Leader in Biomolecular Solutions for Life Science

www.abclonal.com

SAV1 Knockout HeLa Cell Line, Homozygous

Catalog No.: RM01937

Basic Information

Catalog No.
RM01937

Category
Cell Line

Parental Cell line
HeLa

Genotype
Knockout

Background

WW domain-containing proteins are found in all eukaryotes and play an important role in the regulation of a wide variety of cellular functions such as protein degradation, transcription, and RNA splicing. This gene encodes a protein with two WW domains, a SARAH domain, and a coiled-coil region and is ubiquitously expressed in adult tissues. This protein binds to MST1 (mammalian sterile 20-like kinase 1) and promotes MST1-induced apoptosis. It has also been shown to bind to HAX1 (hematopoietic cell-specific protein 1 (HS1)-associated protein X-1) and to attenuate the anti-apoptotic effects of HAX1. Studies in human and mouse suggest this gene acts as a tumor suppressor. [provided by RefSeq, Aug 2012]

Gene Information

Gene Symbol
SAV1

Species
Human

Gene ID
60485

Swiss Prot
Q9H4B6

Synonyms
SAV; WW45; WWP4

Product Information

Description
SAV1 Knockout HeLa Cell Line is engineered from HeLa cell line with Gene-Editing Technology.
Allele-1:exon1 was deleted
Allele-2:exon1 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 line and 1 vial knockout cell line

Shipping Conditions
Dry ice

Amount
1~5x10^6 cells/vial

Storage
Store at -20℃

Protocol
Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at 37℃ with 5% CO_2 condition.
1. Thaw the vial in 37℃ 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℃ incubator with 5% CO_2.
7. A subcultivation ratio of 1:2-1:4 is recommended.

Contact

400-999-6126

cn.market@abclonal.com.cn

www.abclonal.com.cn

Sequencing data

```
WT    CCCGGCGCGGAGCC**************TCGCGTGAAATACT
Mut   CCCGGCGCGGAGCC***Deletion***TCGCGTGAAATACT
Allele-1: exon1 was deleted

WT    CCCGGCGCGGAGCC**************TCGCGTGAAATACT
Mut   CCCGGCGCGGAGCC***Deletion***TCGCGTGAAATACT
Allele-2: exon1 was deleted
```

Genome sequence analysis of PCR products from parental (WT) and SAV1 knockout (KO) HeLa cells, using sanger sequencing.