# Human ACTG2 Gene cDNA clone plasmid

Catalog Number: HG10961-M



#### **General Information**

Gene: actin, gamma 2, smooth muscle, enteric

Official Symbol: ACTG2

Synonym: ACT, ACTE, ACTA3, ACTL3, ACTSG

Source: Human

cDNA Size: 1131bp

RefSeq: NM\_001615.3

Plasmid: pMD-ACTG2

### 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:1051 T/G resulting in the amino acid Leu substitution by Val and 390 T/C not causing the amino acid variation.

Vector:

pMD18-T Simple

#### 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 × 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×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, DH5α and TOP10F'.

Fax:+86-10-51029969

● Tel:+86- 400-890-9989 ● http://www.sinobiological.com

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Sino Biological Inc.
Biological Solution Specialist

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#### **Vector Information**

pMD18-T Simple Vector is a high-efficiency TA cloning vector constructed from pUC18, of which the initial multiple cloning sites (MCS) were destroyed. The pMD18-T Simple Vector is 2.6kb in size and contains the amplicin resistance gene for selection. The coding sequence was inserted by TA cloning at site 425.

#### Physical Map of pMD18-T Simple (MCS destroyed):

