CSI066Cp01

Primary Caprine Bladder Stromal Fibroblasts (BSF)

Organism Species: Capra hircus; Caprine (Goat)

Instruction manual

FOR RESEARCH USE ONLY
NOT FOR USE IN CLINICAL DIAGNOSTIC PROCEDURES

2nd Edition (Revised in Aug, 2026)

[DESCRIPTION]

Cell Type: Fibroblasts
Synonyms: BSF
Strain: Boer Goat
Age: 1-2 Years
Tissue Source: Bladder tissue
Disease: Normal
Size: >5×10^5 cells/vial
Growth properties: Adherent
Morphology: Polygonal

[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.

[CONTENTS]

Form & Buffer: Supplied as solution form in frozen stock solution (Protein-free, Chemical Defined Cell Cryopreservation Medium). ***BSF are cryopreserved at P1 and delivered frozen.***

[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+10%FBS+1% Fibroblasts growth supplement+1%Penicillin-Streptomycin Solution
Temperature:37°C
Condition: 95% air, 5% carbon dioxide
Medium Renewal: Every 1 to 2 days
Dissociation Solution: 0.25% Trypsin

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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.
3. 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.

[IMPORTANT NOTE]

1. The culture cycle of Primary Caprine Bladder Stromal Fibroblasts (BSF) is limited in vitro. It is recommended to use the specialized growth medium provided by Cloud-Clone Corp. and follow the correct operational procedures to ensure optimal culture conditions for these cells.
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]

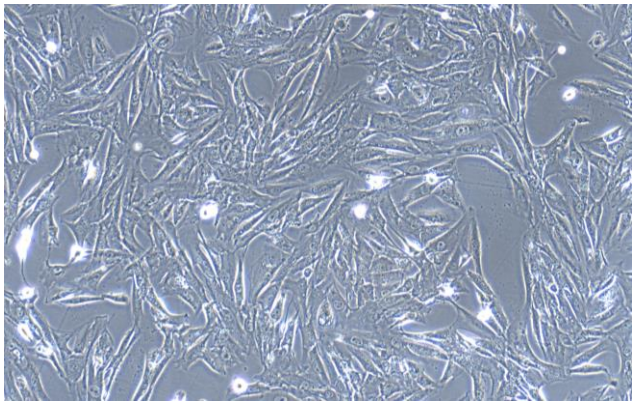

Figure 1

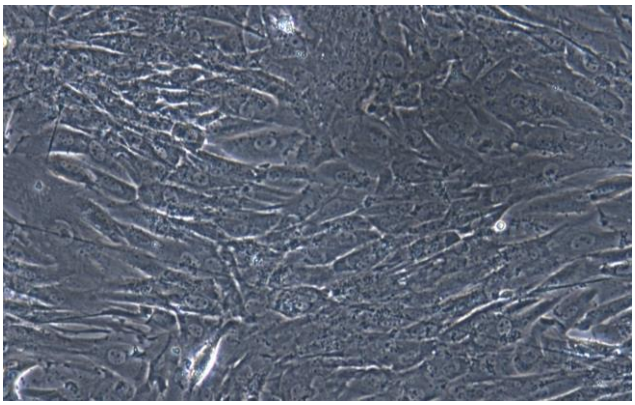

Figure 2

Figure 1 Morphology of Caprine Bladder Stromal Fibroblasts (Optical microscope,×100)

Figure 2 Morphology of Caprine Bladder Stromal Fibroblasts (Optical microscope,×200)

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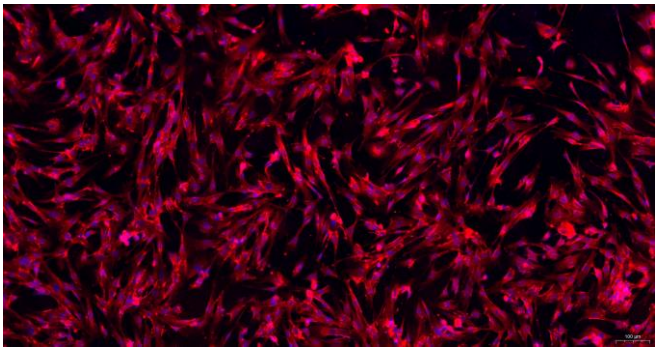

Figure 3

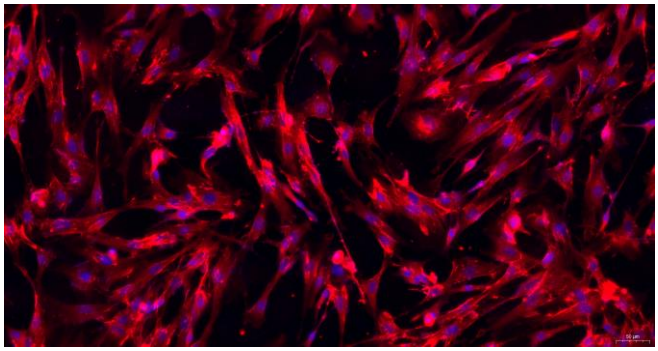

Figure 4

Figure 3 Immunofluorescence identification of Fibronectin specific antibody (×100)

Figure 4 Immunofluorescence identification of Fibronectin specific antibody (×200)

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