

HEK293T-S-117 | 300329

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

Supplements FBS 10%

Dissociation Reagent

Subculturing PBS T25

Split ratio 1:4 1:8

Seeding density 1×10^4 cells/cm²

Fluid renewal 2-3 times

Post-Thaw Recovery 4-24 hours

Freeze medium (FBS) + 10% DMSO

- Thawing and Culturing Cells**
1. Thaw cryovial in 37°C water bath
 2. Transfer cells to cryovial and centrifuge at 300 x g for 3 min
 3. Remove supernatant and resuspend cells in 10 ml of culture medium
 4. Seed cells into T25 flask at 70% confluency
 5. Incubate cells for 15 min at 37°C
 6. Add fresh medium
 7. Monitor cell growth and confluency
 8. Harvest cells when reaching 70-80% confluency

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XXXXXXXXXX HLA
A*: '01:01:01
B*: '37:01:01
C*: '06:02:01
DRB1*: XXXX11:01:01
DQA1*: '05:05:01
DQB1*: '03:01:01
DPB1*: '04:01:01
E: '01:01:01