

# AKT1S1 Knockdown HeLa Cell Line, Heterozygous

Catalog No.: RM01918

## Basic Information

### Catalog No.

RM01918

### Category

Cell Line

### Parental Cell line

HeLa

### Genotype

Knockdown

## Gene Information

### Gene Symbol

AKT1S1

### Species

Human

### Gene ID

84335

### Swiss Prot

Q96B36

### Synonyms

Lobe; PRAS40

## Contact

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## Background

AKT1S1 is a proline-rich substrate of AKT (MIM 164730) that binds 14-3-3 protein (see YWHAH, MIM 113508) when phosphorylated (Kovacina et al., 2003 [PubMed 12524439]).[supplied by OMIM, Mar 2008]

## Product Information

### Description

AKT1S1 Knockdown HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:47bp deletion in exon2

Allele-2:48bp 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

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WT CTATGCTGCCCATG\*\*\*\*\*GCACTGGCCCACAG  
Mut CTATGCTGCCCATG\*\*\*Deletion\*\*\*GCACTGGCCCACAG  
Allele-1: 47bp deletion in exon2  
WT CTATGCTGCCCATG\*\*\*\*\*CACTGGCCCACAGG  
Mut CTATGCTGCCCATG\*\*\*Deletion\*\*\*CACTGGCCCACAGG  
Allele-2: 48bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and AKT1S1 Knockdown (KD) HeLa cells, using sanger sequencing.