

# PPP2CA Knockout 293T Cell Line, Homozygous

Catalog No.: RM02134

## Basic Information

**Catalog No.**

RM02134

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

PPP2CA

**Species**

Human

**Gene ID**

5515

**Swiss Prot**

P67775

**Synonyms**

PP2Ac; PP2CA; PP2Calpha; RP-C

## Contact

 | 400-999-6126 | [cn.market@abclonal.com.cn](mailto:cn.market@abclonal.com.cn) | [www.abclonal.com.cn](http://www.abclonal.com.cn)

## Background

This gene encodes the phosphatase 2A catalytic subunit. Protein phosphatase 2A is one of the four major Ser/Thr phosphatases, and it is implicated in the negative control of cell growth and division. It consists of a common heteromeric core enzyme, which is composed of a catalytic subunit and a constant regulatory subunit, that associates with a variety of regulatory subunits. This gene encodes an alpha isoform of the catalytic subunit. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

PPP2CA Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:130bp deletion in exon2

Allele-2:130bp 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**

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

---

WT AAAAGAATCCAACG\*\*\*\*\*CAGAGGATATTATT  
Mut AAAAGAATCCAACG\*\*\*Deletion\*\*\*CAGAGGATATTATT  
Allele-1: 130bp deletion in exon2  
WT AAAAGAATCCAACG\*\*\*\*\*CAGAGGATATTATT  
Mut AAAAGAATCCAACG\*\*\*Deletion\*\*\*CAGAGGATATTATT  
Allele-2: 130bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and PPP2CA knockout (KO) 293T cells, using sanger sequencing.