# AKT1S1 Knockdown HeLa Cell Lysate, Heterozygous 

## Basic Information

## Catalog No.

RM02045

## Category

Cell Lysate

## Parental Cell line

HeLa
Genotype
Knockdown

## Gene Information

Gene Symbol
AKT1S1

Species
Human
Gene ID
84335

Swiss Prot
Q96B36

## Synonyms

Lobe; PRAS40

## Contact

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## Background

AKT1S1 is a proline-rich substrate of AKT (MIM 164730) that binds 14-3-3 protein (see YWHAH, MIM 113508) when phosphorylated (Kovacina et al., 2003 [PubMed 12524439]).[supplied by OMIM, Mar 2008]

## Product Information

## Description

AKT1S1 Knockdown HeLa Cell Line is engineered from HeLa cell line with Gene-Editing technology.
Allele-1:47bp deletion in exon2
Allele-2:48bp deletion in exon2
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

$4^{\circ} \mathrm{C}$

## Amount

$50 \mu \mathrm{~L}, 2 \mu \mathrm{~g} / \mu \mathrm{L}$.

## Storage

Lysate is stable for 12 months when stored at $-20^{\circ} \mathrm{C}$. 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 CTATGCTGCCCATG***************GCACTGGCCCACAG Mut CTATGCTGCCCATG***Deletion***GCACTGGCCCACAG Allele-1: 47bp deletion in exon2
WT CTATGCTGCCCATG*************** CACTGGCCCACAGG
Mut CTATGCTGCCCATG***Deletion***CACTGGCCCACAGG Allele-2: 48bp deletion in exon2

Genome sequence analysis of PCR products from parental (WT) and AKT1S1 Knockdown (KD) HeLa cells, using sanger sequencing.

