

HEK293T HROG12 T0 M1 | 300882

Dissociation Reagent Trypsin

Subculturing Seed cells into fresh wells with 100 µl of dissociation medium (100 µl of PBS + 100 µl of Trypsin) into T25, 100 µl of 3-5 µl of PBS, 100 µl of 3 µl. Seed cells into fresh wells, 100 µl of dissociation medium (100 µl of PBS + 100 µl of Trypsin) into T25, 100 µl of 3-5 µl of PBS, 100 µl of 3 µl.

Freeze medium DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100), 100 µl

Thawing and Culturing Cells

1. Thaw cells in a 37°C water bath, add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask.
2. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
3. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
4. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
5. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
6. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
7. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.
8. Add 100 µl of DMEM + 10% FBS + 10% DMSO, CM-1 (Cytion 800100) into a 25 cm² flask, incubate at 37°C for 24 hours.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Seed cells into fresh wells with 100 µl of dissociation medium (100 µl of PBS + 100 µl of Trypsin) into T25, 100 µl of 3-5 µl of PBS, 100 µl of 3 µl.

Shipping Conditions Dry ice, -78°C

Storage Conditions Dry ice, -78°C

