

Product sheet

SV40 MES 13 | 305183

Supplements 10% FBS, 14 mM HEPES

Dissociation Reagent

Subculturing 1. Wash cells with PBS. 2. Add 2-3 ml of dissociation reagent. 3. Incubate at 37°C for 5-10 minutes. 4. Add 10 ml of PBS and pipette up and down. 5. Seed cells into a new flask.

Fluid renewal 2-3 times per week

Freeze medium 10% FBS + 10% DMSO

- Thawing and Culturing Cells**
1. Thaw cells in a 37°C water bath.
 2. Add 10 ml of PBS and pipette up and down.
 3. Seed cells into a flask with 10 ml of fresh medium.
 4. Incubate at 37°C with 5% CO₂.
 5. Monitor cell growth and fluid renewal.
 6. Pass cells when they reach 70-80% confluency.
 7. Use cells for experiments or freezing.
 8. Store cells in liquid nitrogen.

Incubation Atmosphere 37°C, 5% CO₂

Flask Coating

Freezing Procedure

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Shipping Conditions -78°C

Storage Conditions -150 °C 196

HLA

Sterility