

P53 Knockout 293T Cell Line, Homozygous

Catalog No.: RM01815

Basic Information

Catalog No.

RM01815

Category

Cell Line

Parental Cell line

293T

Genotype

Knockout

Gene Information

Gene Symbol

TP53

Species

Human

Gene ID

7157

Swiss Prot

P04637

Synonyms

BCC7; LFS1; P53; TRP53

Contact

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Background

This gene encodes a tumor suppressor protein containing transcriptional activation, DNA binding, and oligomerization domains. The encoded protein responds to diverse cellular stresses to regulate expression of target genes, thereby inducing cell cycle arrest, apoptosis, senescence, DNA repair, or changes in metabolism. Mutations in this gene are associated with a variety of human cancers, including hereditary cancers such as Li-Fraumeni syndrome. Alternative splicing of this gene and the use of alternate promoters result in multiple transcript variants and isoforms. Additional isoforms have also been shown to result from the use of alternate translation initiation codons from identical transcript variants (PMIDs: 12032546, 20937277).

Product Information

Description

TP53 Knockout cell line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:exon2 was destroyed

Allele-2:exon2 was destroyed

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

Amount1~5x10⁶ cells/vial.**Storage**

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

ProtocolUpon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at 37°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.

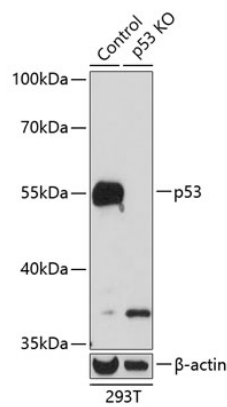
Sequencing data

WT GTGAGCAGCTGGGG*****AAGGAAATTTGCGT
Mut GTGAGCAGCTGGGG***Deletion***AAGGAAATTTGCGT
Allele-1: exon2 was destroyed

WT GTGAGCAGCTGGGG*****AAGGAAATTTGCGT
Mut GTGAGCAGCTGGGG***Deletion***AAGGAAATTTGCGT
Allele-2: exon2 was destroyed

Genome sequence analysis of PCR products from parental (WT) and TP53 knockout (KO) 293T cells, using sanger sequencing.

WB data



Western blot analysis of extracts from parental (Control) and TP53 knockout (KO) 293T cells, using TP53 antibody at 1:1000 dilution.