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EXON2 Knockout 293T Cell Line, Homozygous

Catalog No.: RM02146

Basic Information

Catalog No.

RM02146

Category

Cell Line

Parental Cell line

293T

Genotype

Knockout

Background

This gene is a tumor-specific member of the ECTO-NOX family of genes that encode cell surface NADH oxidases. The encoded protein has two enzymatic activities: catalysis of hydroquinone or NADH oxidation, and protein disulfide interchange. The protein also displays prion-like properties. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Aug 2013]

Gene Information

Gene Symbol

ENOX2

Species

Human

Gene ID

10495

Swiss Prot

Q16206

Synonyms

APK1; COVA1; tNOX

Contact

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Product Information

Description

EXON2 Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:143bp deletion in exon4

Allele-2:143bp deletion in exon4

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

 ${\bf 1}$ vial parental cell line and ${\bf 1}$ vial knockout cell line

Shipping Conditions

Amount

Dry ice

1~5x10⁶ cells/vial

Storage

Stored in liquid nitrogen for a long time less than -130°C. Minimizing freeze-thaw cycles.

Protoco

Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at $37^{\circ}C$ with 5% CO₂ 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% CO₂.
- 7. A subcultivation ratio of 1:2-1:4 is recommended.

Sequencing data

WT TCCCACCTCCTGCA**********************TCGCTTTGCTGAGG
Mut TCCCACCTCCTGCA***Deletion****TCGCTTTGCTGAGG
Allele-1: 143bp deletion in exon4

WT TCCCACCTCCTGCA**********TCGCTTTGCTGAGG
Mut TCCCACCTCCTGCA***Deletion***TCGCTTTGCTGAGG

Allele-2: 143bp deletion in exon4

Genome sequence analysis of PCR products from parental (WT) and ENOX2 knockout (KO) 293T cells, using sanger sequencing.