

ZEB1 Knockout HeLa Cell Lysate, Homozygous

Catalog No.: RM02266

Basic Information

Catalog No.

RM02266

Category

Cell Lysate

Parental Cell line

HeLa

Genotype

Knockout

Gene Information

Gene Symbol

ZEB1

Species

Human

Gene ID

6935

Swiss Prot

P37275

Synonyms

AREB6; BZP; DELTAEF1; FECD6; NIL2A;
PPCD3; TCF8; ZFHEP; ZFHX1A

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Background

This gene encodes a zinc finger transcription factor. The encoded protein likely plays a role in transcriptional repression of interleukin 2. Mutations in this gene have been associated with posterior polymorphous corneal dystrophy-3 and late-onset Fuchs endothelial corneal dystrophy. Alternatively spliced transcript variants encoding different isoforms have been described.[provided by RefSeq, Mar 2010]

Product Information

Description

ZEB1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.

Allele-1:64bp deletion in exon6

Allele-2:64bp deletion in exon6

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

WT AATGGGCGACCAAG*****CGACCACAGATACG
Mut AATGGGCGACCAAG***Deletion***CGACCACAGATACG
Allele-1: 64bp deletion in exon6
WT AATGGGCGACCAAG*****CGACCACAGATACG
Mut AATGGGCGACCAAG***Deletion***CGACCACAGATACG
Allele-2: 64bp deletion in exon6

Genome sequence analysis of PCR products from parental (WT) and ZEB1 Knockout (KO) HeLa cells, using sanger sequencing.