HROC450Met1 T0 M1

300725

GENERAL INFORMATION

Description

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IDENTIFIERS

Citation

HROC450Met1 T0 M1 (□□□□□□□□□□□□□□□□□ Cytion 300725)

NCBI TaxID	9606

HANDLING AND CULTURE

Culture medium

DMEM:Ham's F12 (1:1), w: 3.1 □□□□/□□□□ □□□□□□, w: 2.5 □□□□□□□□□□□□ L-□□□□□□□□□, w: 15 □□□□□□□□□□□□ HEPES, w: 0.5 □□□□□□□□□□□□ □□□□□□□□□□□□□□□□, w: 1.2 □□□□/□□□□ NaHCO3 (□□□□□□□□□□□□□□□ Cytion 820400a)

Incubation atmosphere	37°C, 5%CO2, □□□□□□□□□□□□□

Supplements

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Dissociation reagent	TrypLE Express 15 □□□□ 37°C

Subculturing

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Thawing and culturing

Confirm that the vial remains deeply frozen upon delivery, as cells are shipped on dry ice to maintain optimal temperatures during transit.
Upon receipt, either store the cryovial immediately at temperatures below -150°C to ensure the preservation of cellular integrity, or proceed to step 3 if immediate culturing is required.
For immediate culturing, swiftly thaw the vial by immersing it in a 37°C water bath with clean water and an antimicrobial agent, agitating gently for 40-60 seconds until a small ice clump remains.
Perform all subsequent steps under sterile conditions in a flow hood, disinfecting the cryovial with 70% ethanol before opening.
Carefully open the disinfected vial and transfer the cell suspension into a 15 ml centrifuge tube containing 8 ml of room-temperature culture medium, mixing gently.
Centrifuge the mixture at 200 x g for 5 minutes, carefully discard the supernatant containing freezing medium.
Follow the procedure described under Post-Thaw Recovery

QUALITY CONTROL AND CRYOPRESERVATION

Freeze medium	□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□□□ + 10% DMSO □□□□□□□□□□□□ □□□□□□□□□□□□□□□□□□□□□□□□□□□□

Storage conditions

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LEGAL AND SAFETY

Safety precautions

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