

SOP: Propagation of AG09309
Date modified: 11/12/09
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Ordering Information

AG09309 may be ordered from Coriell Cell Repositories. Proliferating cells are shipped in a T25 flask with 50-60ml of media.

To order starter cultures:

Name/Catalogue #: AG09309 (Adult Toe Fibroblast)

Notes:

This is an adherent cell line.

Materials List

1. Eagle's MEM (Cellgro Cat# 10-010-CM)
2. Characterized Fetal Bovine Serum (HyClone Cat# SH30071)
3. Non-essential Amino Acids, 100X (Invitrogen Cat# 11140-050)
4. T225 culture flasks
5. Graduated pipets (1, 5, 25, 50mL)
6. Penicillin-Streptomycin Solution, 200X (Cellgro Cat# 30-001-CI)
7. Phosphate Buffered Saline (1X PBS) (prepared from 10X stock Cat# by dilution with sterile deionized water)
8. Freezing medium (growth medium containing 6% DMSO)
9. DMSO
10. Cryovials
11. Accutase – Enzyme Cell Detachment Medium (EBioscience, Cat# 00-4555)
12. Hemocytometer
13. Micropipet w/ P20 tips
14. Microscope

Growth Medium for AG09309

Eagle's MEM with Earle's salts and L-Glutamine

15% FBS

Non-essential Amino Acids (1X)

Pen-Strep (1X)

Procedure

A. Receipt of proliferating cells

- 1) Swab down outside of flask with 70% ethanol.
- 2) Equilibrate unopened T25 flask overnight in 37°C, 5% CO₂ humidified incubator to allow cells to recover.

B. Sub-culture

- 1) The next day after receipt, aspirate shipping medium and replace with fresh medium.
- 2) Propagate cells until density reaches 70-80% confluence.
- 3) Aspirate medium.
- 4) Wash cells with warm 1X PBS.
- 5) Add 10mLs of Accutase and return to incubator for 10-15 minutes or until cells detach.
- 6) Immediately remove cells, rinse flask with warm 1X PBS to collect residual cells, and pellet at 500 X g for 5 minutes (4°C).
- 7) Gently re-suspend cell pellet in warm medium.
- 8) Perform 1:2 to 1:5 cell split as needed. Record each subculture event as a passage

C. Maintenance and Generation of Seed Stocks

- 1) Change medium the day after seeding and every 2-3 days thereafter. Use ~50mL of medium per T225 flask.
- 2) Following first or second passage after receipt of cells and with sufficient number of cells to continue maintenance and expansion, a portion of the flasks should be sub-cultured using Accutase as above under “Sub-culture” and the cell pellet resuspended in freezing medium.
- 3) Cells are dispensed into cryovials (2 million cells per 1mL aliquot) and frozen in a -80°C isopropanol bath overnight.
- 4) Cryovials are transferred the next day to liquid nitrogen freezer for long-term storage.

D. Harvest

- 1) Do not use cells that have been passaged more than 8 times.
- 2) Remove cells from flasks according to protocol described above under ‘Sub-culture’.
- 3) Examine viability using Trypan blue staining (SOP TP-7).