

Plasma Membrane GFP-Tag Human Umbilical Vein Endothelial Cells (PM-GFP-hUVECs)

ORDER INFORMATION

Name of Cells: Plasma Membrane GFP-Tag Human Umbilical Vein Endothelial

Cells (PM-GFP-hUVECs)

Catalogue Number: cAP-0001GFP-PM Product Format: Proliferating culture

Cell Number: > 90% confluent ($>5x10^5$ cells) in T25 flask

General Information

hUVECs (cAP-0001) were isolated from normal human umbilical vein and transfected with Lentiviral particles expressing GFP-Tagged plasma membrane protein marker (neuromodulin) at passage one. Puromycin resistant **PM-GPF-hUVECs** (cAP-0001GFP-PM) were selected and shipped in proliferating culture with >90 confluence (the cells are provided @ passage 3). ENDO-Growth Medium (contains 5% serum and growth supplements, cAP-02) is recommended for cell culture and these cells have an average population doubling levels >18 when cultured following the detailed protocol described below).

Characterization of the cells

Cytoplasmic VWF / Factor VIII: >95% positive by immunofluorescence
Cytoplasmic PECAM1 >95% positive by immunofluorescence
>95% positive by immunofluorescence
>95% positive by immunofluorescence

PM-GFP-hUVECs are negative for HIV-1, HBV, HCV, and mycoplasma.

Product Use: PM-GFP-HUVECs are for research use only.

Shipping: Proliferating culture in T25 flask.

Handling of Arriving Cells

When you receive the cells, leave the flask in 37°C CO2 incubator for 1 hour first, and then replace the transport medium with fresh ENDO-Growth medium. Let the cells grow for 24 hour before subculture.

1. Subculture Protocol:

A) Coating T25 flasks: Add 2ml 0.1% Quick Coating Solution (**cAP-01**) into one T25flask and make sure whole surface of the flask is covered with the coating solution. Five minutes later, dispose Quick Coating Solution by aspiration and the flask is ready to be used (no need for overnight incubation when coated with Quick Coating Solution).

Contact & Ordering Information: Angio-Proteomie, 11 Park Drive, Suite 12, Boston, MA 02215, USA. Tel: 6175492665; Fax: (480) 247-4337, angioproteomie@gmail.com



- B) Rinse the cells in T25 flask with 5ml PBS (**Room Temperature, RT**) twice.
- C) Add 2ml of Trypsin/EDTA (**RT**) (Invitrogen Catalogue number: 25300-062) into T25 flask (make sure the whole surface of the T25 flask is covered with Trypsin/EDTA), and gently dispose the Trypsin/EDTA solution **within 10** seconds with aspiration.
- D) Leave the T25 flask with the cells at **RT** for 1 minute (the cells will normally come off the surface within 1 minute).
- E) Suspend the cells with 20ml of ENDO-Growth medium and the cell suspension is transferred directly into 4 x pre-coated T25 flasks (5ml each, and the cells are subcultured at 1:4 ratio)

(Note: No need spin the cells during the subculture process).

2. Cell culture protocol (proliferating):

- A) Culture medium (ENDO-Growth medium) is changed every 2-3 days.
- B) The cells normally become confluent within 7 days (when split with a ratio of 1:4).

3. Preparation of quiescent cells:

A) ENDO-Basal medium containing 0.5% FBS is used to induce quiescent endothelial cells (after 18-24hours).

Other useful information

Items	Company	Cat #
Quick Coating Solution	Angio-Proteomie	cAP-01
ENDO-Growth medium	Angio-Proteomie	cAP-02
ENDO-Basal medium	Angio-Proteomie	cAP-03
ENDO-Growth Supplement	Angio-Proteomie	cAP-04
PBS	Invitrogen	10010
Trypsin/EDTA	Invitrogen	25300-062

Caution: Handling human derived products is potentially bioharzadous. Although each cell strain testes negative for HIV, HBV and HCV DNA, 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 these materials. Never mouth pipette. We recommend following the universal procedures for handling products of human origin as the minimum precaution against contamination.

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