Leader in Biomolecular Solutions for Life Science

www.abclonal.com

# IL6R Knockout HCT116 Cell Line, Homozygous

**Catalog No.: RM01899**

## Basic Information

**Catalog No.**
RM01899

**Category**
Cell Line

**Parental Cell line**
HCT116

**Genotype**
Knockout

## Background

This gene encodes a subunit of the interleukin 6 (IL6) receptor complex. Interleukin 6 is a potent pleiotropic cytokine that regulates cell growth and differentiation and plays an important role in the immune response. The IL6 receptor is a protein complex consisting of this protein and interleukin 6 signal transducer (IL6ST/GP130/IL6-beta), a receptor subunit also shared by many other cytokines. Dysregulated production of IL6 and this receptor are implicated in the pathogenesis of many diseases, such as multiple myeloma, autoimmune diseases and prostate cancer. Alternatively spliced transcript variants encoding distinct isoforms have been reported. A pseudogene of this gene is found on chromosome 9.[provided by RefSeq, May 2011]

## Gene Information

**Gene Symbol**
IL6R

**Species**
Human

**Gene ID**
3570

**Swiss Prot**
P08887

**Synonyms**
CD126; IL-6R-1; IL-6RA; IL6Q; IL6RA; IL6RQ; gp80

## Product Information

**Description**
IL6R Knockout HCT116 Cell Line is engineered from HCT116 cell line with Gene-Editing Technology.
Allele-1:151bp deletion in exon2
Allele-2:151bp 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**
1~5x$10^6$ cells/vial

**Storage**
Stored in liquid nitrogen for a long time less than -130℃. 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℃ with 5% $CO_2$ condition.
1. Thaw the vial in 37℃ 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℃ incubator with 5% $CO_2$.
7. A subcultivation ratio of 1:2-1:4 is recommended.

## Contact


400-999-6126

cn.market@abclonal.com.cn

www.abclonal.com.cn

## Sequencing data

WT GACTCTGACCTGCC**********GGCCGGCCGCCCAG
Mut GACTCTGACCTGCC***Deletion***GGCCGGCCGCCCAG
Allele-1: 151bp deletion in exon2

WT GACTCTGACCTGCC**********GGCCGGCCGCCCAG
Mut GACTCTGACCTGCC***Deletion***GGCCGGCCGCCCAG
Allele-2: 151bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and IL6R knockout (KO) HCT116 cells, using sanger sequencing.