# **CTCF Knockout 293T Cell Line, Homozygous**

Catalog No.: RM01849



## **Basic Information**

Catalog No. RM01849

Category Cell Line

Parental Cell line 293T

Genotype Knockout

## **Gene Information**

Gene Symbol CTCF

Species Human

**Gene ID** 10664

Swiss Prot P49711

Synonyms MRD21

## Contact

6	400-999-6126
$\times$	cn.market@abclonal.com.cn
€	www.abclonal.com.cn

## Background

This gene is a member of the BORIS + CTCF gene family and encodes a transcriptional regulator protein with 11 highly conserved zinc finger (ZF) domains. This nuclear protein is able to use different combinations of the ZF domains to bind different DNA target sequences and proteins. Depending upon the context of the site, the protein can bind a histone acetyltransferase (HAT)-containing complex and function as a transcriptional activator or bind a histone deacetylase (HDAC)-containing complex and function as a transcriptional repressor. If the protein is bound to a transcriptional insulator element, it can block communication between enhancers and upstream promoters, thereby regulating imprinted expression. Mutations in this gene have been associated with invasive breast cancers, prostate cancers, and Wilms' tumors. Alternatively spliced transcript variants encoding different isoforms have been found for this gene.

## **Product Information**

#### Description

CTCF Knockout 293T cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:exon2 was deleted Allele-2:exon2 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  $37^{\circ}$ 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%  $\mbox{CO}_2.$
- 7. A subcultivation ratio of 1:2-1:4 is recommended.

## Sequencing data

WT AGTGCCCCAAAG\*\*\*\*\*Deletion(358bp)\*\*\*AATTCGTTGTT Mut AGTGCCCCAAAG\*\*\*Deletion(358bp)\*\*\*AATTCGTTGTT Allele-1: exon2 was deleted

WT TCGTAFTTCAG\*\*\*\*\*\*\*\*GTAGGACTTCT Mut TCGTAFTTCAG\*\*\*\*\*\*\*\*\*\*\*\*GTAGGACTTCT Allele-2: WT Genome sequence analysis of PCR products from parental (WT) and CTCF knockdown (KD) 293T cells, using sanger sequencing.