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CSK Knockdown 293T Cell Line, Heterozygous

Catalog No.: RM02665

Basic Information

Catalog No. RM02665

Category Cell Line

Parental Cell line 293T

Genotype Knockdown

Gene Information

Gene Symbol CSK

Species Human

Gene ID 1445

Swiss Prot P41240

Contact

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Product Information

Description

CSK Knockdown 293T Cell Line, Heterozygote is engineered from 293T cell line with Gene-Editing technology. Allele-1:84bp deletion in exon2 Allele-2:86bp 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 Dry ice

Amount 1~5x10⁶ 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% CO₂ 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% $\mbox{CO}_2.$
- 7. A subcultivation ratio of 1:2-1:4 is recommended.



Background

www.abclonal.co

Sequencing data

WT CGCCTGGCCATCCG***Deletion***TGGCCGTCACCAAG Mut CGCCTGGCCATCCG***Deletion***TGGCCGTCACCAAG Allele-1: 84bp deletion in exon2

WT CGCCTGGCCATCCG***Deletion***GCCGTCACCAAGGT Mut CGCCTGGCCATCCG***Deletion***GCCGTCACCAAGGT Allele-2: 86bp deletion in exon2 Genome sequence analysis of PCR products from parental (WT) and CSK Knockdown (KD) 293T cells, using sanger sequencing.