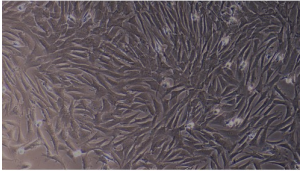
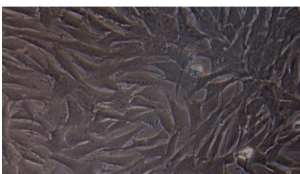


Goat Aortic Endothelial Cells (AEC)

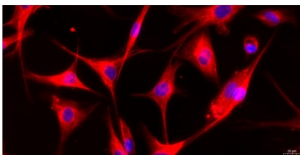
Catalogue No.: abx700022



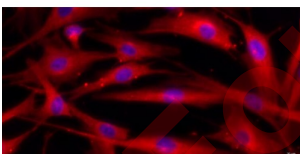
Morphology of Caprine Aortic Endothelial Cells (Optical microscope, 100X).



Morphology of Caprine Aortic Endothelial Cells (Optical microscope, 200X).



Immunofluorescence identification of Coagulation Factor VIII (FVIII) (400X).



Immunofluorescence identification of Von Willebrand Factor (400X).

Goat Aortic Endothelial Cells (AEC) are Adherent Goat Endothelial Cells from Goat Aorta.

Target: Aortic Endothelial Cells (AEC)

Origin: Goat

Host: Goat

Datasheet

Version: 2.0.0
Revision date: 24 Apr 2025



Purity: Negative for HIV-1, HBV, HCV, mycoplasma, bacteria, yeast and fungi.

Storage: Shipped at -70 °C. Upon receipt, store in liquid nitrogen (-196 °C). Avoid repeated freeze/thaw cycles.

Validity: 12 months.

Buffer: Contains 90% FBS and 10% DMSO.

Biological Activity: Cell activity: > 85% (viability by Trypan Blue exclusion)

Note: THIS PRODUCT IS FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC, THERAPEUTIC OR COSMETIC PROCEDURES. NOT FOR HUMAN OR ANIMAL CONSUMPTION.
This product is shipped with dry ice.

Directions for use: Recommended Cell Culture Conditions:

DMEM/F12 + 5% FBS + Goat Endothelial Cell Growth Supplement + 1% Penicillin-Streptomycin Solution @ 37 °C, 95% air, 5% CO₂.

Cell Recovery: Thaw cells in a 37 °C water bath with shaking until the mixture has dissolved. Transfer to a centrifuge tube and add culture medium (see Recommended Cell Culture Conditions above) at a volume 3-5 times the volume of the cells. Centrifuge at 1000 RPM for 5 minutes and discard the supernatant. Transfer to a T25 flask for culture.

Suggested Cell Passage Procedure: Cells should be 85-95% confluent before cell passage is carried out.

1. Discard the medium and wash with PBS 1-2 times.
2. Add 1 ml of Trypsin at 37 °C, then observe the cells under a microscope.
3. When the cells appear retracted and rounded, gently tap the culture flask to detach the cells. Stop the trypsinization by adding 2 ml of culture medium containing 10% serum.
4. Add fresh medium to resuspend the cells. The recommended ratio of primary cells is 1/2. Pipette to obtain a single cell suspension.