

# NDUFAF3 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02686

## **Basic Information**

#### Catalog No.

RM02686

### Category

Cell Line

#### **Parental Cell line**

HeLa

### Genotype

Knockout

# **Background**

This gene encodes a mitochondrial complex I assembly protein that interacts with complex I subunits. Mutations in this gene cause mitochondrial complex I deficiency, a fatal neonatal disorder of the oxidative phosphorylation system. Alternatively spliced transcript variants encoding different isoforms have been identified.

## **Gene Information**

## **Gene Symbol**

NDUFAF3

#### **Species**

Human

# Gene ID

25915

### **Swiss Prot**

Q9BU61

#### **Synonyms**

2P1; E3-3; C3orf60; MC1DN18

#### **Contact**

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## **Product Information**

#### Description

NDUFAF3 Knockout cell line is engineered from HeLa cell line with Gene-Editing Technology. Allele-1:113bp deletion in exon2

Allele-2:113bp deletion in exon2

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  $37^{\circ}\text{C}$  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 GCTCTCGCCGGCGG\*\*\*\*\*\*\*\*\*\*\*CTCGGCCCCTGCGC
Mut GCTCTCGCCGGCGG\*\*\*Deletion\*\*\*CTCGGCCCTGCGC
Allele-1: 113bp deletion in exon2

WT GCTCTCGCCGGCGG\*\*\*\*\*\*\*\*\*\*\*CTCGGCCCCTGCGC
Mut GCTCTCGCCGGCGG\*\*\*Deletion\*\*\*CTCGGCCCCTGCGC
Allele-2: 113bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and NDUFAF3 knockout (KO) HeLa cells, using sanger sequencing.