

# IL6 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM01829

## Basic Information

**Catalog No.**

RM01829

**Category**

Cell Line

**Parental Cell line**

HeLa

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

IL6

**Species**

Human

**Gene ID**


3569

**Swiss Prot**

P05231

**Synonyms**BSF-2; BSF2; CDF; HGF; HSF; IFN-beta-2;  
IFNB2; IL-6

## Contact

 | 400-999-6126 | [cn.market@abclonal.com.cn](mailto:cn.market@abclonal.com.cn) | [www.abclonal.com.cn](http://www.abclonal.com.cn)

## Background

This gene encodes a cytokine that functions in inflammation and the maturation of B cells. In addition, the encoded protein has been shown to be an endogenous pyrogen capable of inducing fever in people with autoimmune diseases or infections. The protein is primarily produced at sites of acute and chronic inflammation, where it is secreted into the serum and induces a transcriptional inflammatory response through interleukin 6 receptor, alpha. The functioning of this gene is implicated in a wide variety of inflammation-associated disease states, including susceptibility to diabetes mellitus and systemic juvenile rheumatoid arthritis. Alternative splicing results in multiple transcript variants.

## Product Information

**Description**

IL6 Knockout cell line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:10bp deletion in exon1

Allele-2:2bp insertion 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 CTGGCAGAAAACAA\*\*\*\*\*CCAAAGATGGCTGA  
Mut CTGGCAGAAAACAA\*\*\*Deletion\*\*\*CCAAAGATGGCTGA  
Allele-1: 10bp deletion in exon1

WT CTGGCAGAAAACAACCTGAAC- -CTTCAAAGATGGCTGA  
Mut CTGGCAGAAAACAACCTGAACATCTTCAAAGATGGCTGA  
Allele-2: 2bp insertion in exon1

Genome sequence analysis of PCR products from parental (WT) and IL6 knockout (KO) HeLa cells, using sanger sequencing.