

# IL1B Knockout A549 Cell Line, Homozygous

Catalog No.: RM02641

## Basic Information

**Catalog No.**

RM02641

**Category**

Cell Line

**Parental Cell line**

A549

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

IL1B

**Species**

Human

**Gene ID**

3553

**Swiss Prot**

P01584

**Synonyms**

IL-1; IL1-BETA; IL1F2

## Contact

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## Background

The protein encoded by this gene is a member of the interleukin 1 cytokine family. This cytokine is produced by activated macrophages as a proprotein, which is proteolytically processed to its active form by caspase 1 (CASP1/ICE). This cytokine is an important mediator of the inflammatory response, and is involved in a variety of cellular activities, including cell proliferation, differentiation, and apoptosis. The induction of cyclooxygenase-2 (PTGS2/COX2) by this cytokine in the central nervous system (CNS) is found to contribute to inflammatory pain hypersensitivity. This gene and eight other interleukin 1 family genes form a cytokine gene cluster on chromosome 2. [provided by RefSeq, Jul 2008]

## Product Information

**Description**

IL1B Knockout A549 Cell Line is engineered from A549 cell line with Gene-Editing technology.

Allele-1:59bp deletion in exon3

Allele-2:59bp deletion in exon3

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 GGCGGCATCCAGCT\*\*\*\*\*CCATGGACAAGCT  
Mut GGCGGCATCCAGCT\*\*\*Deletion\*\*\*CCATGGACAAGCT  
Allele-1: 59bp deletion in exon3

Genome sequence analysis of PCR products from parental (WT) and IL1B knockout (KO) A549 cells, using sanger sequencing.

WT GGCGGCATCCAGCT\*\*\*\*\*CCATGGACAAGCT  
Mut GGCGGCATCCAGCT\*\*\*Deletion\*\*\*CCATGGACAAGCT  
Allele-2: 59bp deletion in exon3