

# EGLN1 Knockout 293T Cell Line, Homozygous

Catalog No.: RM01962

## Basic Information

**Catalog No.**

RM01962

**Category**

Cell Line

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

EGLN1

**Species**

Human

**Gene ID**

54583

**Swiss Prot**

Q9GZT9

**Synonyms**C1orf12; ECYT3; HALAH; HIF-PH2;  
HIFPH2; HPH-2; HPH2; PHD2; SM20;  
ZMYND6

## Contact

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## Background

The protein encoded by this gene catalyzes the post-translational formation of 4-hydroxyproline in hypoxia-inducible factor (HIF) alpha proteins. HIF is a transcriptional complex that plays a central role in mammalian oxygen homeostasis. This protein functions as a cellular oxygen sensor, and under normal oxygen concentration, modification by prolyl hydroxylation is a key regulatory event that targets HIF subunits for proteasomal destruction via the von Hippel-Lindau ubiquitylation complex. Mutations in this gene are associated with erythrocytosis familial type 3 (ECYT3). [provided by RefSeq, Nov 2009]

## Product Information

**Description**

EGLN1 Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing Technology.

Allele-1:73bp deletion in exon1

Allele-2:74bp 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**

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.

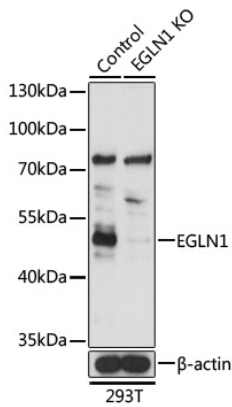
1. Thaw the vial in 37°C water bath, and shake it to melt as soon as possible.
2. Transfer the cell suspension to a 15mL conical tube with pre-warmed 5mL complete medium and centrifuge 1000rpm for approximately 5 minutes at room temperature.
3. Remove and discard the supernatant.
4. Resuspend the cell 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

WT GCCGGGACAACGCC\*\*\*\*\*GTGCGGCCGCCGGC  
Mut GCCGGGACAACGCC\*\*\*Deletion\*\*\*GTGCGGCCGCCGGC  
Allele-1: 73bp deletion in exon1  
WT CGCCGGGACAACGCC\*\*\*\*\*GTGCGGCCGCCGGC  
Mut CGCCGGGACAACGCC\*\*\*Deletion\*\*\*GTGCGGCCGCCGGC  
Allele-2: 74bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and EGLN1 knockout (KO) 293T cells, using sanger sequencing.

## WB data



Western blot analysis of extracts from parental (Control) and EGLN1 Knockout 293T Cell Line, using EGLN1 antibody at 1:3000 dilution.