

Human Dermal Microvascular Endothelial Cells

ORDER INFORMATION

Name of cells: Human Dermal Microvascular Endothelial Cells (**HDMEnds**)
Catalogue number: **cAP-0005n**
Product format: Proliferating culture
Cell number: > 90% confluent in T25 flask

General Information

HDMEnds (**cAP-0005n**) are isolated from normal neonatal human skin tissue. The cells are shipped in proliferating culture with >90 confluence (the cells are provided @ passage 3). EGM™-2 MV full medium (contains 5% serum and growth supplements, LONZA, CC-3202) is recommended for cell culture and these cells have an average additional population doubling levels >**16** when cultured following the detailed protocol described below).

Characterization of the cells

Cytoplasmic VWF / Factor VIII:	> 95% positive by immunofluorescence
Cytoplasmic uptake of Di-I-Ac-LDL:	> 95% positive by immunofluorescence
Cytoplasmic PECAM1:	> 95% positive by immunofluorescence

HDMEnds are negative for HIV-1, HBV, HCV, and mycoplasma.

Product Use

HDMEnds 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 EGM™-2 MV full medium. Let the cells grow for 24 hour before subculture.

1. Subculture Protocol:

- A) Coating T25 flasks: Add 2ml of Quick Coating Solution (**cAP-01**) into one T25 flask 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).
- B) Rinse the cells in T25 flask with 5ml DPBS (**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 Trpsin/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 EGMTM-2 MV full medium and the cell suspension is transferred directly into 4 pre-coated T25 flasks (5 ml each, and the cells are subcultured at 1:4 ratio)

(Note: Never spin the cells during the subculture process).

2. Cell culture protocol (proliferating):

- A) Culture medium (EGMTM-2 MV full medium) is changed every 2 days.
- B) The cells normally become confluent within 7 days (when split at a 1:4 ratio).

3. Preparation of quiescent cells:

- A) EGMTM-2MV medium containing 0.5% FBS is used to induce quiescent endothelial cells (after 18-24hours).

Caution: Handling human derived products is potentially biohazardous. Although each cell strain tests 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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