

AT-1 | 500121

Supplements FBS 10%

Dissociation Reagent

Subculturing PBS T25

Seeding density 1×10^4

Fluid renewal 2-3

Post-Thaw Recovery 48

Freeze medium (FBS) + 10% DMSO

Thawing and Culturing Cells

1. Thaw cryovial in 37°C water bath.
2. Add 10ml of pre-warmed medium to cryovial.
3. Transfer cells to T25 flask.
4. Incubate at 37°C, 5% CO₂.
5. Check for cell attachment after 15-20 minutes.
6. Add 300 µl of medium to each well.
7. Incubate for 10-15 minutes.
8. Add 100 µl of medium to each well.

Incubation Atmosphere 37°C, 5% CO₂

Flask Coating

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