

HEK293T HCT116 | 300195

Thawing and Culturing Cells

1. Thaw the vial immediately in a 37°C water bath. Gently mix the cells by pipetting up and down. Centrifuge at 300 x g for 3 minutes. Resuspend the cells in 10 ml of complete DMEM medium. Seed the cells into a T25 flask.
2. Incubate the cells at 37°C in 5% CO₂ until they reach 70-80% confluency.
3. Harvest the cells by trypsinization. Seed the cells into a T75 flask.
4. Incubate the cells at 37°C in 5% CO₂ until they reach 70-80% confluency.
5. Harvest the cells by trypsinization. Seed the cells into a T75 flask.
6. Incubate the cells at 37°C in 5% CO₂ until they reach 70-80% confluency.
7. Harvest the cells by trypsinization. Seed the cells into a T75 flask.
8. Incubate the cells at 37°C in 5% CO₂ until they reach 70-80% confluency.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Harvest cells by trypsinization. Resuspend in freezing medium. Aliquot into 1 ml vials. Store at -80°C.

Shipping Conditions Dry ice, -78°C

Storage Conditions -150°C, 196 liquid nitrogen

HEK293T / HEK293T / HLA

Sterility HEK293T cells are tested for mycoplasma contamination. PCR testing is performed. HEK293T cells are free of mycoplasma contamination.

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A*: '01:01:01, '02:01:01

B*: 18:01:01, 21:01:01

C*: 05:01:01, 07:01:01

DRB1*: 03:01:01, 11:02:01

DQA1*: 05:01:01, 05:05:01

DQB1*: '02:01:01, '03:19:01

DPB1*: '03:01:01G, '04:02:01G

E: 01:01, 01:03