

SND1 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM02132

Basic Information

Catalog No.

RM02132

Category

Cell Line

Parental Cell line

HeLa

Genotype

Knockout

Gene Information

Gene Symbol

SND1

Species

Human

Gene ID

27044

Swiss Prot

Q7KZF4

Synonyms

TDRD11; Tudor-SN; p100

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Background

This gene encodes a transcriptional co-activator that interacts with the acidic domain of Epstein-Barr virus nuclear antigen 2 (EBNA 2), a transcriptional activator that is required for B-lymphocyte transformation. Other transcription factors that interact with this protein are signal transducers and activators of transcription, STATs. This protein is also thought to be essential for normal cell growth. A similar protein in mammals and other organisms is a component of the RNA-induced silencing complex (RISC). [provided by RefSeq, Jul 2016]

Product Information

Description

SND1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:76bp deletion in exon2

Allele-2:76bp 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

1~5x10⁶ cells/vial

Storage

Stored in liquid nitrogen for a long time less than -130°C. 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°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 GCGCCATCATTGTC*****GCCGGGCAGCCGCC
Mut GCGCCATCATTGTC***Deletion***GCCGGGCAGCCGCC
Allele-1: 76bp deletion in exon2
WT GCGCCATCATTGTC*****GCCGGGCAGCCGCC
Mut GCGCCATCATTGTC***Deletion***GCCGGGCAGCCGCC
Allele-2: 76bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and SND1 knockout (KO) HeLa cells, using sanger sequencing.