

**Manufacturer:**  
Sparmed ApS/CVR.No.: 30898575  
Ryttermarken 2, 3520 Farum, Denmark



**ID:** COA-07603

# Certificate of Analysis

**Date of issue:** 19.02.2016  
**Product ID:** Oosafe® Plasticware: OOPW-HD10  
**LOT No.:** 07603  
**Expiry Date:** 09/2020  
**Storage conditions:** 20<sup>0</sup>C, dry room, no exposal to sun-light  
**Quality Assurance:**

Proven non-embryotoxic by Mouse Embryo Assay Test. 100% embryo development to the expanded blastocyst stage within 96hours. **PASS**  
Proved stable human sperm motility: ≥70% sperm motility after 24hours proven. **PASS**  
Proven non-toxic by Limulus Amebocyte Lysate (LAL) test. Pass criteria <0.03 EU/device **PASS**  
Proven RNase DNase test FREE- **PASS**  
Sterilization by gamma irradiation. Delivered irradiation dose: 8.6kGy-9.5kGy. Specified irradiation dose: 8.0kGy-10.0kG- **PASS**

Quality control according to the ISO 13485:2012

**Final approval:**

A handwritten signature in blue ink, appearing to read 'Katrine Nobel', written over a horizontal line.

Katrine Nobel  
Quality Control Manager

**Stamp:**





SparMED Aps  
 Toppevadvej 34-38  
 DK-3660  
 Stenlose, Denmark



140 Hale Street  
 Haverhill, MA 01830  
[qc@embryotech.com](mailto:qc@embryotech.com)

ELI Accession Number: SPAR-3841-1115

Date of completion: 11-17-2015

Lot numbers: 07603, 07506

Reference numbers: OOPW-FW04, OOPW-HD10  
 OOPW-CT01, OOPW-SC01

**Description of test article(s):**

Oosafe® 4 Well Dish, 100mm Dish, Centrifuge Tube and Sperm Collection Cup

**Assay system requested by customer:** 1mL of culture medium was placed in each of the test articles (3) (OOPW-HD10, OOPW-CT01, OOPW-SC01) and incubated at 37°C for 30-minutes. Post incubation the culture medium was extracted from each test article and pooled. 0.5mL of the extracted culture medium was expelled into each well of the test article (4-well NonTreated dish); 1-cell mouse embryos were then added to each well of the test article and cultured for 96-hours.

**Control assay method and results:** 15 1-cell (B6C3F1 X B6D2F1) embryos were cultured in 0.5mL drops in a non-treated 4-Well Dish using culture medium;

15 / 15 (100 %)  
 15 / 15 (100 %)

1-cell to 2-cell within 24 hr  
 1-cell to expanded blastocyst within 96 hr

*For a valid assay, Embryotech™ requires at least 70% of 1-cell stage control embryos to develop to expanded blastocyst within 96-hours.*

**Test assay method and results:** 21 1-cell (B6C3F1 X B6D2F1) embryos were cultured in one of the test articles using the extracted culture medium:

21 / 21 (100 %)  
 19 / 21 ( 90 %)

1-cell to 2-cell within 24 hr  
 1-cell to expanded blastocyst within 96 hr

**Summary of observations:** All test and control embryos were selected randomly from a common pool of freshly collected embryos and were cultured in the same incubator at 37°C and 5.0% CO<sub>2</sub>. 100 percent of the control embryos developed to the expanded blastocyst stage within 96-hours, 90 percent of the embryos cultured in the test article developed to the expanded blastocyst stage within 96-hours.

signature  
 Study Director

date

11-17-2015

signature  
 Quality Reviewer

date

11-18-2015



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ELI Accession Number: S2291-1115SPAR

Date of completion: 11-17-2015

Lot number: 07506  
 Lot number: 07603

Order numbers: OOPW-IC01, OOPW-SC01, OOPW-ST01  
 Order numbers: OOPW-CT01, OOPW-CW05, OOPW-FW04,  
 OOPW-HD10, OOPW-IC03, OOPW-TF02,  
 OOPW-TF03

Description of test article(s): Oosafe® ICSI Dish, Sperm Collection Cup, 60mm Dish, Centrifuge Tube, Center Well Dish, 4 Well Dish-NonTreated, 100mm Dish, ICSI/IMSI Dish, 35mm Dish


Assay system requested by customer: 1mL of sperm wash medium was added to the test articles (10 test articles pooled) for 30 minutes. Post incubation the sperm wash medium from the test articles was pooled and 200µl of the medium was added to the 4-well with the sperm for 24-hour incubation.

Test Assay method and results: A non-frozen donor sperm specimen was obtained and used for this assay. The sperm was prepared and the motile fraction separated using a sperm gradient and centrifuge cycle. The motility was noted at the beginning of the assay and again at 24-hours using a measure depth Makler Chamber System. Analyses were performed in sequence each time, with no more than 5-minutes between the test and the control samples.

**Results:**

Test method:	Specification	Initial	Result % 24hr	SMI Value	Pass/Fail
SOP/TSG/ELI/008					
Test Article	SMI ≥ 0.75	90%	90%	1.00	Pass
Control	≥ 70%	90%	90%	N/A	Pass

Summary of observations: The motility remained consistent in the test article extract and control while in an incubator atmosphere of 32°C and 5% CO<sub>2</sub>. Neither the test nor the control showed any signs that the motility was affected during the course of the assay.

  
 signature  
 Study Director

11-18-2015  
 date

  
 signature  
 Quality Reviewer

11-18-2015  
 date





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ELI Accession Number: E6685-1115SPAR

Date of completion: 11-13-2015

Lot number: 07506  
 Lot number: 07603

Order numbers: OOPW-IC01, OOPW-SC01, OOPW-ST01  
 Order numbers: OOPW-CT01, OOPW-CW05, OOPW-FW04,  
 OOPW-HD10, OOPW-IC03, OOPW-TF02,  
 OOPW-TF03

Description of test article(s): Cosafe® ICSI Dish, Sperm Collection Cup, 60mm Dish, Centrifuge Tube, Center Well Dish, 4 Well Dish-NonTreated, 100mm Dish, ICSI/IMSI Dish, 35mm Dish


Assay system requested by customer: Endotoxin titer and interference screening using the Gel-Clot method.

LAL lot number: 513-05-647  
 CSE lot number: 143  
 LRW lot number: AYE162370

Sensitivity ( $\lambda$ ) = 0.03 EU/mL

Control Standard Series			Test Sample Dilutions	NPC		PPC	
2 $\lambda$ .06	+	+	Undiluted	-	-	+	+
$\lambda$ .03	+	+	1:2	-	-	+	+
$\frac{1}{2}\lambda$ .015	-	-	1:4	-	-	+	+
$\frac{1}{4}\lambda$ .0075	-	-	1:8	-	-	+	+
NWC	-	-	1:16	-	-	+	+

Summary of observations: The error for the Gel-Clot assay is +/- one two-fold dilution. The test article in this assay indicates an Endotoxin Concentration of < 0.03 EU/device.

  
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 Study Director

11-18-2015  
 date

  
 signature  
 Quality Reviewer

11-19-2015  
 date

Amended: 11-18-2015

## RNase Test Data and Results

Date: 12/07/2015  
Company: Sparmed ApS  
Date received: 12/01/2015

Project #: 113288A  
Contact: Onur Ozturk  
Technician: Chase Wong

PO#: 130214-1  
Phone: 45- 39 40 2503

Products tested:	Product code:	Lot #:
Oosafe 4-WELL DISH	OOPW-FW04	07603
Oosafe 35 MM DISH	OOPW-TF03	07603
Oosafe 60MM DISH	OOPW-ST03	07603
Oosafe CENTER WELL	OOPW-CW05	07603
Oosafe ICSI DISH	OOPW-IC03	07603
Oosafe 100 MM DISH	OOPW-HD10	07603
Oosafe 60MM DISH	OOPW-ST01	07506
Oosafe 35 MM DISH	OOPW-TF02	07603
Oosafe 15 mL tube	OOPW-CT01	07603
Oosafe ICSI DISH	OOPW-IC01	07506

### Extraction:

Extract solution: DEPC Treated Water  
Lot #: DW15A6  
Volume: 1000µl

Number of test items exposed to extract solution: 10  
Special extraction instructions: Tested products according to extraction protocol #155.

### Procedure and Controls:

RNA: 6.0 kb Poly (A)-tailed  
RNA lot #: 1310020  
Salts: MgCl<sub>2</sub> and NaCl  
Salt lot #: S15G2

RNA standard pool: 3µl of RNA + 12µl Salts.  
Volume of each standard reaction: 5µl  
Volume of extract added to the standard: 10µl  
Total volume: 15µl

Negative Control (-): RNA and salt standards with 10 µl of unexposed extract solution added

Positive Control (+): RNA and salt standards with 10 µl of extract solution exposed to RNase from a tip touched by un-gloved hands

Incubation periods: 1 hr @ 37°C, followed by 5 minutes at 65°C

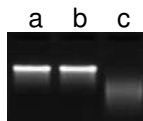
### Gel Electrophoresis:

2µl gel loading dye + 15µl reaction is loaded on a 1.2% agarose in ½ X TAE gel

Gel loading dye lot #: DD006

Electrophoresis: 20 minutes @ 80 volts

### Photographic Results:



Lane (a) product samples, (b) unexposed RNA standard as a negative control, (c) RNA standard exposed to RNase as a positive control.

### Conclusions:

There is no visible degradation in lanes (a) and (b). Lane (a) represents the product samples and lane (b) represents the negative control. Lane (c), which represents the RNA standard, exposed to RNase as a positive control shows degradation of the RNA. The results suggest that the product sample is free of detectable RNase contamination.

### Recommendations:

Based on this experimental procedure, we can show a definite risk of RNase contamination if your product is touched by un-gloved hands. We suggest that all operations are monitored and personnel are instructed in the importance of avoiding RNase contamination.



Chase Wong  
Lab Technician

12/08/2015  
Date



Carl Tsang  
Q.A.

12/08/2015  
Date



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### RNase FREE CERTIFICATE OF ANALYSIS

12/08/2015

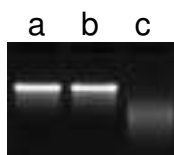
The following sample obtained from **Sparmed ApS** on **12/01/2015** is free of any detectable RNase contamination.

<u>PRODUCTS TESTED</u>	<u>PRODUCT CODE</u>	<u>LOT NUMBER</u>
Oosafe 4-WELL DISH	OOPW-FW04	07603
Oosafe 35 MM DISH	OOPW-TF03	07603
Oosafe 60MM DISH	OOPW-ST03	07603
Oosafe CENTER WELL	OOPW-CW05	07603
Oosafe ICSI DISH	OOPW-IC03	07603
Oosafe 100 MM DISH	OOPW-HD10	07603
Oosafe 60MM DISH	OOPW-ST01	07506
Oosafe 35 MM DISH	OOPW-TF02	07603
Oosafe 15 mL tube	OOPW-CT01	07603
Oosafe ICSI DISH	OOPW-IC01	07506

Product was tested for RNase activity by the following protocol:

Product was extracted in RNase free water. The extract was then added to an RNA standard. The RNA standard was incubated at 37° C for 1 hour then heated to 65° C for 5 minutes. RNA samples were then run on an agarose gel, photographed, and evaluated for degradation.

FIGURE 1.



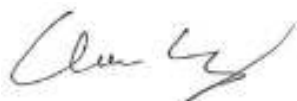
Lane (a) **product samples**, (b) unexposed RNA standard as a negative control, (c) RNA standard exposed to RNase as a positive control.

**Conclusions:**

No visible degradation is present in the product sample. The product can therefore be considered RNase free.

**Comments:**

The Test Sensitivity is 10<sup>-9</sup> Kunitz Units/ µl.



Certified by: Chase Wong, 12/08/2015



Q.A. Carl Tsang, 12/08/2015

## DNase Test Data and Results

Date: 12/07/2015  
Company: Sparmed ApS  
Date Received: 12/01/2015

Project #: 113288B  
Contact: Onur Ozturk  
Technician: Chase Wong

PO#: 130214-1  
Phone: 45- 39 40 2503

Products tested:	Product code:	Lot #:
Oosafe 4-WELL DISH	OOPW-FW04	07603
Oosafe 35 MM DISH	OOPW-TF03	07603
Oosafe 60MM DISH	OOPW-ST03	07603
Oosafe CENTER WELL	OOPW-CW05	07603
Oosafe ICSI DISH	OOPW-IC03	07603
Oosafe 100 MM DISH	OOPW-HD10	07603
Oosafe 60MM DISH	OOPW-ST01	07506
Oosafe 35 MM DISH	OOPW-TF02	07603
Oosafe 15 mL tube	OOPW-CT01	07603
Oosafe ICSI DISH	OOPW-IC01	07506

### Extraction:

Extract solution: DEPC Treated Water  
Lot #: DW15A6  
Volume: 1000µl

Number of test items exposed to extract solution: 10  
Special extraction instructions: Tested products according to extraction protocol #155.

### Procedure and Controls:

DNA: 1 kb Ladder  
DNA lot #: 1506872  
Salts: MgCl<sub>2</sub> and NaCl  
Salt lot #: S15G2

DNA standard pool: 3µl of DNA + 12µl Salts.  
Volume of each standard reaction: 5µl  
Volume of extract added to the standard: 10µl  
Total volume: 15µl

Negative Control (-): DNA and salt standards with 10 µl of unexposed extract solution added

Positive Control (+): DNA and salt standards with 10 µl of extract solution exposed to DNase from a tip exposed to human saliva

Incubation periods: 1 hr @ 37°C, followed by 5 minutes at 65°C

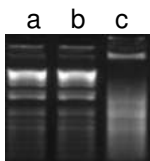
### Gel Electrophoresis:

2µl gel loading dye + 15µl reaction is loaded on a 1.2% agarose in ½ X TAE gel

Gel loading dye lot #: DD006

Electrophoresis: 30 minutes @ 80 volts

### Photographic Results:



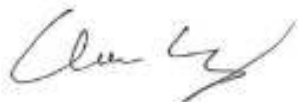
Lane (a) product samples, (b) unexposed DNA standard as a negative control, (c) DNA standard exposed to DNase as a positive control.

### Conclusions:

There is no visible degradation in lanes (a) and (b). Lane (a) represents the product samples and lane (b) represents the negative control. Lane (c), which represents the DNA standard, exposed to DNase as a positive control shows degradation of the DNA. The results suggest that the product sample is free of detectable DNase contamination.

### Recommendations:

Based on this experimental procedure, we can show a definite risk of DNase contamination if your product is exposed to saliva. We suggest that all operations are monitored and personnel are instructed in the importance of avoiding DNase contamination.



Chase Wong  
Lab Technician

12/08/2015  
Date



Carl Tsang  
Q.A.

12/08/2015  
Date



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### DNase FREE CERTIFICATE OF ANALYSIS

12/08/2015

The following sample obtained from **Sparmed ApS** on **12/01/2015** is free of any detectable DNase contamination.

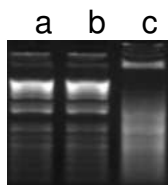
<u>PRODUCTS TESTED</u>	<u>PRODUCT CODE</u>	<u>LOT NUMBER</u>
Oosafe 4-WELL DISH	OOPW-FW04	07603
Oosafe 35 MM DISH	OOPW-TF03	07603
Oosafe 60MM DISH	OOPW-ST03	07603
Oosafe CENTER WELL	OOPW-CW05	07603
Oosafe ICSI DISH	OOPW-IC03	07603
Oosafe 100 MM DISH	OOPW-HD10	07603
Oosafe 60MM DISH	OOPW-ST01	07506
Oosafe 35 MM DISH	OOPW-TF02	07603
Oosafe 15 mL tube	OOPW-CT01	07603
Oosafe ICSI DISH	OOPW-IC01	07506

Product was tested for DNase activity by the following protocol:

Product was extracted in DNase free water. The extract was then added to a DNA standard.

The DNA standard was incubated at 37° C for 1 hour then heated to 65° C for 5 minutes. DNA samples were then run on an agarose gel, photographed, and evaluated for degradation.

**FIGURE 1.**



Lane (a) **product samples**, (b) unexposed DNA standard as a negative control, (c) DNA standard exposed to DNase as a positive control.

**Conclusions:**

No visible degradation is present in the product sample. The product can therefore be considered DNase free.

**Comments:**

The Test Sensitivity is  $10^{-7}$  Kunitz Units/  $\mu$ l.



Certified by: Chase Wong, 12/08/2015



Q.A. Carl Tsang, 12/08/2015