# Rhesus CCN3 / NOV (IGFBP9) Gene ORF cDNA clone in cloning vector

Catalog Number: CG90081-G



### **General Information**

Gene: nephroblastoma overexpressed gene

Official Symbol: NOV

Synonym: NOV

Source: Rhesus

cDNA Size: 1074bp

**RefSeq:** XM\_001097136.2

Plasmid: pGEM-cynoNOV

## **Description**

Lot: Please refer to the label on the tube

#### **Sequence Description:**

Identical with the Gene Bank Ref. ID sequence except for the point mutations: 991T/C(C331R); 813C/G not causing the amino acid variation. Please check the sequence information before order.

Vector:

pGEM-T

#### Quality control:

The plasmid is confirmed by full-length sequencing with primers in the sequencing primer list.

#### Sequencing primer list:

M13-47: 5' GCCAGGGTTTTCCCAGTCACGAC 3'

RV-M: 5' GAGCGGATAACAATTTCACACAGG3'

Other M13 primers can also be used as sequencing primers.

#### Shipping carrier:

Each tube contains approximately 10 µg of lyophilized plasmid.

#### Storage:

The lyophilized plasmid can be stored at ambient temperature for three months.

# **Plasmid Resuspension protocol**

- 1. Centrifuge at 5,000 × g for 5 min.
- 2. Carefully open the tube and add 100  $\mu l$  of sterile water to dissolve the DNA.
- Close the tube and incubate for 10 minutes at room temperature.
- 4. Briefly vortex the tube and then do a quick spin to concentrate the liquid at the bottom. Speed is less than  $5000\times g$ .
- 5. Store the plasmid at  $-20 \,^{\circ}$ C.

#### The plasmid is ready for:

- Restriction enzyme digestion
- PCR amplification
- E. coli transformation
- DNA sequencing

# *E.coli* strains for transformation (recommended but not limited)

Most commercially available competent cells are appropriate for the plasmid, e.g. TOP10, DH5 $\alpha$  and TOP10F $^{\prime}$ .

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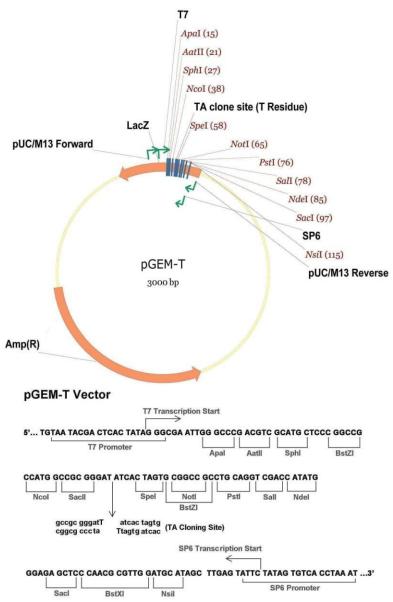


#### **Vector Information**

The pGEM-T vector is a high-efficiency TA cloning vector which contains multiple cloning sites as shown below. The pGEM-T vector is 3.0kb in size and contains the ampicillin resistance gene for selection. The coding sequence was inserted by TA cloning.

Notes: The direction of cDNA insertion into the TA-cloning vector is random, maybe forward or reverse. For insert orientation information, please feel free to contact us.

### Physical Map of pGEM-T:



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