

SK-OV-3 | 300342

SK-OV-3 - SK-OV-3

Isoenzymes PGM3, 1, PGM1, 1-2, ES-D, 1, Me-2, 1, AK-1, 1, GLO-1, 1-2, G6PD, B, 0.0311

Tumorigenic SK-OV-3 is tumorigenic in nude mice.

Karyotype (P16) 46,XX,t(11;17)(p11;p11)

SK-OV-3

Culture Medium DMEM:Ham's F12 (1:1), w: 3.1 g/L $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$, w: 2.5 mM L- $\text{CaCl}_2 \cdot 2\text{H}_2\text{O}$, w: 15 mM HEPES, w: 0.5 mM $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$, w: 1.2 g/L NaHCO_3 820400a)

Supplements 10% FBS

Dissociation Reagent Trypsin

Subculturing SK-OV-3 cells are cultured in DMEM:Ham's F12 (1:1) supplemented with 10% FBS. For subculturing, cells are trypsinized and resuspended in DMEM:Ham's F12 (1:1) supplemented with 10% FBS. Cells are seeded into T25 flasks at a density of 1×10^4 cells per flask. Cells are harvested at 70-80% confluency.

Split ratio 1:2 or 1:3

Seeding density 1×10^4 cells/flask

Post-Thaw Recovery SK-OV-3 cells are thawed in a water bath at 37°C. Cells are washed with PBS and resuspended in DMEM:Ham's F12 (1:1) supplemented with 10% FBS. Cells are seeded into T25 flasks and allowed to recover for 24 hours before use.

Freeze medium DMEM:Ham's F12 (1:1) supplemented with 10% FBS + 10% DMSO

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Thawing and Culturing Cells

1. Thaw the cells rapidly in a water bath at 37°C. Do not allow the cells to reach room temperature. After thawing, centrifuge at 300 x g for 3 minutes. Remove the supernatant and resuspend the cells in 15 ml of fresh medium. Seed the cells into 8 wells of a 96-well plate.
2. Incubate the cells at 37°C in 5% CO₂ for 24 hours. After 24 hours, the cells should be at 70% confluency.
3. Harvest the cells and count them. Seed the cells into 8 wells of a 96-well plate.
4. Incubate the cells at 37°C in 5% CO₂ for 24 hours. After 24 hours, the cells should be at 70% confluency.
5. Harvest the cells and count them. Seed the cells into 8 wells of a 96-well plate.
6. Incubate the cells at 37°C in 5% CO₂ for 24 hours. After 24 hours, the cells should be at 70% confluency.
7. Harvest the cells and count them. Seed the cells into 8 wells of a 96-well plate.
8. Incubate the cells at 37°C in 5% CO₂ for 24 hours. After 24 hours, the cells should be at 70% confluency.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Harvest cells and resuspend in freezing medium. Store at -80°C.

Shipping Conditions Store at -80°C.

Storage Conditions Store at -150°C for 196 weeks.

SK-OV-3 / SK-OV-3 / HLA

Sterility The cells are free of mycoplasmas and PCR detectable. The cells are free of mycoplasmas and PCR detectable.

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STR

Amelogenin: x,x

CSF1PO: 11

D13S317: 8,11

D16S539: 12

D5S818: 11

D7S820: 13,14

TH01: 9,9.3

TPOX: 8,11

vWA: 17,18

D3S1358: 14

D21S11: 30, 31, 31.2

D18S51: 16, 17, 18

Penta E: 5,13

Penta D: 12,13

D8S1179: 14,15

FGA: 24, 25, 26

HLA

A*: '03:01:01, '68:01:02

B*: '18:01:01, '35:01:01

C*: '04:01:01, '05:01:01

DRB1*: '01:01:01, '03:01:01

DQA1*: '01:01:01, '05:01:01

DQB1*: '02:01:01, '05:01:01

DPB1*: '02:01:02G, '04:01:01G

E: '01:01:01, '01:06:01