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IFNAR2 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM01885

Basic Information

Catalog No.
RM01885

Category
Cell Line

Parental Cell line
HeLa

Genotype
Knockout

Gene Information

Gene Symbol
IFNAR2

Species
Human

Gene ID
3455

Swiss Prot
P48551

Synonyms
IFN-R; IFN-alpha-REC; IFNABR; IFNARB; IMD45

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Background

The protein encoded by this gene is a type I membrane protein that forms one of the two chains of a receptor for interferons alpha and beta. Binding and activation of the receptor stimulates Janus protein kinases, which in turn phosphorylate several proteins, including STAT1 and STAT2. Multiple transcript variants encoding at least two different isoforms have been found for this gene. [provided by RefSeq, Jul 2008]

Product Information

Description
IFNAR2 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.
Allele-1:2bp deletion in exon3
Allele-2:1bp 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
1~5x10^6 cells/vial

Storage
Store at -20℃

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.

Sequencing data

WT TTAAAAAACCACTCCATTGTACCAACTCACTATACATT
Mut TTAAAAAACCACTCCATT- -ACCAACTCACTATACATT
Allele-1: 2bp deletion in exon3

WT TAAAAAACCACTCCATTGTACCAACTCACTATACATTG
Mut TAAAAAACCACTCCATTGT- CCAACTCACTATACATTG
Allele-2: 1bp deletion in exon3

Genome sequence analysis of PCR products from parental (WT) and IFNAR2 knockout (KO) HeLa cells, using sanger sequencing.

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