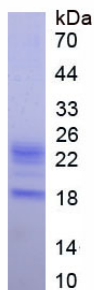


Figure. SDS-PAGE

[**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.