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ARHGEF1 Knockout 293T Cell Line, Homozygous

Catalog No.: RM02500

Basic Information

Catalog No.
RM02500

Category
Cell Line

Parental Cell line
293T

Genotype
Knockout

Background

Rho GTPases play a fundamental role in numerous cellular processes that are initiated by extracellular stimuli that work through G protein coupled receptors. The encoded protein may form complex with G proteins and stimulate Rho-dependent signals. Multiple alternatively spliced transcript variants have been found for this gene, but the full-length nature of some variants has not been defined. [provided by RefSeq, Jul 2008]

Gene Information

Gene Symbol
ARHGEF1

Species
Human

Gene ID
9138

Swiss Prot
Q92888

Synonyms
GEF1; LBCL2; LSC; P115-RHOGEF; SUB1.5

Product Information

Description
ARHGEF1 Knockout 293T Cell line is engineered from 293T cell line with Gene-Editing Technology.
Allele-1:exon6 was deleted
Allele-2:exon6 was deleted
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
1~5x10^6 cells/vial

Storage
Stored in liquid nitrogen for a long time less than -130℃. 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℃ with 5% CO_2 condition.
1. Thaw the vial in 37℃ 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℃ incubator with 5% CO_2.
7. A subcultivation ratio of 1:2-1:4 is recommended.

Contact

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Sequencing data

Genome sequence analysis of PCR products from parental (WT) and ARHGEF1 knockout (KO) 293T cells, using sanger sequencing.