





# CHO-CXCR7 CHO-CXCR7 | 305412L

## Thawing and Culturing Cells

1. Thaw the vial rapidly in a 37°C water bath. Remove the vial and centrifuge at 300 × g for 3 minutes. Discard the supernatant and resuspend the cells in 10 ml of pre-warmed complete medium.
2. Seed the cells into a T75 flask containing 70% pre-warmed complete medium.
3. Incubate the cells at 37°C in a humidified atmosphere with 5% CO<sub>2</sub>.
4. Once the cells have reached confluence, passage them into a new T75 flask.
5. Repeat the seeding and incubation process until the cells are ready for use.
6. For long-term storage, harvest the cells and cryopreserve them in liquid nitrogen.
7. Thaw the cells and culture them as described above.
8. Verify the identity and purity of the cells using PCR and HLA typing.

## Incubation Atmosphere

37°C, 5% CO<sub>2</sub>, humidified atmosphere.

## Shipping Conditions

Cryopreserved cell lines are shipped on dry ice in validated, insulated packaging with sufficient refrigerant to maintain approximately -78 °C throughout transit. On receipt, inspect the container immediately and transfer vials without delay to appropriate storage.

## Storage Conditions

For long-term preservation, place vials in vapor-phase liquid nitrogen at about -150 to -196 °C. Storage at -80 °C is acceptable only as a short interim step before transfer to liquid nitrogen.

## Genotyping / HLA

## Sterility

Genotyping and HLA typing are performed to verify the identity and purity of the cells. PCR analysis is used to confirm the presence of the CXCR7 gene. HLA typing is used to ensure that the cells are free of contamination from other cell lines.