

CSI271Mu01

Primary Mouse Tubal Epithelial Cells (TuEC)

Organism Species: Mus musculus (Mouse)

Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1st Edition (Revised in Apr, 2024)

### [ DESCRIPTION ]

**Cell Type:** Epithelium **Synonyms:** TuEC

Species: Mus musculus (Mouse)
Tissue Source: Fallopian tube

Size: >5×10<sup>5</sup>cell/vial

## [PROPERTIES]

Cell activity: >85% (Viability by Trypan Blue Exclusion).

Formulation: Frozen 1 mL or T25 flask.

Biosafety: Negative for HIV-1, HBV, HCV, mycoplasma, bacteria, yeast and fungi.

Applications: For research use only. It is not approved for human or animal use, or for application in

clinical diagnostic procedures. **Growth Properties:** Adherent

## [CONTENTS]

Form & Buffer: Supplied as solution form in frozen stock solution, containing 90% FBS+10% DMSO.

# [USAGE]

Upon receiving the cells in a T-25 flask at room temperature, immediately transfer the cells to 37°C, 5% incubator; the cells in vials, directly and immediately transfer the cells from dry ice to liquid nitrogen.

#### **Culture conditions:**

DMEM/F12+5% FBS+1% Epithelial Cell Growth Supplement+1% Penicillin-Streptomycin Solution

Temperature: 37°C

Condition: 95% air, 5% carbon dioxide

### Cell recovery:

After receiving the cells, shake at 37°C in a water bath until completely dissolved, transfer to a 15 ml centrifuge tube, add 3-5 times complete culture solution, 1000 rpm for 5 min, discard the supernatant, and place in a T25 flask for culture.

#### Cell passage:

- 1. Cell passage when cell growth at 85-95%.
- 2. Discard the medium and wash with PBS 1-2 times.

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- Add 1 ml of Trypsin at 37°C, observe the cell under the microscope. If the cells are retracted and rounded, pat the culture flask to let the cells fall off. Stop digestion by adding 2 ml of complete medium containing 10% serum. Make it a single cell suspension.
- 4. Add the fresh medium to resuspend the cells. Unless otherwise stated, the recommended ratio of primary cells is 1/2.

## [Shipping]

Dry ice.

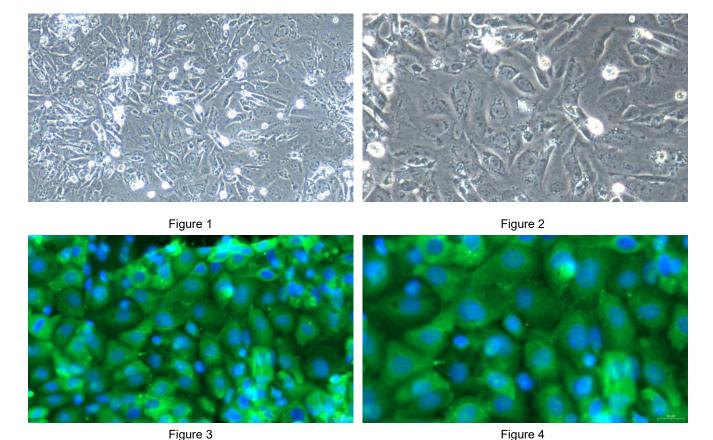
## [STORAGE]

Upon receiving, directly and immediately transfer the cells from dry ice to liquid nitrogen and keep the cells in liquid nitrogen until they are needed for experiments.

## [IMPORTANTNOTE]

- 1. The cultured cycle of Primary Mouse Tubal Epithelial Cells is limited in *vitro*. It is suggested that after cell resuscitation, the special growth medium and correct operation method recommended by us should be used for culture, and it should be used for follow-up experiments as soon as possible.
- 2. The cell is for research use only, and we will not be responsible for any issue if the cell was used in clinical diagnostic or any other procedures.

## [Figure]



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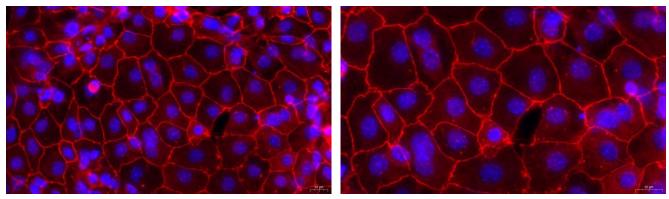


Figure 5 Figure 6

- Figure 1 Morphology of Mouse Tubal Epithelial Cells (Optical microscope,×100)
- Figure 2 Morphology of Mouse Tubal Epithelial Cells (Optical microscope,×200)
- Figure 3 Immunofluorescence identification of Cytokeratin 18 specific antibody (×400)
- Figure 4 Immunofluorescence identification of Cytokeratin 18 specific antibody (×600)
- Figure 5 Immunofluorescence identification of ZO-1 specific antibody (×400
- Figure 6 Immunofluorescence identification of ZO-1 specific antibody (×600)