

CDKN1A Knockout 293T Cell Line, Homozygous

Catalog No.: RM01804

Basic Information

Catalog No.

RM01804

Category

Cell Line

Parental Cell line

293T

Genotype

Knockout

Gene Information

Gene Symbol

CDKN1A

Species

Human

Gene ID

1026

Swiss Prot

P38936

SynonymsCAP20; CDKN1; CIP1; MDA-6; P21; SDI1;
WAF1; p21CIP1

Contact

 | 400-999-6126 | cn.market@abclonal.com.cn | www.abclonal.com.cn

Background

This gene encodes a potent cyclin-dependent kinase inhibitor. The encoded protein binds to and inhibits the activity of cyclin-cyclin-dependent kinase2 or -cyclin-dependent kinase4 complexes, and thus functions as a regulator of cell cycle progression at G1. The expression of this gene is tightly controlled by the tumor suppressor protein p53, through which this protein mediates the p53-dependent cell cycle G1 phase arrest in response to a variety of stress stimuli. This protein can interact with proliferating cell nuclear antigen, a DNA polymerase accessory factor, and plays a regulatory role in S phase DNA replication and DNA damage repair. This protein was reported to be specifically cleaved by CASP3-like caspases, which thus leads to a dramatic activation of cyclin-dependent kinase2, and may be instrumental in the execution of apoptosis following caspase activation. Mice that lack this gene have the ability to regenerate damaged or missing tissue. Multiple alternatively spliced variants have been found for this gene.

Product Information

Description

CDKN1A Knockout cell line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:1bp insertion and 51bp deletion in exon1

Allele-2:50bp deletion in exon1

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

Amount1~5x10⁶ 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% CO₂ 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% CO₂.
7. A subcultivation ratio of 1:2-1:4 is recommended.

Sequencing data

WT GGAGGCCCGTG*****CCTGGGAGCGT
Mut GGAGGCCCGTGC***Deletion(51bp)***CCTGGGAGCGT
Allele-1: 1bp Insertion and 51bp deletion in exon1
WT GGAGGCCCGTGA*****CCTGGGAGCGT
Mut GGAGGCCCGTGA***Deletion(50bp)***CCTGGGAGCGT
Allele-2: 50bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and CDKN1A knockout (KO) 293T cells, using sanger sequencing.