

# TMEM65 Knockdown NIH/3T3 Cell Line, Heterozygous

Catalog No.: RM02196

# **Basic Information**

#### Catalog No.

RM02196

### Category

Cell Line

#### **Parental Cell line**

NIH/3T3

#### Genotype

Knockdown

## **Gene Information**

#### **Gene Symbol**

TMEM65

# **Species**

Mouse

# Gene ID

74868

#### **Synonyms**

2610029013Rik; 4930438D12Rik

## **Contact**

2	400-999-6126
$\bowtie$	cn.market@abclonal.com.cn
•	www.abclonal.com.cn

# **Background**

## **Product Information**

#### **Description**

TMEM65 Knockdown NIH/3T3 cell line is engineered from NIH/3T3 cell line with Gene-Editing Technology.

Allele-1:97bp deletion in exon1

Allele-2:WT

Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

#### **Packaging**

1 vial parental cell line and 1 vial knockout cell line

#### **Shipping Conditions**

#### **Amount**

Dry ice

1~5x10<sup>6</sup> cells/vial

#### Storage

Stored in liquid nitrogen for a long time less than -130°C. Minimizing freeze-thaw cycles.

#### **Protocol**

Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at  $37^{\circ}C$  with 5% CO<sub>2</sub> condition.

- 1. Thaw the vial in 37°C water bath ,and shake it to melt as soon as possible.
- 2. Transfer the cell suspension to a 15mL conical tube with pre-warmed 5mL complete medium and centrifuge 1000rpm for approximately 5 minutes at room temperature.
- 3. Remove and discard the supernatant.
- 4. Resuspend the cell pellet with 1mL pre-warmed complete medium and seed in 10cm dish.
- 5. Add 8-10mL of complete medium.
- 6. Incubate the culture at 37°C incubator with 5% CO<sub>2</sub>.
- 7. A subcultivation ratio of 1:2-1:4 is recommended.

# Sequencing data

WT CGCTGAACACGGCG\*\*\*\*\*\*\*\*\*\*\*\*GTGGGGCCGCCTGG
Mut CGCTGAACACGGCG\*\*\*Deletion\*\*\*GTGGGGCCGCCTGG
Allele-1: 97bp deletion in exon1

WT CGCTGAACACGGCG\*\*\*\*\*\*\*\*\*GTGGGGCCGCCTGG Mut CGCTGAACACGGCG\*\*\*\*\*\*\*GTGGGGCCGCCTGG Allele-2: WT Genome sequence analysis of PCR products from parental (WT) and TMEM65 knockdown (KD) NIH3T3 cells, using sanger sequencing.