

**CSI034Mu01**

**Pulmonary Fibroblasts (PF)**

**Organism Species: Mus musculus (Mouse)**

**Instruction manual**

FOR IN VITRO AND RESEARCH USE ONLY  
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

1<sup>st</sup> Edition (Revised in Jan, 2016)

## **[Description]**

**Product Name:** Pulmonary Fibroblasts (PF)

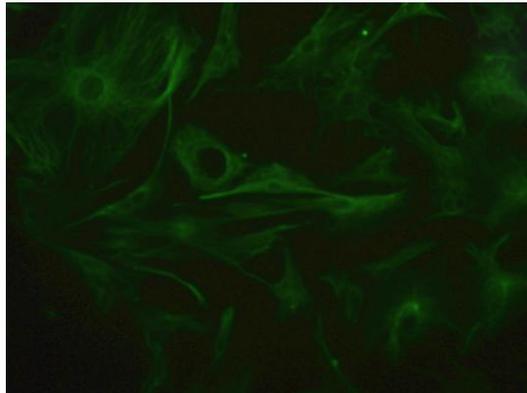
**Catalog Number:** CSI034Mu01

**Tissue Source:** Lung tissue

**Strain:** Balb/c

**Disease:** Normal

**Numbers of Cell:**  $>5 \times 10^5$  cells



Immunofluorescence Staining (IF) for Vimentin  
Pulmonary Fibroblasts

## **[Properties]**

**Cell Activity:**  $>90\%$

**Cell Adherence:** Adherence

**Bio-safety:** Negative for HIV-1, HBV, HCV, mycoplasma, bacteria, yeast and fungi.

## **[Contents]**

**Form & Buffer:** Supplied as solution form in cryopreserved medium (liquid nitrogen) or Supplied as solution form in Pericardial Fibroblasts culture Medium (RT).

## **[Shipping]**

Room temperature (T-25 flask) or Dry ice.

## **【Cell Resuscitation】**

1. Place the frozen vial in a 37°C water bath. Hold and rotate the vial gently until the contents completely thaw. Centrifuge the tube at 1000 rpm for 5mins. Resuspend cells in culture medium. A seeding density of 7,500 cells/cm<sup>2</sup> is recommended.
2. Upon receiving the cells in a T-25 flask at room temperature, immediately disinfect T-25 flask with 75% alcohol, transfer the cells to 37°C, 5% incubator.

**Caution:** *Cryopreserved cells are very delicate. Thaw the vial in a 37 °C water bath and return the cells to culture as quickly as possible with minimal handling!*

## **【Subculture】**

1. Subculture when the culture reaches 90% confluency. Warm complete medium, Trypsin/EDTA solution ( Ca<sup>2+</sup> and Mg<sup>2+</sup>-free ,T/E), T/E neutralization solution (TNS), and DPBS ( Ca<sup>2+</sup> and Mg<sup>2+</sup>-free ) to room temperature.
2. Rinse the cells with DPBS. Add 1mL 0.25% T/E solution into culture plate. Gently rock the plate complete coverage of cells by T/E solution. Incubate the plate in a 37°C incubator for 5 min or until cells completely round up.
3. Add 1 mL TNS solution to the plate and transfer detached cells to the 15 mL centrifuge tube. Rinse the plate with another 2 mL TNS to collect the residual cells. Centrifuge the 15 mL tube at 1000 rpm for 5 min.
4. Resuspend cells in culture medium. Count and plate cells in a new culture plate with the recommended cell density.

**Caution:** *Handling animal-derived products is potentially biohazardous. Always wear gloves and safety glasses when working with these materials.*

## **【Recommended Medium】**

It is recommended to use Pericardial Fibroblasts culture Medium for culturing *in vitro*.