

# FAS Knockout HeLa Cell Line, Homozygous

Catalog No.: RM01874

## Basic Information

### Catalog No.

RM01874

### Category

Cell Line

### Parental Cell line

HeLa

### Genotype

Knockout

## Gene Information

### Gene Symbol

FAS

### Species

Human

### Gene ID

355

### Swiss Prot

P25445


### Synonyms

ALPS1A; APO-1; APT1; CD95; FAS1;  
FASTM; TNFRSF6

## Contact

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## Background

The protein encoded by this gene is a member of the TNF-receptor superfamily. This receptor contains a death domain. It has been shown to play a central role in the physiological regulation of programmed cell death, and has been implicated in the pathogenesis of various malignancies and diseases of the immune system. The interaction of this receptor with its ligand allows the formation of a death-inducing signaling complex that includes Fas-associated death domain protein (FADD), caspase 8, and caspase 10. The autoproteolytic processing of the caspases in the complex triggers a downstream caspase cascade, and leads to apoptosis. This receptor has been also shown to activate NF-kappaB, MAPK3/ERK1, and MAPK8/JNK, and is found to be involved in transducing the proliferating signals in normal diploid fibroblast and T cells. Several alternatively spliced transcript variants have been described, some of which are candidates for nonsense-mediated mRNA decay (NMD). The isoforms lacking the transmembrane domain may negatively regulate the apoptosis mediated by the full length isoform. [provided by RefSeq, Mar 2011]

## Product Information

### Description

FAS Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.  
Allele-1:61bp deletion in exon2  
Allele-2:61bp 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 line and 1 vial knockout cell line

### Shipping Conditions

Dry ice

### Amount

1~5x10<sup>6</sup> cells/vial

### Storage

Stored in liquid nitrogen for a long time less than -130°C. Minimizing freeze-thaw cycles.

### Protocol

Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at 37°C with 5% CO<sub>2</sub> condition.

1. Thaw the vial in 37°C water bath, and shake it to melt as soon as possible.
2. Transfer the cell suspension to a 15mL conical tube with pre-warmed 5mL complete medium and centrifuge 1000rpm for approximately 5 minutes at room temperature.
3. Remove and discard the supernatant.
4. Resuspend the cell pellet with 1mL pre-warmed complete medium and seed in 10cm dish.
5. Add 8-10mL of complete medium.
6. Incubate the culture at 37°C incubator with 5% CO<sub>2</sub>.
7. A subcultivation ratio of 1:2-1:4 is recommended.

## Sequencing data

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WT CATCAACTCCAAGG\*\*\*\*\*TGATGGCCAATTCT  
Mut CATCAACTCCAAGG\*\*\*Deletion\*\*\*TGATGGCCAATTCT  
Allele-1: 61bp deletion in exon2  
WT CATCAACTCCAAGG\*\*\*\*\*TGATGGCCAATTCT  
Mut CATCAACTCCAAGG\*\*\*Deletion\*\*\*TGATGGCCAATTCT  
Allele-2: 61bp deletion in exon2

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