# ABclonal® www.abclonal.com

# HK1 Knockout 293T Cell Line, Homozygous

Catalog No.: RM01822

## **Basic Information**

#### Catalog No.

RM01822

#### Category

Cell Line

#### **Parental Cell line**

293T

#### Genotype

Knockout

## **Background**

Hexokinases phosphorylate glucose to produce glucose-6-phosphate, the first step in most glucose metabolism pathways. This gene encodes a ubiquitous form of hexokinase which localizes to the outer membrane of mitochondria. Mutations in this gene have been associated with hemolytic anemia due to hexokinase deficiency. Alternative splicing of this gene results in several transcript variants which encode different isoforms, some of which are tissue-specific.

#### **Gene Information**

#### **Gene Symbol**

HK1

# Species

Human

# Gene ID

3098

### **Swiss Prot**

P19367

#### **Synonyms**

HK; HK1-ta; HK1-tb; HK1-tc; HKD; HKI; HMSNR; HXK1; hexokinase

## **Contact**

<u>a</u>	400-999-6126
$\bowtie$	cn.market@abclonal.com.cn
•	www.abclonal.com.cn

#### **Product Information**

#### Description

HK1 Knockout cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:4bp deletion in exon2

Allele-2:4bp 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**

Amount

Dry ice

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  $37^{\circ}$ C with 5% CO<sub>2</sub> condition.

- 1. Thaw the vial in  $37^{\circ}$ 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

TCATAGATATCATGACTCGCTTCAGGAAGGAGATGAAGAA Mut TCATAGATATCATGACTCGC----GGAAGGAGATGAAGAA Allele-1: 4bp deletion in exon2

WT TCATAGATATCATGACTCGCTTCAGGAAGGAGATGAAGAA Mut TCATAGATATCATGACTCGC----GGAAGGAGATGAAGAA

Allele-2: 4bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and HK1 knockout (KO) 293T cells, using sanger sequencing.