abcam

Product datasheet

Human SMAD6 knockout A549 cell line ab277891

1 Image

Overview

Product name Human SMAD6 knockout A549 cell line

Parental Cell Line A549
Organism Human

Mutation description Knockout achieved by using CRISPR/Cas9, Homozygous: 109 bp deletion in exon 1

Passage number <20

Knockout validation Sanger Sequencing

Biosafety level

General notes Recommended control: Human wild-type A549 cell line (ab275)

Recommended control: Human wild-type A549 cell line (<u>ab275463</u>). Please note a wild-type cell line is not automatically included with a knockout cell line order, if required please add recommended wild-type cell line at no additional cost using the code WILDTYPE-TMTK1.

Cryopreservation cell medium: Cell Freezing Medium-DMSO Serum free media, contains 8.7% DMSO in MEM supplemented with methyl cellulose.

Culture medium: F-12K + 10% FBS

Initial handling guidelines: Upon arrival, the vial should be stored in liquid nitrogen vapor phase and not at -80°C. Storage at -80°C may result in loss of viability.

- 1. Thaw the vial in 37°C water bath for approximately 1-2 minutes.
- 2. Transfer the cell suspension (0.8 mL) to a 15 mL/50 mL conical sterile polypropylene centrifuge tube containing 8.4 mL pre-warmed culture medium, wash vial with an additional 0.8 mL culture medium (total volume 10 mL) to collect remaining cells, and centrifuge at 201 x g (rcf) for 5 minutes at room temperature. 10 mL represents minimum recommended dilution. 20 mL represents maximum recommended dilution.
- 3. Resuspend the cell pellet in 5 mL pre-warmed culture medium and count using a haemocytometer or alternative cell counting method. Based on cell count, seed cells in an appropriate cell culture flask at a density of 2x10³-1x10⁴ cells/cm². Seeding density is given as a guide only and should be scaled to align with individual lab schedules.
- 4. Incubate the culture at 37°C incubator with 5% CO₂. Cultures should be monitored daily.

Subculture quidelines:

All seeding densities should be based on cell counts gained by established methods. A guide seeding density of $6x10^4$ cells/cm² is recommended.

A partial media change 24 hours prior to subculture may be helpful to encourage growth, if required.

Cells should be passaged when they have achieved 80-90% confluence.

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Do not exceed 7x10⁴ cells/cm².

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We will provide viable cells that proliferate on revival.

Properties

Number of cells 1 x 10⁶ cells/vial, 1 mL

Adherent /Suspension Adherent

Tissue Lung

Cell type epithelial

Disease Carcinoma

Gender Male

Mycoplasma free Yes

Storage instructions Shipped on Dry Ice. Store in liquid nitrogen.

Storage buffer Constituents: 8.7% Dimethylsulfoxide, 2% Cellulose, methyl ether

Target

Function Antagonist of signaling by TGF-beta (transforming growth factor) type 1 receptor superfamily

members; has been shown to inhibit selectively BMP (bone morphogenetic proteins) signaling by competing with the co-SMAD SMAD4 for receptor-activated SMAD1. SMAD6 is an inhibitory SMAD (I-SMAD) or antagonistic SMAD. Binds to regulatory elements in target promoter regions.

Tissue specificityUbiquitous in various organs, with higher levels in lung. Isoform B is up-regulated in diseased

heart tissue.

Sequence similarities Belongs to the dwarfin/SMAD family.

Contains 1 MH1 (MAD homology 1) domain. Contains 1 MH2 (MAD homology 2) domain.

Post-translational

Phosphorylated by BMP type 1 receptor kinase and by PRKX.

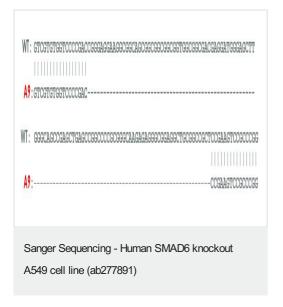
modifications

Ubiquitinated by WWP1.

Cellular localization

Nucleus.

Images



109 bp deletion in exon 1.

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