# VHL Knockout 293T Cell Line, Homozygous

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Catalog No.: RM50098

# **Basic Information**

#### Catalog No.

RM50098

#### Category

Cell Line

#### **Parental Cell line**

293T

#### Genotype

Knockout

# **Background**

This gene encodes a component of a ubiquitination complex. The encoded protein is involved in the ubiquitination and degradation of hypoxia-inducible-factor (HIF), which is a transcription factor that plays a central role in the regulation of gene expression by oxygen. In addition to oxygen-related gene expression, this protein plays a role in many other cellular processes including cilia formation, cytokine signaling, regulation of senescence, and formation of the extracellular matrix. Variants of this gene are associated with von Hippel-Lindau syndrome, pheochromocytoma, erythrocytosis, renal cell carcinoma, and cerebellar hemangioblastoma.

# **Gene Information**

## **Gene Symbol**

VHL

#### **Species**

Human

# Gene ID

7428

# **Swiss Prot**

P40337

#### **Synonyms**

RCA1; VHL1; pVHL; HRCA1; VHL

#### **Contact**

<u>a</u>	400-999-6126
$\bowtie$	cn.market@abclonal.com.cn
•	www.abclonal.com.cn

# **Product Information**

#### Description

VHL Knockout cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:149bp deletion in exon1

Allele-2:28bp 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**

**Amount** 

Dry ice

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%  $CO_2$  condition.

- 1. Thaw the vial in  $37^{\circ}\text{C}$  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

WT AGTCCGGCCCGGAA\*\*\*\*\*\*\*\*\*\*CGACGGCGAGCCGC
Mut AGTCCGGCCCGGAA\*\*\*Deletion\*\*\*CGACGGCGAGCCGC
Allele-1: 149bp deletion in exon1

WT CGGA\*\*\*\*\*\*\*\*GGAA\*\*CCGT\*\*\*\*\*\*\*GACG
Mut CGGA\*\*Deletion\*\*\*GGAA\*\*CCGT\*\*Deletion\*\*\*GACG
Allele-2: 28bp deletion in exon1

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