

MAP1LC3B Knockdown 293T Cell Lysate, Heterozygous

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Catalog No.: RM50064

Basic Information

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RM50064

Category

Cell Lysate

Parental Cell line

293T

Genotype

Knockdown

Gene Information

Gene Symbol

MAP1LC3B

Species

Human

Gene ID

81631

Swiss Prot

Q9GZQ8

Synonyms

ATG8F; LC3B; MAP1A/1BLC3; MAP1LC3B-a

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Background

The product of this gene is a subunit of neuronal microtubule-associated MAP1A and MAP1B proteins, which are involved in microtubule assembly and important for neurogenesis. Studies on the rat homolog implicate a role for this gene in autophagy, a process that involves the bulk degradation of cytoplasmic component. [provided by RefSeq, Jul 2008]

Product Information

Description

MAP1LC3B Knockdown 293T Cell Lysate is engineered from 293T cell line with Gene-Editing technology.

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 CTGGCCCTGAT*****ATTAGGTATTC
Mut CTGGCCCTGAT***Deletion(137bp)***ATTAGGTATTC
Allele-1: 102 bp deletion in exon3

WT CTGGCCCTGAT*****AATTAGGTATTC
Mut CTGGCCCTGAT***Deletion(136bp)***AATTAGGTATTC
Allele-2: 101 bp deletion in exon3

Genome sequence analysis of PCR products from parental (WT) and MAP1LC3B Knockdown (KD) 293T cells, using sanger sequencing.