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# FLOT1 Knockdown 293T cell line, Heterozygous

Catalog No.: RM50173

#### **Basic Information**

#### Catalog No.

RM50173

#### Category

Cell Lysate

#### **Parental Cell line**

293T

#### Genotype

Knockdown

## **Background**

This gene encodes an protein that localizes to the caveolae, which are small domains on the inner cell membranes. This protein plays a role in vesicle trafficking and cell morphology. Alternative splicing results in multiple transcript variants.

#### **Gene Information**

#### **Gene Symbol**

FLOT1

#### **Species**

Human

# Gene ID

10211

#### **Swiss Prot**

075955

#### **Synonyms**

FLOT1; 1

#### **Contact**

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#### **Product Information**

#### Description

FLOT1 Knockdown cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:63bp deletion in exon4

Allele-2:64bp deletion in exon4

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°C with 5%  $CO_2$  condition.

- 1. Thaw the vial in  $37^{\circ}$ 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 GCGGCCGCCTGTCA\*\*\*\*\*\*\*\*\*\*\*CCAGAGGGCCATCA
Mut GCGGCCGCCTGTCA\*\*\*Deletion\*\*\*CCAGAGGGCCATCA
Allele-1: 63bp deletion in exon4

WT GCGGCCGCCTGTCA\*\*\*\*\*\*\*\*\*\*\*\*\*\*CAGAGGGCCATCAT
Mut GCGGCCGCCTGTCA\*\*\*Deletion\*\*\*CAGAGGGCCATCAT
Allele-2: 64bp deletion in exon4

Genome sequence analysis of PCR products from parental (WT) and FLOT1 knockdown (KD) 293T cells, using sanger sequencing.