

OPM-VERO SFM6C3

Serum-free Cell Culture Medium

— For Vaccine Production

OPM-VERO SFM6C3 is a serum-free cell culture medium designed for adherent culture of Vero cells and contains L-Glutamine. This medium supports the production of human vaccines such as COVID-19 vaccine, polio vaccine, smallpox vaccine, rabies vaccine, Japanese B encephalitis vaccine, rotavirus vaccine, enterovirus Type 71 vaccine and oncolytic virus, and the production of veterinary vaccines such as porcine diarrhea virus vaccine and small ruminant vaccine, etc.

Application

OPM-VERO SFM6C3 is intended for large scale manufacturing of therapeutic biomolecules, as well as for research purposes, but not for human or any therapeutic use.

Storage & Transportation

Store at 2~8°C, dark and dry
Ship at Room temperature (Liquid), Blue ice (Dry powder)

Shelf Life

OPM-VERO SFM6C3 Medium Liquid: 12 months
OPM-VERO SFM6C3 Dry Powder: 24 months

Reconstitution Method for Dry Powder

1. Measure out 90% of final required volume of purified water intended for cell culture use, e.g. WFI. Recommended water temperature is 25~35°C (minimum final volume \geq 1L) .
2. Slowly add dry powder medium at 19.86 g/L while stirring.
3. Adjust pH to 7.2 with 1N NaOH or 1N HCl.
4. Add cell culture grade purified water to 100% final volume.
5. Continue to stir for 10 minutes. Sterile filter using a membrane filter with a pore size of 0.22 micron.

Quality Specifications

Specifications	OPM-VERO SFM6C3 Medium	OPM-VERO SFM6C3 DPM
Appearance	Red clear liquid	Off-white or light yellow, homogenous dry powder
pH	7.2~7.7	7.2~7.7
Osmolality (mOsm/kg)	285~315	285~315
Solubility	—	Good by following the reconstitution instructions
Endotoxin (EU/mL)	<1.0	<1.0
Sterility test	Negative	—

Cell Culture Conditions

36°C~38°C, 5~8%CO₂

Cell Recovery

1. Rapidly thaw (<2 min) a vial of frozen cells in a 37 °C water bath.
2. Transfer the entire contents aseptically into a shake flask containing 15 mL mL prewarmed OPM-VERO SFM6C3 cell culture medium without antibiotics.
3. Incubate at 37 °C in a humidified atmosphere of 5%~8% CO₂ in air.
4. Passage cells when the confluency reaches 80-100%.

Cell Culture Passaging

1. Aspirate medium from cell monolayer and rinse the flask three times with prewarmed DPBS without Ca²⁺ or Mg²⁺.
2. Add 5 mL 0.25% Trypsin-EDTA to the flask and incubate until cells have detached (~2-5 minutes at room temperature).
3. Add prewarmed 9 mL OPM-VERO SFM6C3 medium to stop the dissociation reaction.
Note: If using 0.25% Trypsin-EDTA, addition of 500 µg/mL Soybean Trypsin Inhibitor is required.
4. Centrifuge cell suspension at 100 ×g for 5 minutes and discard the supernatant.
5. Resuspend the cell pellet in 10mL OPM-VERO SFM6C3 medium.
6. Seed flasks at 1~5 × 10⁴ viable cells/cm².
7. Incubate cells in a humidified 37 °C incubator with 5-8% CO₂ until the confluency reaches 80-100%.

Cryopreservation

1. Harvest the desired quantity of cells in log phase of growth with viability over 90%. Save the conditioned medium to prepare freezing medium.
2. Determine VCD to ensure that the final cell density is 1~5×10⁶ cells/ mL.
3. Harvest cells when the confluency reaches 80%.
4. Prepare the freezing medium consisting of 92.5% OPM-VERO SFM6C3 (50:50 ratio of fresh to conditioned media) +7.5% DMSO, and let the freezing medium cool down to 4°C.
5. Harvest cells and centrifuge at 100 × g for 5-10 minutes. Resuspend the cell pellet in the pre-determined volume of 4°C freezing medium.
6. Dispense aliquots of this suspension into cryo-vials ((1.5~2 mL per vial).
7. Place the vials in a cryo-box or a controlled rate freezing apparatus following standard procedures (1°C decrease per minute).
8. For long-term storage, transfer the vials to liquid nitrogen.

Order Information

Cell Culture Media

Name	Cat No.	Type	Volume
OPM-VERO SFM6C3 Medium	C216306	Liquid	500ml /1000 ml
OPM-VERO SFM6C3 DPM	C210863	Dry powder	10L/50L/100L

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