

# CETN2 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02130

## Basic Information

**Catalog No.**

RM02130

**Category**

Cell Line

**Parental Cell line**

HeLa

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

CETN2

**Species**

Human

**Gene ID**

1069

**Swiss Prot**

P41208

**Synonyms**

CALT; CEN2

## 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

Caltractin belongs to a family of calcium-binding proteins and is a structural component of the centrosome. The high level of conservation from algae to humans and its association with the centrosome suggested that caltractin plays a fundamental role in the structure and function of the microtubule-organizing center, possibly required for the proper duplication and segregation of the centrosome. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

CETN2 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:80bp deletion in exon1

Allele-2:80bp 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

**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 AGAATGAGCCCTAA\*\*\*\*\*ATGTTAAGAAGCTG  
Mut AGAATGAGCCCTAA\*\*\*Deletion\*\*ATGTTAAGAAGCTG  
Allele-1: 80bp deletion in exon1  
WT AGAATGAGCCCTAA\*\*\*\*\*ATGTTAAGAAGCTG  
Mut AGAATGAGCCCTAA\*\*\*Deletion\*\*\*ATGTTAAGAAGCTG  
Allele-2: 80bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and CETN2 knockout (KO) HeLa cells, using sanger sequencing.