

HROC313 HROC313Met | 300849

Thawing and Culturing Cells

1. **Thawing:** Thaw the vial rapidly in a 37°C water bath. Transfer the cells to a pre-warmed tube and centrifuge at 300 × g for 3 minutes. Remove the supernatant and resuspend the cells in 10 mL of pre-warmed medium. Seed the cells into a 75 cm² flask containing 50 mL of pre-warmed medium. Incubate at 37°C in 5% CO₂.
2. **Counting:** After 24 hours, count the cells and seed into a new flask at a density of 1 × 10⁶ cells per flask.
3. **Expansion:** Expand the cells in 100 mL of medium in a 250 cm² flask. Once the cells reach confluence, seed them into a 75 cm² flask.
4. **Passaging:** Pass the cells into a new flask when they reach 70-80% confluence. Use a trypsin-EDTA solution to detach the cells.
5. **Freezing:** For long-term storage, freeze the cells in a freezing medium and store at -150°C.
6. **Thawing:** Thaw the cells rapidly in a 37°C water bath and resuspend in 10 mL of medium.
7. **Seeding:** Seed the cells into a 75 cm² flask at a density of 1 × 10⁶ cells per flask.
8. **Expansion:** Expand the cells in 100 mL of medium in a 250 cm² flask.

Incubation Atmosphere 37 °C, 5% CO₂

Flask Coating Non-adherent

Freezing Procedure Freeze at -80°C

Shipping Conditions -78°C

Storage Conditions -150°C to -196°C

HLA

Sterility Sterile