

# ENO1 Knockdown HeLa Cell Lysate, Heterozygous

Catalog No.: RM01776

## Basic Information

**Catalog No.**

RM01776

**Category**

Cell Lysate

**Parental Cell line**

HeLa

**Genotype**

Knockdown

## Gene Information

**Species**

Human

**Gene ID**

2023

**Swiss Prot**

P06733

**Synonyms**

ENO1L1; HEL-S-17; MPB1; NNE; PPH

## Contact

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## Background

This gene encodes alpha-enolase, one of three enolase isoenzymes found in mammals. Each isoenzyme is a homodimer composed of 2 alpha, 2 gamma, or 2 beta subunits, and functions as a glycolytic enzyme. Alpha-enolase in addition, functions as a structural lens protein (tau-crystallin) in the monomeric form. Alternative splicing of this gene results in a shorter isoform that has been shown to bind to the c-myc promoter and function as a tumor suppressor. Several pseudogenes have been identified, including one on the long arm of chromosome 1. Alpha-enolase has also been identified as an autoantigen in Hashimoto encephalopathy. [provided by RefSeq, Jan 2011]

## Product Information

**Description**

ENO1 Knockdown HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.

Allele-1:WT

Allele-2:exon3 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× 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 CAGTGGTTCTCTCT\*\*\*\*\*TAATGCCACCAGAG  
Mut CAGTGGTTCTCTCT\*\*\*\*\*TAATGCCACCAGAG  
Allele-1: WT  
WT CGCGTCGGCCTCAA\*\*\*\*\*TCCCAGGCCAGGG  
Mut CGCGTCGGCCTCAA\*\*\*Deletion\*\*\*TCCCAGGCCAGGG  
Allele-2: exon3 was deleted

Genome sequence analysis of PCR products from parental (WT) and ENO1 Knockdown (KD) HeLa cells, using sanger sequencing.