

Panc-1 | 300228

Thawing and Culturing Cells

1. Thaw the vial rapidly 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.
3. Incubate the cells at 37°C in 5% CO₂. The cells should reach 70% confluency within 7-10 days.
4. Harvest the cells by trypsinization. Seed the cells into a new flask.
5. Repeat the process for subsequent passages.
6. Store the cells in liquid nitrogen for long-term storage.
7. Thaw the cells and seed them into a new flask.
8. Repeat the process for subsequent passages.

Incubation Atmosphere 37°C, 5% CO₂, humidified air

Flask Coating None

Freezing Procedure Harvest cells by trypsinization, wash with PBS, resuspend in freezing medium, and store in liquid nitrogen.

Shipping Conditions Dry ice, -78°C

Storage Conditions -150°C, 196 Krypton

Genotype / HLA

Sterility PCR genotyping, negative results.

