



SNL 76/7 Feeder Cells, P14, Mitomycin-C-treated

Product Information

Product Specifications

Catalog Number	Cells per Vial	Number of Vials
ASF-1327	5 x 10 ⁶	1
ASF-1328	5 x 10 ⁶	5

Description

Mouse embryonic stem cells (ESCs) and induce pluripotent stem cells (iPSCs) generally require culture on feeder cells as well as medium containing Leukemia Inhibitory Factor (LIF) to maintain pluripotency and the capacity to self-renew. The SNL 76/7 cell line, derived from a mouse fibroblast STO cell line transformed with neomycin resistance and murine LIF genes, can be used as feeder cells for mouse and human ESC and iPSC culture. The SNL cells are sensitive to HAT selection (hypoxanthine, aminoprotein and thymidine) and negative for HPRT (hypoxanthine guanine phosphoribosyl transferase). The cells are derived from a representative cross section of the Carworth CF-1 colony from Charles River.

Passage

P14

Treatment

Mitomycin C

Shipping

Dry ice

Storage and Stability

Store in liquid nitrogen freezer immediately upon receipt.

Biosafety Level

BSL-1

Safety Precaution

PLEASE READ BEFORE HANDLING ANY FROZEN VIALS. Please wear the appropriate Personal Protection Equipment (lab coat, thermal gloves, safety goggles and a face shield) when handling the cells. Handle the frozen vials with due caution. Please be aware that the following scenario can occur: Liquid nitrogen can leak into the vials when the vials are submerged in liquid nitrogen. Upon thawing, the liquid nitrogen returns to the gas phase, resulting in a dangerous build-up of pressure within the vial. This can result in the vial exploding and expelling not only the vial contents but also the vial cap and plastic fragments of the vial.

Restricted Use

This product is for research use only and not intended for human or animal diagnostic or therapeutic uses.

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Media and Material

Medium (I)

Component	Concentration	Vendor
ESC-Sure™ DMEM		Applied StemCell, #ASM-5001
ESC-Sure™ FBS	10%	Applied StemCell, #ASM-5007
Nonessential amino acids	0.1 mM	Life Technologies, #11140-050
Sodium Pyruvate	1 mM	Life Technologies, #11360-070
L-Glutamine	2 mM	Life Technologies, #25030-164

Suggested plating density (II)

Dish Size	Surface Area*	Working volume	MEF per dish / well
100 mm	55 cm ²	11 - 16.5 ml	1.7 - 2.8 x 10 ⁶
60 mm	21 cm ²	4.2 - 6.3 ml	0.65 – 1.1 x 10 ⁶
35 mm	9 cm ²	1.8 - 2.7 ml	0.27 – 0.45 x 10 ⁶
T25	25 cm ²	5 – 7.5 ml	0.75 - 1.25 x 10 ⁶
T75	75 cm ²	15 – 22.5 ml	2.25 - 3.75 x 10 ⁶
T175	175 cm ²	35 – 52 ml	5.25 - 8.75 x 10 ⁶
6-well	9.5 cm ²	1.9 - 2.9 ml	0.29 – 0.48 x 10 ⁶
12-well	3.8 cm ²	0.8 - 1.2 ml	0.11 – 0.19 x 10 ⁶
24-well	1.9 cm ²	0.4 - 0.6 ml	57,000 – 95,000
48-well	0.95 cm ²	0.2 - 0.3 ml	22,500 – 47,500
96-well	0.32 cm ²	100 - 200 µl	9,600 – 16,000

*Approximate growth surface areas. Numbers can vary between plastic ware from different suppliers

Protocol

1. Remove a vial of frozen cells from liquid nitrogen and place it onto dry ice for 5' before thawing it at 37 °C water bath. As soon as the majority of the content of the vial thawed, transfer it to a conical tube containing 10x volume of pre-warmed medium.
2. Spin at 1000 rpm for 5 min, discard medium, resuspend the cells in growth medium and plate them at an appropriate density in a gelatin-coated tissue-culture dish (generally 25,000-50,000 cells/cm², Appendix III). Optimal density is to be determined by the user for specific applications.