

# RHOC Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02239

## Basic Information

**Catalog No.**

RM02239

**Category**

Cell Line

**Parental Cell line**

HeLa

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

RHOC

**Species**

Human

**Gene ID**

389

**Swiss Prot**

P08134

**Synonyms**

ARH9; ARHC; H9; RHOH9

## Contact

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## Background

This gene encodes a member of the Rho family of small GTPases, which cycle between inactive GDP-bound and active GTP-bound states and function as molecular switches in signal transduction cascades. Rho proteins promote reorganization of the actin cytoskeleton and regulate cell shape, attachment, and motility. The protein encoded by this gene is prenylated at its C-terminus, and localizes to the cytoplasm and plasma membrane. It is thought to be important in cell locomotion. Overexpression of this gene is associated with tumor cell proliferation and metastasis. Multiple alternatively spliced variants, encoding the same protein, have been identified. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

RHOC Knockout cell line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:70bp deletion in exon1

Allele-2:70bp 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 AAGACCTGCCTCCT\*\*\*\*\*ATTGAGGTGGACGG  
Mut AAGACCTGCCTCCT\*\*\*Deletion\*\*\*ATTGAGGTGGACGG  
Allele-1: 70bp deletion in exon1  
WT AAGACCTGCCTCCT\*\*\*\*\*ATTGAGGTGGACGG  
Mut AAGACCTGCCTCCT\*\*\*Deletion\*\*\*ATTGAGGTGGACGG  
Allele-2: 70bp deletion in exon1

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