

# Sperm preparation

## Density gradient centrifugation method

### Day -1

The day before oocyte pick-up



Pre-equilibrate Gx - IVF at

**37°C 6 % CO<sub>2</sub>  
overnight**

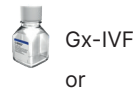
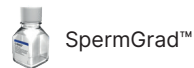


Pre-rinse all utensils, including tubes and dishes, with G-RINSE.

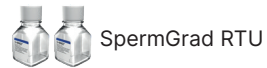
### Day 0

#### 1. Assess the semen sample

#### 2. Prepare gradient solutions



or



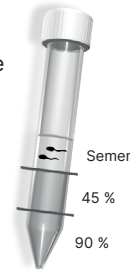
If you use SpermGrad RTU go to paragraph 3.

#### Dilution of SpermGrad

Mix SpermGrad with Gx-IVF in separate rinsed tubes to obtain 90 % and 45 % stock solutions. For 90 % stock solution, mix 9.0 mL SpermGrad with 1.0 mL Gx-IVF and for 45 % stock solution, mix 4.5 mL SpermGrad with 5.5 mL Gx-IVF.

#### 3. Prepare gradients

Pipette 1.5 mL of the 90 % solution into the rinsed tube first and then slowly pipette 1.5 mL of the 45 % solution on top of it. Finally, 1.0 mL of the semen is layered on the top.



Make up 2-4 gradient tubes. Before use, allow the stock solutions to warm to ambient temperature.

#### 4. Centrifuge the gradients at

**300-600g 10-20 min**

#### 5. Wash I

Remove the two top layers. Transfer the pellets to new rinsed tubes and re-suspend with 5 mL equilibrated Gx-IVF and centrifuge again at



**300-600g 10 min**

#### 6. Wash II

Aspirate and discard the supernatant. Transfer the pellets to new rinsed tubes and re-suspend with 5 mL equilibrated Gx-IVF and centrifuge again at



**300-600g 10 min**

#### 7. Assess sperm preparation

Aspirate and discard the supernatants. Combine all pellets in a new rinsed tube and re-suspend in 0.5-1.0 mL of equilibrated Gx-IVF depending on sample quality.



Determine motility and concentration of spermatozoa in the washed sample.

#### 8. Dilution

Dilute with equilibrated Gx-IVF to a final concentration of 75,000-200,000 motile sperms/mL.

#### 9. Preparation of insemination dishes

Prepare rinsed insemination centre well dishes with 0.5-1.0 mL of sperm solution and pre-equilibrate.



If oil overlay is used, droplets of at least 100 µL volume are recommended. Equilibrate the dishes at

**37°C 6 % CO<sub>2</sub> >2 h**

#### 10. Insemination

Transfer the oocytes to the insemination centre well dishes and leave at

**37°C 6 % CO<sub>2</sub>  
overnight**

**Alternatively:** Add equilibrated sperm suspension to equilibrated centre well dishes with the oocytes already present.