

# LDHB Knockout 293T Cell Line, Homozygous

**Catalog No.:** RM02737

## Basic Information

**Catalog No.**

RM02737

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

LDHB

**Species**

Human

**Gene ID**

3945

**Swiss Prot**

P07195

**Synonyms**

LDH-B; LDH-H; LDHBD; TRG-5; HEL-S-281; LDHB

## Contact

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## Background

This gene encodes the B subunit of lactate dehydrogenase enzyme, which catalyzes the interconversion of pyruvate and lactate with concomitant interconversion of NADH and NAD<sup>+</sup> in a post-glycolysis process. Alternatively spliced transcript variants have been found for this gene. Recent studies have shown that a C-terminally extended isoform is produced by use of an alternative in-frame translation termination codon via a stop codon readthrough mechanism, and that this isoform is localized in the peroxisomes. Mutations in this gene are associated with lactate dehydrogenase B deficiency. Pseudogenes have been identified on chromosomes X, 5 and 13.

## Product Information

**Description**

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

Allele-1:134bp deletion in exon3

Allele-2:134bp deletion in exon3

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**

1~5x10<sup>6</sup> 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% CO<sub>2</sub> 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% CO<sub>2</sub>.
7. A subcultivation ratio of 1:2-1:4 is recommended.

## Sequencing data

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WT CAGTGTAGCTCAAG\*\*\*\*\*TACAGTCCTGATTG  
Mut CAGTGTAGCTCAAG\*\*\*Deletion\*\*\*TACAGTCCTGATTG  
Allele-1: 134bp deletion in exon3

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

WT CAGTGTAGCTCAAG\*\*\*\*\*TACAGTCCTGATTG  
Mut CAGTGTAGCTCAAG\*\*\*Deletion\*\*\*TACAGTCCTGATTG  
Allele-2: 134bp deletion in exon3