# MAP1LC3B Knockout 293T Cell Lysate, Homozygous

Catalog No.: RM50133



## **Basic Information**

Catalog No. RM50133

Category Cell Lysate

Parental Cell line 293T

Genotype Knockout

## **Gene Information**

Gene Symbol MAP1LC3B

Species Human

Gene ID 81631

Swiss Prot Q9GZQ8

Synonyms LC3B; ATG8F; MAP1LC3B-a; MAP1A/1BLC3; 3B

### Contact

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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.

## **Product Information**

#### Description

MAP1LC3B Knockout cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:exon2 was deleted Allele-2:exon2 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 Lysate and 1 vial knockout cell Lysate

## **Shipping Conditions** 4°C

**Amount** 50μL, 2μg/μL.

#### Storage

Lysate is stable for 12 months when stored at -20°C. Minimizing freeze-thaw cycles.

#### Protocol

To be used as WB control. Lysate is supplied in  $1 \times$  SDS sample buffer (2% SDS, 60 mM Tris-HCl pH 6.8, 10% Glycerol, 0.02% Bromophenol blue, 60 mM beta-mercaptoethanol). Lysate should be boiled for 3 - 5 minutes before loading onto gel.

## Sequencing data

WT TGTGCCACAGC\*\*\*\*Deletion(189bp)\*\*\*AGAGGAGAGAGA Mut TGTGCCACAGC\*\*\*Deletion(189bp)\*\*\*AGAGGAGAGAGA Allele-1: exon2 was deleted

WT TCTGCTGTGCC\*\*\*\*\*\*\*\*\*\*\*\*\*\*\*CAGAGGAGAGC Mut TCTGCTGTGCC\*\*\*Deletion(193bp)\*\*CAGAGGAGAGC Allele-2: exon2 was deleted Genome sequence analysis of PCR products from parental (WT) and MAP1LC3B knockout (KO) 293T cells, using sanger sequencing.