

HK-2xCRISPR-CAP-D2-mEGFP | 301572

Thawing and Culturing Cells

1. Thaw the cells rapidly in a water bath at 37°C. Transfer the cells to a pre-warmed medium.
2. Seed the cells into a flask containing 10 mL of pre-warmed medium. Incubate at 37°C with 5% CO₂.
3. Once the cells have reached confluence, they can be used for experiments or passaged.
4. For passaging, remove the medium and wash the cells with PBS. Add 1 mL of trypsin solution and incubate at 37°C for 5 minutes.
5. Add 1 mL of medium to stop the trypsin reaction. Pipette up the cells and transfer to a new flask.
6. Add 10 mL of fresh medium to the flask. Incubate at 37°C with 5% CO₂.
7. When the cells reach 70-80% confluence, they can be passaged again.
8. For long-term storage, seed cells into a flask with 10 mL of medium. Once confluent, add 1 mL of freezing medium and incubate at 37°C for 24 hours.

Incubation Atmosphere 37°C, 5% CO₂, humidified

Flask Coating None

Freezing Procedure Seed cells into a flask with 10 mL of medium. Once confluent, add 1 mL of freezing medium and incubate at 37°C for 24 hours. Then, freeze the cells in a liquid nitrogen vapor phase.

Shipping Conditions Dry ice, -78°C

Storage Conditions -150°C, 196 K

Genotype / HLA

Sterility

PCR genotyping of the CRISPR-Cas9 system. The results show that the cells are free of the Cas9 gene and contain the expected CRISPR-Cas9 system.

Flow cytometry analysis of the cells shows that they are free of GFP expression, indicating that the CRISPR-Cas9 system is not active.