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# CAPN2 Knockout HeLa Cell Lysate, Homozygous

Catalog No.: RM02367

# **Basic Information**

#### Catalog No.

RM02367

# Category

Cell Lysate

# **Parental Cell line**

HeLa

#### Genotype

Knockout

# **Background**

The calpains, calcium-activated neutral proteases, are nonlysosomal, intracellular cysteine proteases. The mammalian calpains include ubiquitous, stomach-specific, and muscle-specific proteins. The ubiquitous enzymes consist of heterodimers with distinct large, catalytic subunits associated with a common small, regulatory subunit. This gene encodes the large subunit of the ubiquitous enzyme, calpain 2. Multiple heterogeneous transcriptional start sites in the 5' UTR have been reported. Two transcript variants encoding different isoforms have been found for this gene. [provided by RefSeq, Mar 2009]

#### **Gene Information**

# **Gene Symbol**

CAPN2

# **Species**

Human

# **Gene ID**

824

#### **Swiss Prot**

P17655

# **Synonyms**

CANP2; CANPL2; CANPml; mCANP

# **Contact**

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# **Product Information**

#### **Description**

CAPN2 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.

Allele-1:exon2 was deleted

Allele-2:exon2 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**

Amount

4°C

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 TTGTGGGTCTCGTT\*\*\*\*\*\*\*\*\*\*TGTAAGGCTGTGGG
Mut TTGTGGGTCTCGTT\*\*\*Deletion\*\*\*TGTAAGGCTGTGGG

Allele-1: exon2 was deleted

WT TTGTGGGTCTCGTT\*\*\*\*\*\*\*\*\*\*\*TGTAAGGCTGTGGG
Mut TTGTGGGTCTCGTT\*\*\*Deletion\*\*\*TGTAAGGCTGTGGG

Allele-2: exon2 was deleted

Genome sequence analysis of PCR products from parental (WT) and CAPN2 knockout (KO) HeLa cells, using sanger sequencing.