# Porcine reproductive and respiratory syndrome virus (PRRSV) (type 2, strain ATCC VR2332 (NA)) N natural ORF mammalian expression plasmid

Sino Biological Inc. **Biological Solution Specialist** 

**Catalog Number:** VG40337-G

# **General Information**

Gene: PRRSV (type 2, strain ATCC VR2332

(NA)) N

Official Symbol: PRRSV-N

PRRSV-N Synonym:

Source: Porcine reproductive and respiratory

syndrome virus (PRRSV)

cDNA Size: 372bp

U87392.3 RefSeq:

Plasmid: pGEM-PRRSV2-AAD12131-N

# **Description**

Lot: Please refer to the label on the tube

# **Sequence Description:**

Identical with the Gene Bank Ref. ID sequence.

Vector:

pGEM-T

# Shipping carrier:

Each tube contains approximately 10 µg of lyophilized plasmid.

### Storage:

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

## **Quality control:**

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

### Sequencing primer list:

5' GCCAGGGTTTTCCCAGTCACGAC 3' M13-47:

RV-M: 5' GAGCGGATAACAATTTCACACAGG 3'

Other M13 primers can also be used as sequencing primers.

# **Plasmid Resuspension protocol**

- 1. Centrifuge at  $5,000 \times g$  for 5 min.
- 2.Carefully open the tube and add 100 µl of sterile water to dissolve the DNA.
- 3. 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 °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'.

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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 amplicin resistance gene for selection. The coding sequence was inserted by TA cloning.

### Physical Map of pGEM-T:

