# SpermCryo<sup>M</sup> All-round

# PRODUCT INFORMATION

# SpermCryo<sup>TM</sup> All-round

Catalogue no.	
SPCA-0005	5 ml
SPCA-0010	10 ml
SPCA-0020	20 ml

#### Application

SpermCryo<sup>™</sup> All-round is a four times concentrated ready-to-use HEPES buffered cryopreservation medium for freezing human semen at ultra low temperatures (-196°C) in liquid nitrogen.

#### Composition

SpermCryo<sup>™</sup> All-round is a HEPES buffered freezing medium for human sperm. It contains 26,7% glycerol in order to protect the sperm from damage due to the freezing process and it contains 0,4% HSA.

#### Material not included

- Sperm freezing straws (e.g. CBS high security sperm straws)
- Freezing tank with liquid nitrog
- LAF Bench (ISO Class 5)
- Sterile pipettes

#### **Quality Control**

- pH: (20 7,90) (Release criteria: 70 – 7,60)
- Endotoxin: < 25 EU/ml
- Sterility: steril SAL 10<sup>-3</sup>
- Sprin Survival Test ≥ 80% structured by Exposure of untraited sperm to the test
- Chemical co
- Use of Ph Eur or OSP grade products if available
- Not MEA tested
- Certificate of Analysis available apon request

### Sterility

SpermCryo<sup>TM</sup> All-round is sterilized by sterile filtration.

# STERILE A

#### Precautions and warnings

Always work under hygienic conditions (LAF-bench, ISO Class 5) to avoid possible contamination.

Always wear protective clothing when working with specimens.

Handle specimens as if capable of transmitting HIV or hepatitis.

Standard measures to prevent infections resulting from the use of medicinal products prepared from human blood or plasma include selection of donors, screening of individual donations and plasma pools for specific markers of infection and the inclusion of effective manufacturing steps for the inactivation/removal of viruses. Despite this, when medicinal products prepared from human blood or plasma are administered, the possibility of transmitting infective agents cannot be completely excluded. This also applies to unknown or emerging viruses and other pathogens. There are no reports of proven virus transmissions with albumin manufactured to European Pharmacopoeia specifications by established processes.

The above media do not contain antibiotics.

# Pre-use checks

Domesuse if the seal on the both eist roken or open when the group of is delivered.

Do not use if the product shows any sgns of microbial contamination or becomes cloudy.

# Storage Condi

tore refrigerated (2° - 8°C

not use after expir

Do not freeze before use

Keep away from sunlight.

After operation the container, do not use the product onger than the Stabile conditions must be maintained and must auct must be stored at 2° - 8° C. the leafter transport (up to 5 days) at elevated to maratures & 37°C).

## Technical Support



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**C€** 

Distributor:	

# SpermCryo<sup>TM</sup> All-round

# Instructions for use

### **Sperm Preparation**

 $\overline{SpermCryo^{TM}}$  All-round can be used in combination with  $\overline{SpermFilter}^{\otimes}$  ready-to-use gradients.

# Before freezing

It is recommended to concentrate the sperm before freezing in case of very low sperm concentrations. This may increase the sperm quality after thawing and will reduce the number of straws to be frozen.

#### After thawing

In case needed, use sperm preparation techniques after thawing to eliminate dead sperm cells and debris. Dilute the concentrated sperm in a washing medium to your use.

# **Method**

Ensure that all media is well mixed before use.

#### Freezing

- Allow the semen to aquefy at room temperature for 30 minutes.
- Add 1 part SpermCryo M All-round to
- Acta Sparn Cryo<sup>TM</sup> All-round dropwise while gently swifting in sample.

  Caution: To accompled suck make sure Sperm Cryo<sup>TM</sup> All-round sparn temperature.
- Leave the sample for 10 minutes at room temperature for equilibration.
- Suck the mixture into the freezing straws, leaving approximately 1.5 cm of air at the end of the straw. Seal the straws.
- Dry each straw with a linen free cloth and shake in order to move the air bubble to the centre of the straw.
- Freeze the straws vertically just above the liquid nitrogen level for 15 minutes.
- · Store the straws in liquid nitrogen.

#### **Thawing**

- Retrieve the required straws from the liquid nitrogen.
- Place the straws in tap water for 5 minutes.

a container (e.g. test tube) and tap straw against the side of the container in order to allow container available of the mixture into the

- idute the concentrated sperm in a suitable inscrination medium, at least 3 ml per 0.5 ml section pix thoroughly.
- Centralinge at 300-350g for 15 minutes.
- Resuspend the pellet in a phtable insemination medium (e.g. SpennWash<sup>6</sup>).

SYMBOL	MEANING
REF	<ul> <li>Catalogue number</li> </ul>
LOT	Batch code
$\overline{\underline{\Sigma}}$	Use by (expiry date)
X***	Temperature limitations
1	Sterile medical device processed using aseptic technique (filtration)
[]i	Consult instructions for use
<b>C €</b> 0344	CE mark

#### Bibliography

- 1 Mahadevan M, Trounson AD. Effect of cryoprotective media and dilution methods on the preservation of human spermatozoa. Andrologia, 1983; 15: 355-66.
- 2 Mahadevan M, Trounson AD, Leeton JF. Successful use of human semen cryobanking for in vitro fertilization, Fertil Steril, 1983; 15: 355-66.
- 3 Brotherton J. Cryopreservation of human semen. Archives of Andrology, 1990; 25: 181-95.

Ref. nr. 319/1

- 4 Kobayashi T, Kaneko S, Hara I, Park YJ, et al. Concentrating human sperm before cryopreservation. Andrologia, 1991; 23: 25-8.
- 5 Graczykowski JW, Siegel MS. Influence of sperm processing on the fertilizing capacity and recovery of motile sperm from thawed human semen. Archives of Andrology, 1991; 26: 155-61.

