

SV40-GFP Mouse Liver Microvascular Endothelial Cells (SV40-GFP-mLivMVECs)

ORDER INFORMATION

Name of Cells: **Catalogue Number: Product Format:** Cell Number:

SV40t-GFP Mouse Liver Microvascular Endothelial Cells (SV40-GFP-mLivMVECs) cAP-m0006-SV40-GFP Frozen Vial > 5 x 10⁵/vial

General Information

Mouse Liver Microvascular Endothelial Cells (mLivMVECs) were initially isolated from mouse liver tissues of 3-week old C57/Black6 mice and SV40-GFPmLivMVECs were selected from mLivMVECs infected with SV40 and GFP expressing lentiviruses with puromycin. Endothelial Growth Medium (cAP-02) is recommended for the expansion of SV40-GFP-mLivMVECs). And we warranty a minimum of 30 passages of the cells if cultured following the detailed protocol described below).

Characterization:

CD31, VEGFR2, CD144 and CD146 Positive:

SV40-GFP-mhtMVECs are tested negative for common animal pathogen and mycoplasma.

Product Use: SV40-GFP-mHtMVECs are for Research Use Only.

Shipping: Frozen Vial with $> 5 \times 10^5$ cells/vial

Handling of Arriving Cells

When you receive the cells in a frozen vial, you can transfer the vial of cells into a -80°C freezer for short term storage or a liquid nitrogen tank for long term storage. Thaw the cells in a 37°C water bath, and then transfer the cells into a T25 flask Quick Coating Solution (cAP-01) pre-coated flasks as described in details in Subculture Protocol.

Subculture Protocol

Pre-coating of T25 flasks: Add 2ml of the Quick Coating Solution Coating Solution (cAP-01) into one T25 flask and make sure the whole surface of the flask is covered with the Coating Solution; Leave the T25 flask with the Quick Coating Solution for 5 minutes; Aspirate off the Coating Solution after 5 minutes and the flask is ready to be used.

- When Cells are nearly confluent, rinse the cells in T25 flask with 5ml HBSS (Room Temperature, RT) twice. A)
- Add 2ml of Trypsin/EDTA (RT) (cAP-23) into one T25 flask (make sure the whole surface of the T25 flask is covered with Trypsin/EDTA), and B) gently dispose the excessive Trypsin/EDTA solution within 30 seconds with aspiration.
- C) Leave the T25 flask with the cells at RT for 3-5 minute (the cells usually will detach from the surface within 1-2 minutes). You must monitor the cells under microscope and when most of cells become rounded up, hit the flask gently against the bench surface, and the cells will move on the surface of the flask when monitoring under microscope.
- Add 5ml Trypsin Neutralization Buffer (cAP-28) and spin the cells down with 800g for 5 minutes. D)
- Re-suspend the cell pellet with 15ml of full cell culture medium and the cell suspension is transferred directly into 3 pre-coated T25 flasks (5ml E) each, and the cells are sub-cultured at 1:3 ratios)
- F) Change medium every 2-3 days and cells usually become confluent within 5-7days (when split at a 1:3 ratio).

NOTE: The purchase of this product conveys to the buyer the nontransferable right to use the purchased amount of the product and all replicates and derivatives for research purposes conducted by the buyer in his laboratory only (whether the buyer is an academic or for-profit entity). The buyer cannot sell or otherwise transfer (a) this product (b) its components or (c) materials made using this product or its components to a third party or otherwise use this product or its components or materials made using this product or its components to a third party or otherwise use this product or its components or materials made using this product or its components to a third party or otherwise use this product or its components or materials made using this product or its components to a third party or otherwise use this product or its components or materials made using this product or its components or otherwise use this product or its components or materials made using this product or its components or otherwise use this product or its components or materials made using this product or its components or otherwise use this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using this product or its components or materials made using the product or its components or materials made using the product or its components or materials made using the product or its components or materials made using the product or its components or materials made us components for Commercial Purposes Related Products:

| Quick Coating Solution | cAP-01 | 240ml | Angio-Proteomie |
|--|---------|--------|-----------------|
| Endothelial Growth Medium | cAP-02 | 500ml | Angio-Proteomie |
| Endothelial Basal Medium | cAP-02B | 500ml | Angio-Proteomie |
| HBSS w/o Ca ²⁺ , Mg ²⁺ | cAP-11 | 100ml | Angio-Proteomie |
| Cell Freezing Solution (FBS) | cAP-22 | 50ml | Angio-Proteomie |
| Cell Freezing Solution (Non-FBS) | cAP-22B | 50ml | Angio-Proteomie |
| Trypsin/EDTA Solution | cAP-23 | 100ml | Angio-Proteomie |
| Trypsin Neutralization Solution | cAP-28 | 100ml | Angio-Proteomie |
| ITS (100x) | cAP-26 | 10ml | Angio-Proteomie |
| L-Glutamine-MAXIMUM (100x) | cAP-27 | 100ml | Angio-Proteomie |
| Human Plasma Fibronectin Solution | cAP-42 | 1mg/ml | Angio-Proteomie |

THESE PRODUCTS ARE FOR RESEARCH USE ONLY

Caution: Handling human and animal tissue derived products is potentially bio-hazardous. Although each cell strain is tested negative for HIV, HBV and HCV DNA, or pathogens, diagnostic tests are not necessarily 100% accurate; therefore proper precautions must be taken to avoid inadvertent exposure. Always wear gloves and safety glasses when working with these materials. Never mouth pipette. We recommend following the universal procedures for handling products of human origin as the minimum precaution against contamination.