

CRYOVIVE™ 5% or 2% DMSO

STERILE, CHEMICALLY DEFINED, ANIMAL ORIGIN-FREE (cdAOF) CELL CRYOPRESERVATION MEDIUM

50 mls Liquid Per Bottle. Contains 5 or 2 percent DMSO. Product number: AVBP-001 (5% DMSO) and Product number: AVBP-090 (2% DMSO), respectively.

Customer Support: Call +1 (206) 823-5895, email info@avantbio.com or visit www.avantbio.com.

PRODUCT DESCRIPTION

CRYOVIVE™ is a physiologically-buffered, sterile, defined, animal origin–free, serum–free liquid cryopreservation medium containing 5% or 2% dimethylsulfoxide (DMSO). CRYOVIVE™ contains only USP-certified and other high-quality ingredients. CRYOVIVE™ does not contain undefined products, or any animal or human-derived products.

INTENDED USE

CRYOVIVE™ is intended for low temperature freezing and storage of mammalian cells or tissue in the liquid nitrogen (LN₂) vapor-phase (–196 °C). This product is for laboratory research use only and is not intended for use in animals, humans, or for use in diagnostic procedures. CAUTION: CONTAINS DMSO. Take appropriate precautions regarding the proper use and disposal of DMSO containing solutions. When handling this product, wear of protective clothing and eye wear. Dispose of CRYOVIVE™ properly.

STORAGE AND STABILITY

CRYOVIVE™ should be stored at – 20 °C and thawed before use. Upon thawing CRYOVIVE should be

mixed by swirling the bottle, and then stored at 2–8 $^{\circ}$ C, in the dark thereafter. CRYOVIVE $^{\text{TM}}$ should not be freeze thawed. Protect CRYOVIVE $^{\text{TM}}$ from excessive heat and UV or visible light. CRYOVIVE is guaranteed for 6 months at 2–8 $^{\circ}$ C and 18 months at – 20 $^{\circ}$ C.

INSTRUCTIONS FOR USE

- 1. Store CRYOVIVE™ at 2–8 °C until it is ready for use. To help maintain sterility, wipe down outside of bottle in a sterile field with 70% ethanol before removing the lid. The contents are sterile. If the seal has been broken or tampered, do not use. Gently mix (swirl) CRYOVIVE™ medium prior to use.
- **2.** If your cells are adherent, enzymatically remove the cells from their subculture surface using dAOF cell dissociation reagents (TrypLE $^{\rm TM}$ Select, Life Technologies cat. no. 12563-029). Collect and pellet the cells using centrifugation. After removing the supernatant, resuspend the cell pellet in cold (2–8 °C) CRYOVIVE $^{\rm TM}$ medium and store cells temporarily at 2–8 °C (e.g., on ice). If required, rapidly count viable and nonviable cells, and adjust volume to obtain the desired cell concentration, typically 4 x 10 $^{\rm 5}$ to 4 x 10 $^{\rm 6}$ cells/ml.
- **3**. Quickly distribute the cell suspension (typically 0.5–2 mls) into cooled cryopreservation vials and immediately cool each vial of cells to 2–8 °C.
- **4.** Cryopreserve the cells by controlled temperature reduction (descent) to the LN_2 vapor–phase temperature. It is recommended that you follow a standard slow–rate (e.g., 1 °C/min), controlled cooling protocol to the LN_2 vapor-phase temperature. If precise controlled rate cooling is not possible in your laboratory, the cells can be cryopreserved by placement of the cells in a controlled cooling enclosure (e.g., Nalgene-Nunc Mr. Frosty) preequilibrated at 4°C that is placed at -80°C or -78.5 °C for 3-4 hrs., which is then followed by storage of the cells in the vapor–phase of LN_2 .
- **5.** Following cryopreservation in CRYOVIVE™, for recovery, the cells should be rapidly warmed with gentle shaking in a 37 °C water bath, just until all visible ice melts (2-3 minutes maximum). Do not allow cells to incubate for prolonged periods of time at 37 °C or room temperature. **Immediately dilute cells (2–fold) in cell culture medium** and temporarily store cells at 2–8 °C (e.g., on ice) before placing into subculture. If required, rapidly count viable and non-viable cells (e.g., trypan blue exclusion) while the cells on are on ice and subculture cells at the desired density using your individual protocol. When placed into subculture, CRYOVIVE™ cell suspensions should be diluted at least a 75-fold.





AVANTBIO LIMITED USE AGREEMENT

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