

MYSM1 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM50051

Basic Information

Catalog No.

RM50051

Category

Cell Line

Parental Cell line

HeLa

Genotype

Knockout

Gene Information

Gene Symbol

MYSM1

Species

Human

Gene ID

114803

Swiss Prot

Q5VVJ2

Synonyms

2ADUB; BMFS4; 2A-DUB; MYSM1

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Background

Enables histone binding activity; peptidase activity; and transcription coactivator activity. Involved in several processes, including chromatin remodeling; monoubiquitinated histone H2A deubiquitination; and positive regulation of transcription by RNA polymerase II. Located in nucleolus and nucleoplasm. Part of protein-containing complex. Implicated in diabetic retinopathy.

Product Information

Description

MYSM1 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

1~5x10⁶ 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 TCGGGA GGAC*****TCCGGCAAGGGAAA
Mut TCGGGA**TCTT**GGAC***Deletion***TCCGGCAAGGGAAA
Allele-1: exon1 was deleted

WT TCTCGGGAGGACGG*****GCGGGCCCTGGGAA
Mut TCTCGGGAGGACGG***Deletion***GCGGGCCCTGGGAA
Allele-2: exon1 was deleted

Genome sequence analysis of PCR products from parental (WT) and MYSM1 knockout (KO) HeLa cells, using sanger sequencing.