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# **BCL10 Knockout 293T Cell Lysate, Homozygous**

Catalog No.: RM02487

### **Basic Information**

## Catalog No.

RM02487

#### Category

Cell Lysate

#### **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**

095999

# **Synonyms**

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

#### **Contact**

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#### **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 Lysate and 1 vial knockout cell Lysate

**Shipping Conditions** 4°C

**Amount** 50μL, 2μg/μL.

#### Storage

Lysate is stable for 12 months when stored at -20°C. Minimizing freeze-thaw cycles.

#### **Protocol**

To be used as WB control. Lysate is supplied in  $1\times$  SDS sample buffer (2% SDS, 60 mM Tris-HCl pH 6.8, 10% Glycerol, 0.02% Bromophenol blue, 60 mM beta-mercaptoethanol). Lysate should be boiled for 3 - 5 minutes before loading onto gel.

# Sequencing data

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

Genome sequence analysis of PCR products from parental (WT) and BCL10 knockout (KO) 293T cells, using sanger sequencing.