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BCL10 Knockout 293T Cell Line, Homozygous

Catalog No.: RM02449

Basic Information

Catalog No.
RM02449

Category
Cell Line

Parental Cell line
293T

Genotype
Knockout

Background

This gene was identified by its translocation in a case of mucosa-associated lymphoid tissue (MALT) lymphoma. The protein encoded by this gene contains a caspase recruitment domain (CARD), and has been shown to induce apoptosis and to activate NF-kappaB. This protein is reported to interact with other CARD domain containing proteins including CARD9, 10, 11 and 14, which are thought to function as upstream regulators in NF-kappaB signaling. This protein is found to form a complex with MALT1, a protein encoded by another gene known to be translocated in MALT lymphoma. MALT1 and this protein are thought to synergize in the activation of NF-kappaB, and the deregulation of either of them may contribute to the same pathogenetic process that leads to the malignancy. Alternative splicing results in multiple transcript variants. [provided by RefSeq, Mar 2016]

Gene Information

Gene Symbol
BCL10

Species
Human

Gene ID
8915

Swiss Prot
O95999

Synonyms
CARMEN; CIPER; CLAP; IMD37; c-E10; mE10

Product Information

Description
BCL10 Knockout 293T Cell line is engineered from 293T cell line with Gene-Editing Technology.
Allele-1:exon1 was deleted
Allele-2:exon1 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

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WT    CCCGAGCTCCCGGA**********GGTCGGAGGAGCGG
Mut   CCCGAGCTCCCGGA***Deletion***GGTCGGAGGAGCGG
Allele-1:   exon1 was deleted

WT    CCCGAGCTCCCGGA**********GGTCGGAGGAGCGG
Mut   CCCGAGCTCCCGGA***Deletion***GGTCGGAGGAGCGG
Allele-2:   exon1 was deleted
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Genome sequence analysis of PCR products from parental (WT) and BCL10 knockout (KO) 293T cells, using sanger sequencing.