## Cryopreservation

Cleavage stage embryos

Directions for supplementation of un-supplemented G-Series<sup>™</sup> media can be found in the G-Series Manual on www.vitrolife.com. Once supplemented, the media should be used as the G-Series PLUS media described below.

Day 2 or 3 Freezing: FreezeKit <sup>™</sup> Cleave 1. Prepare dishes with 0.5-1.0	4. Seal the straws and place into the freezing machine at room temperature and commence the freezing program, see G-Series manual.	Thawing: ThawKit <sup>™</sup> Cleave 1. Prepare dishes with 0.5-1.0 ml of TS1, TS2 and ES into respec- tive dishes at	4. Open the straw using aseptic technique and gently expel the embryos into TS1 and leave at ambient temp 5 min	For embryo transfer, see separate Embryo transfer protocol
ml each of Equilibration solu- tion (ES) and Freezing solution (FS) at ambient temp	5. Manually seed the straws with liquid nitrogen (LN <sub>2</sub> ) held forceps close to the cotton plug at	ambient temp and atmosphere	5. Move the embryos into TS2 and expose for	
ambient temp and atmosphere	-6°C	One dish per patient and	ambient temp 5 min	
One dish per patient and solution		solution2. Remove the straw from liquid nitrogen and expose to air	6. Move the embryos into ES and expose for	
2. Place the embryos in ES and rinse properly. The embryos can stay in ES for 10 minutes.		30 sec	ambient temp and atmosphere 5 min	
3. Transfer the embryos into the FS and load the embryos into		Place the straw in a water bath at	7. Rinse embryos properly in G-1	
straws immediately. Total expo- sure time in FS is	6. Continue the freezing program.	30°C 45 sec	PLUS™ or G-2 PLUS™ depending on developmental stage and culture until transfer at	
10 min	7. Plunge the straw into liquid nitrogen when the freezing pro- gram has ended and arrange for storage at	<b>3.</b> Remove the straw and wipe it carefully.	37°C 6 % CO2	
Pre-rinse all utensils, includ- ing dishes and pipettes when applicable with G-RINSE™	-196°C			

