# BloodStor® 100

### **Biopreservation reagent for cells and tissues**



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Catalog #	07951	50 mL	INFO@STEMICELL.COM • TECHSUPPORT@STEMICELL.COM	
-	07939	100 mL	FOR GLOBAL CONTACT DETAILS VISIT OUR WEBSITE	
	07938	5 x 100 mL		

## Product Description

BloodStor® is a series of generic cGMP freezing media products used to cryopreserve stem cells and other cells isolated from umbilical cord blood, peripheral blood, bone marrow, and other biologics. BloodStor® 100 contains 100% dimethyl sulfoxide (DMSO) USP.

- Sterile, USP vial
- USP grade/highest-quality components
- Filled under cGMP
- Sterility and endotoxin testing

### **Properties**

Storage: Store at 20 - 30°C.

Shelf Life: Stable until expiry date (EXP) on label. Protect product from prolonged exposure to light.

Contains: • 100% Dimethyl sulfoxide (DMSO)

Please refer to the Safety Data Sheet (SDS) for hazard information.

BloodStor® 100 will solidify at temperatures below 18°C. If BloodStor® 100 is solidified upon receipt, allow to thaw at room temperature (18 - 25°C) and mix gently. Product performance will not be affected.

# Handling / Directions For Use

### FREEZING

- 1. Wipe down the outside of the cryopreservation reagent container and desired culture medium bottle with 70% ethanol or isopropanol before opening.
- 2. Prepare desired crypreservation medium (e.g. 1 part BloodStor® 100 in 9 parts culture medium) and keep cold (2 8°C).
- 3. Obtain a cell suspension using a cell-specific protocol and centrifuge cells to obtain a cell pellet.
- 4. Carefully remove the supernatant with a pipette, leaving a small amount of medium to ensure the cell pellet is not disturbed. Resuspend the cell pellet by gently flicking the tube.
- 5. Add cold (2 8°C) cryopreservation medium, mix thoroughly, and transfer the suspension to a cryovial.
- Freeze cells using a standard slow rate-controlled cooling protocol (approximately -1°C/minute) or an isopropanol freezing container and store at liquid nitrogen temperature (-135°C).

NOTE: Long-term storage at -80°C is not recommended.

### THAWING

- 1. Warm medium of choice in a 37°C water bath.
- 2. Wipe the outside of the vial of cells with 70% ethanol or isopropanol.
- 3. In a biosafety hood, twist the cap a quarter-turn to relieve internal pressure and then retighten.
- 4. Quickly thaw cells in a 37°C water bath by gently shaking the vial. Remove the vial when a small frozen cell pellet remains. Do not vortex cells.
- 5. Wipe the outside of the vial with 70% ethanol or isopropanol.
- 6. Dilute in warmed medium of choice at a ratio of 1 part sample to 9 parts medium.
- 7. Centrifuge the cell suspension at 300 x g for 10 minutes at room temperature (15  $25^{\circ}$ C).
- 8. Carefully remove the supernatant with a pipette, leaving a small amount of medium to ensure the cell pellet is not disturbed. Resuspend the cell pellet by gently flicking the tube.
- 9. Gently add medium to the tube.
- 10. Repeat steps 7 and 8.



THIS PRODUCT IS MANUFACTURED UNDER A cGMP QUALITY MANAGEMENT SYSTEM COMPLIANT TO 21 CFR 820.

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