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NDUFB6 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02669

Basic Information

Catalog No.
RM02669

Category
Cell Line

Parental Cell line
HeLa

Genotype
Knockout

Background

The protein encoded by this gene is a subunit of the multisubunit NADH:ubiquinone oxidoreductase (complex I). Mammalian complex I is composed of 45 different subunits. It locates at the mitochondrial inner membrane. This protein has NADH dehydrogenase activity and oxidoreductase activity. It transfers electrons from NADH to the respiratory chain. The immediate electron acceptor for the enzyme is believed to be ubiquinone. Alternative splicing occurs at this locus and three transcript variants encoding distinct isoforms have been identified.

Gene Information

Gene Symbol
NDUFB6

Species
Human

Gene ID
4712

Swiss Prot
O95139

Synonyms
CI; B17

Product Information

Description
NDUFB6 Knockout cell line is engineered from HeLa cell line with Gene-Editing Technology.
Allele-1:exon1 was deleted
Allele-2:exon1 was deleted
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.

Contact

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Sequencing data

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WT    GTCCTTTTGCGTTG*************CGCTCAGTGCTCTG
Mut   GTCCTTTTGCGTTG***Deletion***CGCTCAGTGCTCTG
Allele-1:    exon1 was deleted

WT    CGTTGGTACCAGCG*************CAAGGACTGATAA
Mut   CGTTGGTACCAGCG***Deletion***CAAGGACTGATAA
Allele-2:    exon1 was deleted
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Genome sequence analysis of PCR products from parental (WT) and NDUFB6 knockout (KO) HeLa cells, using sanger sequencing.

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