

# CASP9 Knockout 293T Cell Lysate, Homozygous

**Catalog No.:** RM02303

## Basic Information

**Catalog No.**

RM02303

**Category**

Cell Lysate

**Parental Cell line**

293T

**Genotype**

Knockout

## Gene Information

**Gene Symbol**

CASP9

**Species**

Human

**Gene ID**

842

**Swiss Prot**

P55211

**Synonyms**APAF-3; APAF3; ICE-LAP6; MCH6;  
PPP1R56

## Contact

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## Background

This gene encodes a member of the cysteine-aspartic acid protease (caspase) family. Sequential activation of caspases plays a central role in the execution-phase of cell apoptosis. Caspases exist as inactive proenzymes which undergo proteolytic processing at conserved aspartic residues to produce two subunits, large and small, that dimerize to form the active enzyme. This protein can undergo autoproteolytic processing and activation by the apoptosome, a protein complex of cytochrome c and the apoptotic peptidase activating factor 1; this step is thought to be one of the earliest in the caspase activation cascade. This protein is thought to play a central role in apoptosis and to be a tumor suppressor. Alternative splicing results in multiple transcript variants. [provided by RefSeq, May 2013]

## Product Information

**Description**

CASP9 Knockout 293T Cell Line is engineered from 293T cell line with Gene-Editing technology.

Allele-1:53bp deletion in exon1

Allele-2:77bp deletion in exon1

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× 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

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WT TGCTCAGACCAGAG\*\*\*\*\*TTCTGGAGGATTG  
Mut TGCTCAGACCAGAG\*\*\*Deletion\*\*\*TTCTGGAGGATTG  
Allele-1: 53bp deletion in exon1

WT GTGCTCAGACCAGA\*\*\*\*\*ATTTAATTTTAGCA  
Mut GTGCTCAGACCAGA\*\*\*Deletion\*\*\*ATTTAATTTTAGCA  
Allele-2: 77bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and CASP9 Knockout (KO) 293T cells, using sanger sequencing.