

MIA PaCa-2 | 300438

Thawing and Culturing Cells

1. Thaw the vial immediately in a water bath at 37°C. Do not allow the cells to warm to room temperature. Transfer the cells to a pre-warmed medium.
2. Centrifuge the cells at 300 x g for 3 minutes. Resuspend the cells in 15 µl of medium. Seed the cells into a 96-well plate at 1500 cells per well.
3. Incubate the cells in a humidified incubator at 37°C with 5% CO₂. Change the medium after 24 hours.
4. After 24 hours, the cells should be visible. If not, check the viability of the cells.
5. Seed the cells into a 96-well plate at 1500 cells per well. Incubate for 24 hours.
6. After 24 hours, the cells should be visible. If not, check the viability of the cells.
7. After 24 hours, the cells should be visible. If not, check the viability of the cells.
8. After 24 hours, the cells should be visible. If not, check the viability of the cells.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Seed cells into a 96-well plate at 1500 cells per well. Freeze the cells in a liquid nitrogen vapor phase.

Shipping Conditions Ship the cells in a dry ice container at -78°C.

Storage Conditions Store the cells at -150°C for up to 196 days.

Genotype / HLA

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

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■■■■ HLA

- A*: '01.01.1900 00:02
- B*: 14:02:01
- C*: 08:02:01
- DRB1*: 01:02:01
- DQA1*: 01:01:02
- DQB1*: 05:01:01
- DPB1*: 02:01:02
- E: 01:01:01