

# CYCS Knockout 293T Cell Line, Homozygous

Catalog No.: RM01938

## Basic Information

**Catalog No.**

RM01938

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

CYCS

**Species**

Human

**Gene ID**

54205

**Swiss Prot**

P99999

**Synonyms**

CYC; HCS; THC4

## Contact

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## Background

This gene encodes a small heme protein that functions as a central component of the electron transport chain in mitochondria. The encoded protein associates with the inner membrane of the mitochondrion where it accepts electrons from cytochrome b and transfers them to the cytochrome oxidase complex. This protein is also involved in initiation of apoptosis. Mutations in this gene are associated with autosomal dominant nonsyndromic thrombocytopenia. Numerous processed pseudogenes of this gene are found throughout the human genome.[provided by RefSeq, Jul 2010]

## Product Information

**Description**

CYCS Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:exon1 was deleted

Allele-2:exon1 was deleted

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**

Dry ice

**Amount**

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<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

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WT TAAGTGGCTAGAGTGGTCATTCATTACACTGTATTTG  
Mut TAAGTGGCTAGAGTGGTCATTCATTACACTGTATTTG  
Allele-1: WT  
WT TGGATTGGTAATTA\*\*\*\*\*CTATCAGGAGTGTG  
Mut TGGATTGGTAATTA\*\*\*Deletion\*\*\*CTATCAGGAGTGTG  
Allele-2: exon1 was deleted

Genome sequence analysis of PCR products from parental (WT) and CYCS Knockout (KO) 293T cells, using sanger sequencing.