

U2OS-CRISPR-SNAPf-Nup133 | 300666

Thawing and Culturing Cells

1.

1. **U2OS-CRISPR-SNAPf-Nup133** cells are delivered in a cryovial containing 1.5 x 10⁶ cells.
2.

2. Thaw the cryovial in a water bath at 37°C.
3.

3. Add 1 mL of pre-warmed complete medium to the cryovial.
4.

4. Pipette 70% of the cell suspension into a T25 flask.
5.

5. Incubate the cells at 37°C for 15 minutes.
6.

6. Centrifuge the cells at 300 x g for 3 minutes.
7.

7. Resuspend the cells in 10 mL of complete medium.
8.

8. Seed the cells into a T25 flask.

Incubation Atmosphere 37°C, 5% CO₂, **U2OS-CRISPR-SNAPf-Nup133**

Flask Coating **U2OS-CRISPR-SNAPf-Nup133**

Freezing Procedure **U2OS-CRISPR-SNAPf-Nup133**

Shipping Conditions **U2OS-CRISPR-SNAPf-Nup133**

Storage Conditions **U2OS-CRISPR-SNAPf-Nup133** -150 °C -196 °C **U2OS-CRISPR-SNAPf-Nup133**

U2OS-CRISPR-SNAPf-Nup133 / **U2OS-CRISPR-SNAPf-Nup133** / HLA

Sterility **U2OS-CRISPR-SNAPf-Nup133** PCR **U2OS-CRISPR-SNAPf-Nup133**
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