

**EPX274Ge61 50µg**

**Eukaryotic Nucleoprotein (NP)**

**Organism Species: *Pan-species (General)***

***Instruction manual***

FOR RESEARCH USE ONLY

NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

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12th Edition (Revised in Aug, 2016)

**[ PROPERTIES ]**

**Source:** Eukaryotic expression

**Host:** 293F cell

**Residues:** Met1~Ala419

**Tags:** N-terminal His Tag

**Subcellular Location:** Secreted

**Purity:** > 95%

**Traits:** Freeze-dried powder

**Buffer formulation:** PBS, pH7.4, containing 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:** 10.7

**Predicted Molecular Mass:** 49.3kDa

**Accurate Molecular Mass:** 60kDa 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 affect 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** ]

```
MSDNGPQNQRNAPRITFGGSPDSTGSNQNGERSGARSKQRRPQGLPNNTASWFTALTQHGKEDLKFPRGQGVPIINTNSSPDDQIGYYRR
ATRRIRGGDGKMKDLSPRWYFYLLGTGPEAGLPYGANKDGI IHWATEGALNTPKDHIGTRNPANNAIVLQLPQGTTLPKGFYAEGSRG
GSQASSRSSSRNSSRNSTPGSSRGTS PARMAGNGGDAALALLLLDRLNQLLESKMSGKQQQQGQTVTKKSAEASKKPRQKRTATKA
YNVTQAFGRRGPEQTQGNFGDQELIRQGT DYKHWPIAQFAPSASAFFGMSRIGMEVTPSGTWLTYTGAIKLDKDPNFKDQVILLNKH
IDAYKTFPPTPEPKDKKKKKADETQALPQRKQKQTVTLLPAADLDDFSKQLQQSMSSADSTQA
```

[ **IDENTIFICATION** ]

AGTTCGATTCATGAGGTAACGTCGGAAACGGGAGCCGCGGATTCCTTGGGGTCGGGAGTGGAGCTGGGACAAAGCGGAGGTAAGCGGGGGGAAAGCGGCGCGGAGGCTGCGAGAGCCCGGCGTGGTACCGCGTGGTTCACCGCGTGGTGGACCGAAAGAGGCGCGAGTCCCGCGGAGCGAGGCGCGGATTCAGCGAGG

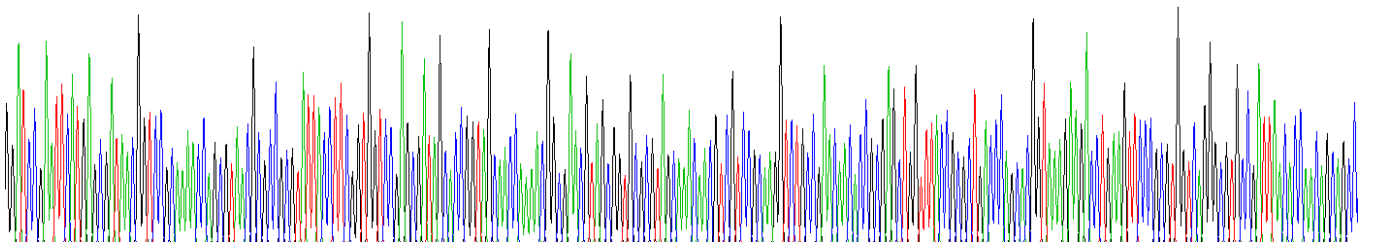


Figure . Gene Sequencing (extract)

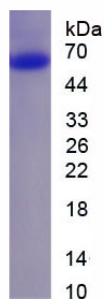


Figure. SDS-PAGE

**[ 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.