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GDI2 Knockout 293T Cell Line, Homozygous

Catalog No.: RM02237

Basic Information

Catalog No.
RM02237

Category
Cell Line

Parental Cell line
293T

Genotype
Knockout

Background

GDP dissociation inhibitors are proteins that regulate the GDP-GTP exchange reaction of members of the rab family, small GTP-binding proteins of the ras superfamily, that are involved in vesicular trafficking of molecules between cellular organelles. GDIs slow the rate of dissociation of GDP from rab proteins and release GDP from membrane-bound rabs. GDI2 is ubiquitously expressed. The GDI2 gene contains many repetitive elements indicating that it may be prone to inversion/deletion rearrangements. Alternative splicing results in multiple transcript variants encoding distinct isoforms. [provided by RefSeq, Jul 2008]

Gene Information

Gene Symbol
GDI2

Species
Human

Gene ID
2665

Swiss Prot
P50395

Synonyms
HEL-S-46e; RABGDIB

Product Information

Description
GDI2 Knockout cell line is engineered from 293T cell line with Gene-Editing Technology.
Allele-1:82bp deletion in exon2
Allele-2:82bp 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
1~5x10^6 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

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WT     AATGTATCCTGTCA**************CATTGGAAGATGTA
Mut    AATGTATCCTGTCA***Deletion***CATTGGAAGATGTA
Allele-1：82bp deletion in exon2
WT     AATGTATCCTGTCA**************CATTGGAAGATGTA
Mut    AATGTATCCTGTCA***Deletion***CATTGGAAGATGTA
Allele-2：82bp deletion in exon2
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Genome sequence analysis of PCR products from parental (WT) and GDI2 knockout (KO) 293T cells, using sanger sequencing.