

# ARHGDI<sup>A</sup> Knockout 293T Cell Line, Homozygous

Catalog No.: RM02227

## Basic Information

**Catalog No.**

RM02227

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**ARHGDI<sup>A</sup>**Species**

Human

**Gene ID**

396

**Swiss Prot**

P52565

**Synonyms**GDIA1; HEL-S-47e; NPHS8; RHOGDI;  
RHOGDI-1

## Contact

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## Background

This gene encodes a protein that plays a key role in the regulation of signaling through Rho GTPases. The encoded protein inhibits the disassociation of Rho family members from GDP (guanine diphosphate), thereby maintaining these factors in an inactive state. Activity of this protein is important in a variety of cellular processes, and expression of this gene may be altered in tumors. Mutations in this gene have been found in individuals with nephrotic syndrome, type 8. Alternate splicing results in multiple transcript variants. [provided by RefSeq, Jul 2014]

## Product Information

**Description**

ARHGDI<sup>A</sup> Knockout cell line is engineered from 293T cell line with Gene-Editing Technology.  
Allele-1:91bp deletion in exon1  
Allele-2:91bp 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 TGCAGCGGAGAACG\*\*\*\*\*AAAGTACAAGGAGG  
Mut TGCAGCGGAGAACG\*\*\*Deletion\*\*\*AAAGTACAAGGAGG  
Allele-1: 91bp deletion in exon1  
WT TGCAGCGGAGAACG\*\*\*\*\*AAAGTACAAGGAGG  
Mut TGCAGCGGAGAACG\*\*\*Deletion\*\*\*AAAGTACAAGGAGG  
Allele-2: 91bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and ARHGDI1 knockout (KO) 293T cells, using sanger sequencing.