

CCND1 Knockout HeLa Cell Lysate, Homozygous

Catalog No.: RM01985

Basic Information

Catalog No.

RM01985

Category

Cell Lysate

Parental Cell line

HeLa

Genotype

Knockout

Gene Information

Gene Symbol

CCND1

Species

Human

Gene ID

595

Swiss Prot

P24385

Synonyms

BCL1; D11S287E; PRAD1; U21B31

Contact

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Background

The protein encoded by this gene belongs to the highly conserved cyclin family, whose members are characterized by a dramatic periodicity in protein abundance throughout the cell cycle. Cyclins function as regulators of CDK kinases. Different cyclins exhibit distinct expression and degradation patterns which contribute to the temporal coordination of each mitotic event. This cyclin forms a complex with and functions as a regulatory subunit of CDK4 or CDK6, whose activity is required for cell cycle G1/S transition. This protein has been shown to interact with tumor suppressor protein Rb and the expression of this gene is regulated positively by Rb. Mutations, amplification and overexpression of this gene, which alters cell cycle progression, are observed frequently in a variety of tumors and may contribute to tumorigenesis. [provided by RefSeq, Jul 2008]

Product Information

Description

CCND1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.

Allele-1:29bp deletion in exon1

Allele-2:29bp 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

WT AGTGGAACCATCC*****AACGACCGGGTGCT
Mut AGTGGAACCATCC***Deletion***AACGACCGGGTGCT
Allele-1: 29bp deletion in exon1
WT AGTGGAACCATCC*****AACGACCGGGTGCT
Mut AGTGGAACCATCC***Deletion***AACGACCGGGTGCT
Allele-2: 29bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and CCND1 knockout (KO) HeLa cells, using sanger sequencing.