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# **TEAD1** Knockout HeLa Cell Line, Homozygous

Catalog No.: RM01923

#### **Basic Information**

#### Catalog No.

RM01923

#### Category

Cell Line

#### **Parental Cell line**

HeLa

#### Genotype

Knockout

## **Background**

This gene encodes a ubiquitous transcriptional enhancer factor that is a member of the TEA/ATTS domain family. This protein directs the transactivation of a wide variety of genes and, in placental cells, also acts as a transcriptional repressor. Mutations in this gene cause Sveinsson's chorioretinal atrophy. Additional transcript variants have been described but their full-length natures have not been experimentally verified. [provided by RefSeq, May 2010]

#### **Gene Information**

#### **Gene Symbol**

TEAD1

#### **Species**

Human

## Gene ID

7003

#### **Swiss Prot**

P28347

#### **Synonyms**

AA; NTEF-1; REF1; TCF-13; TCF13; TEAD-1; TEF-1

### **Contact**

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#### **Product Information**

#### Description

TEAD1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.

Allele-1:23bp insertion and 118bp deletion in exon1

Allele-2:70bp deletion in exon1

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**

 ${\bf 1}$  vial parental cell line and  ${\bf 1}$  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.

#### **Protoco**

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°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

WT TGACTCTG\*GAAGGCAA
Mut TGACTCTG\*\*\*Insertion\*\*\*\*Deletion\*\*\*GAAGGCAA
Allele-1: 23bp insertion and 118bp deletion in exon1

WT TCTGGAGCCCCGAC\*\*\*\*\*\*\*\*\*\*ACGAAGGCAAAATG
Mut TCTGGAGCCCCGAC\*\*\*Deletion\*\*\*ACGAAGGCAAAATG
Allele-2: 70bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and TEAD1 knockout (KO) HeLa cells, using sanger sequencing.