

Jurkat E6.1 | 300223

Thawing and Culturing Cells

1. Thaw the vial rapidly in a water bath at 37°C. Do not allow the cells to warm to room temperature. Transfer the cells to a pre-warmed T25 flask containing 10 mL of complete medium.
2. Allow the cells to settle for 15 minutes. Add 10 mL of complete medium to the flask. Incubate at 37°C in 5% CO₂.
3. After 24 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.
4. After 48 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.
5. After 72 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.
6. After 96 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.
7. After 120 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.
8. After 144 hours, check the cell density. If the cell density is 1×10^5 cells/mL, add 10 mL of complete medium to the flask.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Harvest cells into a 15 mL centrifuge tube. Spin at 300 x g for 3 minutes. Resuspend the pellet in 1 mL of freezing medium. Aliquot into 1 mL vials. Store at -80°C.

Shipping Conditions Store at -80°C. Ship on dry ice.

Storage Conditions Store at -150°C for 196 days.

HLA

Sterility The cells are free of mycoplasmas and PCR detectable. The cells are free of endotoxins.

XXXX Jurkat E6.1 | 300223

XXXXXX HLA

A*: 03:01:01
B*: 07:02:01, 35:03:01
C*: 04:01:01, 07:02:01
DRB1*: 07:01:01, 15:01:01
DQA1*: '01:02:01, '02:01:01
DQB1*: 02:02:01, 06:03:01
DPB1*: '02:01:02G, '04:02:01G