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-IC03

LOT No.: 07603

Expiry Date: 09/2020

Storage conditions: 20°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**

42, 3520 FARU

Quality control according to the ISO 13485:2012

Stamp:

Final approval:

Katrine Nobel

Quality Control Manager



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



ELI Accession Number: SPAR-3841-1115 Date of completion: 11-17-2015

Lot number: 07603, 07506 Reference number: OOPW-IC03, OOPW-IC01

Description of test article(s): Oosafe® ICSI/IMSI Dish for Sperm Selection, ICSI Dish

Assay system requested by customer: 1mL of culture medium was placed in the test article (OOPW-IC03) and incubated at 37°C for 30-minutes. Post incubation three 20µl of the culture medium was extracted from the test article (OOPW-IC01) and placed into another test article. 1-cell mouse embryos were then added to each drop of extract medium in the second test article and cultured for 96-hours.

Control assay method and results: 15 1-cell (BeC3F1 X BeD2F1) 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 (BeC3F1 X BeD2F1) embryos were cultured in one of the test articles using the culture medium from the other test article:

21 / 21 (100 %) 18 / 21 (86 %)

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% CO₂, 100 percent of the control embryos developed to the expanded blastocyst stage within 96-hours. 86 percent of the embryos cultured in the test article developed to the expanded blastocyst stage within 96-hours.

signature Study Director 11-17-2015

11-18-2015

signature Quality Reviewer date



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



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: SOP/TSG/ELI/008	Specification	Initial	Result % 24hr	SMI Value	Pass/Fail
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% CO2. Neither the test nor the control showed any signs that the motility was affected during the course of the assay.

signature Study Director

signature Cluality Reviewer 11-18-2015



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



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): 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: 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 St	andard s	Series	Test Sample Dilutions	N	PC	PI	PC
2λ.06	+	+	Undiluted		-	+	+
λ.03	+	*	1:2	-	12	+	+
½λ .015		*	1:4	-	22	+	+
1/4λ .0075	8.5		1:8	-	- 3	+	+
NWC		-	1:16	*	100	+	+

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.

signature

Study Director

signature Quality Reviewer

Amended: 11-18-2015

11-19-2015

RNase Test Data and Results

Date: 12/07/2015 Project #: 113288A PO#: 130214-1

Company: Sparmed ApS Contact: Onur Ozturk Phone: 45- 39 40 2503

Date received: 12/01/2015 Technician: Chase Wong

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

Extraction:

Extract solution: DEPC Treated Water Number of test items exposed to extract solution: 10

Lot #: DW15A6 Special extraction instructions: Tested products according to extraction

Volume: 1000ul protocol #155.

Procedure and Controls:

RNA: 6.0 kb Poly (A)-tailed RNA standard pool: 3μ l of RNA + 12μ l Salts. RNA lot #: 1310020 Volume of each standard reaction: 5μ l

Salts: MgCl₂ and NaCl Volume of extract added to the standard: 10µl

Salt lot #: S15G2 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

ungloved 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.

Clu J

Chase Wong12/08/2015Carl Tsang12/08/2015Lab TechnicianDateQ.A.Date



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info@mobio.com | www.mobio.com | Tel. 800-606-6246 | 2746 Loker Avenue West, Carlsbad, CA 92010

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.

PRODUCTS TESTED	PRODUCT CODE	LOT NUMBER
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^{-9} Kunitz Units/ μ l.

Certified by: Chase Wong, 12/08/2015

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

Col Tag

DNase Test Data and Results

Date: 12/07/2015 Project #: 113288B PO#: 130214-1

Company: Sparmed ApS Contact: Onur Ozturk Phone: 45- 39 40 2503

Date Received: 12/01/2015 Technician: Chase Wong

Products tested: Product code: Lot #: Oosafe 4-WELL DISH OOPW-FW04 07603 Oosafe 35 MM DISH OOPW-TF03 07603 OOPW-ST03 Oosafe 60MM DISH 07603 Oosafe CENTER WELL OOPW-CW05 07603 07603 **Oosafe ICSI DISH** OOPW-IC03 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 Number of test items exposed to extract solution: 10

Lot #: DW15A6 Special extraction instructions: Tested products according to extraction

Volume: 1000μl protocol #155.

Procedure and Controls:

DNA: 1 kb Ladder

DNA standard pool: 3μl of DNA + 12μl Salts.

DNA lot #: 1506872

Volume of each standard reaction: 5μl

Salts: MgCl₂ and NaCl Volume of extract added to the standard: 10µl

Salt lot #: S15G2 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:

2_{ul} gel loading dye + 15_{ul} 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:

a b c

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 Wong12/08/2015Carl Tsang12/08/2015Lab TechnicianDateQ.A.Date





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DNase FREE CERTIFICATE OF ANALYSIS

12/08/2015

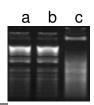
The following sample obtained from **Sparmed ApS** on **12/01/2015** is free of any detectable DNase contamination.

PRODUCT CODE	LOT NUMBER
OOPW-FW04	07603
OOPW-TF03	07603
OOPW-ST03	07603
OOPW-CW05	07603
OOPW-IC03	07603
OOPW-HD10	07603
OOPW-ST01	07506
OOPW-TF02	07603
OOPW-CT01	07603
OOPW-IC01	07506
	OOPW-FW04 OOPW-TF03 OOPW-ST03 OOPW-CW05 OOPW-IC03 OOPW-HD10 OOPW-ST01 OOPW-TF02 OOPW-CT01

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/ μ l.

Certified by: Chase Wong, 12/08/2015

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