

# PAX3 Knockdown 293T Cell Line, Heterozygous

Catalog No.: RM01900

## Basic Information

**Catalog No.**

RM01900

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockdown

## Gene Information

**Gene Symbol**

PAX3

**Species**

Human

**Gene ID**

5077

**Swiss Prot**

P23760

**Synonyms**

CDHS; HUP2; WS1; WS3

## Contact

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## Background

This gene is a member of the paired box (PAX) family of transcription factors. Members of the PAX family typically contain a paired box domain and a paired-type homeodomain. These genes play critical roles during fetal development. Mutations in paired box gene 3 are associated with Waardenburg syndrome, craniofacial-deafness-hand syndrome, and alveolar rhabdomyosarcoma. The translocation t(2;13)(q35;q14), which represents a fusion between PAX3 and the forkhead gene, is a frequent finding in alveolar rhabdomyosarcoma. Alternative splicing results in transcripts encoding isoforms with different C-termini. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

PAX3 Knockdown 293T Cell Line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:168bp deletion in exon2

Allele-2:169bp 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 CGCGTCAACCAGCT\*\*\*\*\*TCCTGGTGCCATCG  
Mut CGCGTCAACCAGCT\*\*\*Deletion\*\*\*TCCTGGTGCCATCG  
Allele-1: 168bp deletion in exon2  
WT CGCGTCAACCAGCT\*\*\*\*\*CCTGGTGCCATCGG  
Mut CGCGTCAACCAGCT\*\*\*Deletion\*\*\*CCTGGTGCCATCGG  
Allele-2: 169bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and PAX3 Knockdown (KD) 293T cells, using sanger sequencing.