

HEK293-CLDN6 | 305985

Culture Medium RPMI 1640, w: 2.0 mM β -mercaptoethanol, w: 2.0 mM/1000 NaHCO₃ (Cytion 820700a)

Supplements FBS 10%, 1 mM HEPES 10, NEAA 1%

Dissociation Reagent

Subculturing

Fluid renewal 2-3

Post-Thaw Recovery

Freeze medium

Thawing and Culturing Cells

1. Thaw cells in a 37°C water bath.
2. Add 10 ml of pre-warmed culture medium to the cryovial.
3. Centrifuge at 300 x g for 3 minutes.
4. Resuspend cells in 10 ml of culture medium.
5. Seed cells into a T25 flask.
6. Incubate cells in a 37°C incubator with 5% CO₂.
7. Monitor cell growth and confluency.
8. Harvest cells when they reach 70-80% confluency.

Incubation Atmosphere 37°C, 5% CO₂

Shipping Conditions

