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CellosaurusAccession CVCL_1714

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SUP-T1 | 305642

**Thawing and
Culturing Cells**

1. Remove the vial from liquid nitrogen storage and allow it to warm to room temperature.
2. Dilute the cell suspension into 10 ml of pre-warmed medium.
3. Seed the cells into a T75 flask containing 50 ml of pre-warmed medium.
4. Incubate the cells at 37 °C in 5% CO₂ atmosphere.
5. Monitor cell growth and confluency.
6. Once cells reach 70-80% confluency, they can be passaged.
7. For passage, trypsinize the cells and seed them into a new flask.
8. Repeat the process for subsequent passages.

**Incubation
Atmosphere**

37 °C, 5% CO₂

**Shipping
Conditions**

Shipped on dry ice, -78 °C

**Storage
Conditions**

Store at -150 °C to -196 °C in liquid nitrogen

Supernatant **Media** **Cells** **Reagents**

Sterility

Cells are tested for mycoplasma contamination using PCR. Media and supernatant are tested for mycoplasma contamination using PCR.