

# JUN Knockdown HeLa Cell Line, Heterozygous

**Catalog No.:** RM02092

## Basic Information

**Catalog No.**

RM02092

**Category**

Cell Line

**Parental Cell line**

HeLa

**Genotype**

Knockdown

## Gene Information

**Gene Symbol**

JUN

**Species**

Human

**Gene ID**

3725

**Swiss Prot**

P05412

**Synonyms**

AP-1; AP1; c-Jun

## Contact

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## Background

This gene is the putative transforming gene of avian sarcoma virus 17. It encodes a protein which is highly similar to the viral protein, and which interacts directly with specific target DNA sequences to regulate gene expression. This gene is intronless and is mapped to 1p32-p31, a chromosomal region involved in both translocations and deletions in human malignancies. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

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

Allele-1:98bp deletion in exon1

Allele-2:99bp 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 CGTTCCTCCCGTCC\*\*\*\*\*CGCCAAGAACTCGG  
Mut CGTTCCTCCCGTCC\*\*\*Deletion\*\*\*CGCCAAGAACTCGG  
Allele-1: 98bp deletion in exon1  
WT CGTTCCTCCCGTCC\*\*\*\*\*GCCAAGAACTCGGA  
Mut CGTTCCTCCCGTCC\*\*\*Deletion\*\*\*GCCAAGAACTCGGA  
Allele-2: 99bp deletion in exon1

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