

SpermMar Test IgG

SpermMar Test IgG

Positive and Negative Controls

For *in vitro* diagnostic use only.
Reagent for professional use only.

INTRODUCTION

As sperm does not come into contact with the blood circulation, the male reproductive system contains no antisperm antibodies in normal conditions. However, when the blood-testis barrier is breached, the immune system can detect mature sperm as antigenic and form antisperm antibodies that cause sub- or infertility. Antisperm antibodies belong to two immunological classes: immunoglobulin (Ig)A and IgG antibodies, and can be present in the semen sample as well as in male blood serum. In addition, antisperm antibodies are sometimes also found in blood serum of women. Antisperm IgG antibodies are clinically associated with immunological infertility (1-3), and screening can therefore provide help in assessing couple's infertility.

INTENDED PURPOSE

The SpermMar Test IgG is a semi-quantitative, non-automated, diagnostic kit for detecting antisperm antibodies of the IgG class on spermatozoa in human semen or serum. It is a rapid, easy-to-use microscopic test with an intended testing population of infertile couples. The test can be performed on fresh, untreated human semen sample when applying the direct SpermMar Test IgG, or on human blood serum (from men and women) when using the indirect SpermMar Test IgG. The SpermMar Test IgG may help in assessing the diagnosis and management of couple infertility.

The SpermMar Test IgG Positive Control and SpermMar Test IgG Negative Control are designed to verify the performance of the indirect SpermMar Test IgG.

PRINCIPLE OF THE TEST

The direct SpermMar Test IgG is performed by mixing fresh, untreated semen with latex particles that have been coated with human IgG, and antihuman IgG antiserum. The formation of agglutinates between the latex particles and motile spermatozoa indicates the presence of IgG antibodies on the spermatozoa.

In the indirect SpermMar Test IgG, washed motile donor spermatozoa are incubated with diluted and de-complemented patient serum of male or female origin. If the serum contains antisperm antibodies, these will cover the donor spermatozoa which will react positively in a subsequent SpermMar Test IgG.

The SpermMar Test IgG Positive and Negative Control are used as control material of the indirect SpermMar Test IgG and contain ready-to use patient serum with antisperm antibodies levels respectively higher than 80% for Positive Control and lower than 40% for Negative Control.

MATERIALS INCLUDED WITH THE TEST

SpermMar Test IgG:

- 1 vial containing 0.7 ml SpermMar Test IgG Latex Particles
- 1 vial containing 0.7 ml SpermMar Test IgG Antiserum
- Micro Slides 76 x 26 mm*
- Cover-glasses 24 x 40 mm*
- Microcapillary pipettes calibrated at 10 microliters*
- Rubber bulb*

* complete kit only

SpermMar Test IgG Positive and Negative Controls:

- 1 vial with 2.5 ml de-complemented patient serum diluted in Ferticult Flushing medium without human serum albumin

A certificate of analysis and MSDS are available on request or can be downloaded from our website (www.fertipro.com).

MATERIALS REQUIRED, BUT NOT PROVIDED

- Light microscope (with 400x to 600x magnification, bright field, dark field or phase contrast)
- For performing indirect SpermMar Test IgG: Isotonic pH buffered salt solution without protein supplement (e.g. PBS, EBSS, HTF Hepes, Ham's F10...)
- For performing indirect SpermMar Test IgG: Motile donor sperm tested negative for IgG
- Non spermicidal condom (if required)
- Microtiter plate (e.g. Kima 650 101) / Eppendorf tubes

Note that the SpermMar Test IgG Positive and Negative Control are not included in the SpermMar Test IgG and need to be purchased separately.

METHOD

Scan barcode (or download link on www.fertipro.com) to view the demonstration video.



Specimen collection and preparation

Semen collection

Standard semen collection containers should be used, typically in polypropylene and sperm survival/sperm motility tested, when semen is collected by masturbation. Non semen-toxic plastic condoms should be used when semen collection by masturbation is discouraged. Keep the semen collection container at room temperature before adding the semen sample in order to avoid large changes in temperature that may affect spermatozoa. Ideally, semen should be examined within 1 hour after ejaculation.

Serum collection

The blood sample should be collected in standard blood serum collection tubes. It is important to follow the instructions of the manufacturer of the collection tubes. Each serum tube should be inverted 10 times after collection, after waiting 30 minutes to allow coagulation the tube should be centrifuged (e.g. 10 minutes at 1000 g) to separate the serum. Serum can be stored at 2-8 °C for a maximum of 7 days.

Reagent preparation

SpermMar Test IgG Latex Particles are ready to use, however, they should be thoroughly mixed before use to provide a homogeneous suspension. SpermMar Test IgG Antiserum is ready to use.

SpermMar Test IgG Positive and Negative Controls are ready to use. Allow to adjust to room temperature before use.

Direct SpermMar Test IgG

- 1 Allow the reagents and specimens to adjust to room temperature.
- 2 On a micro slide, place:
 - 10 µl of fresh untreated semen
 - 10 µl of SpermMar Test IgG Latex Particles
 - 10 µl of SpermMar Test IgG Antiserum

This can be done by means of the provided 10 microliters capillary pipettes (complete kit).

Note: To use the microcapillary pipettes: Insert the end of the pipette marked with the heavy black line into the rubber bulb (approximately 5 mm). Allow the pipette to fill by capillary action to the first mark (10 microlitres). Do not draw liquid into the rubber bulb. Holding the bulb between the thumb and the middle finger, gently squeeze the bulb to expel the liquid from the pipette.

- 3 Mix the sample and the Latex Particles 5 times with the edge of a cover glass.
- 4 Mix the Antiserum with the Sample-Latex mixture.
- 5 Put the cover glass on the mixture and observe the mixture under a light microscope using a 400x or a 600x magnification (phase contrast or dark field illumination may facilitate reading of the slides).
- 6 Read the result after 2-3 minutes. Observe for latex particles attached to motile sperm. Count 100 spermatozoa to determine the percentage reactive sperm. If no attachment of latex particles to sperm is observed, read again after 10 minutes.

Note: Keep the preparation in a damp chamber (e.g. a Petri dish containing a moistened piece of filter paper).

Indirect SpermMar Test IgG

- 1 Allow all reagents and specimens to adjust to room temperature.
- 2 Inactivate the serum specimens by heating them at 56 °C for 30 minutes if glass test-tubes are used, 45 minutes if plastic test-tubes are used.
- 3 Adjust the pH (by adding 0.1N NaOH or HCl) of the isotonic pH buffered salt solution to 7.4 - 7.5.

- 4 Wash the motile donor spermatozoa by letting them swim up in the pH adjusted medium (pH = 7.4 - 7.5). Swim up can be done in 5 ml glass or sterile plastic test-tubes with round bottom at 37 °C for 1 hour. Adjust the sperm concentration to 20x10⁶ spermatozoa/ml with the isotonic pH buffered salt solution (pH = 7.4 - 7.5).
- 5 Serially dilute the inactivated serum specimen 1/16 with isotonic pH buffered salt solution (pH = 7.4 - 7.5) in a titer plate or Eppendorf tube.
- 6 Mix 50 µl of the (1/16) diluted, inactivated serum specimen (step 5) with 50 µl of the washed motile donor sperm (step 4) in a free well on the titer plate. Incubate for 60 minutes at 37 °C.
- 7 On a micro slide, place :
 - 10 µl of the sperm-serum mixture (step 6)
 - 10 µl of SpermMar IgG Latex Particles
 - 10 µl of SpermMar IgG Antiserum
- 8 Mix the sample and the Latex Particles 5 times with the edge of a cover glass.
- 9 Mix the Antiserum with the Sample-Latex mixture.
- 10 Put the cover glass on the mixture and observe the mixture under a light microscope using a 400x or 600x magnification (phase contrast or dark field illumination may also be used to facilitate reading).
- 11 Read the results after 2-3 minutes. Observe for latex particles attached to motile sperm. Count 100 spermatozoa to determine the percentage reactive sperm. If no attachment of particles to sperm is observed, read again after 10 minutes.

Note: Keep the preparation in a damp chamber (e.g. a Petri dish containing a moistened piece of filter paper).

Method of SpermMar Test IgG Positive and Negative Control

- 1 Allow all reagents and specimens to adjust to room temperature.
- 2 Wash the motile donor spermatozoa by letting them swim up in the pH adjusted isotonic pH buffered salt solution (pH = 7.4 - 7.5). Swim up can be done in 5 ml glass or sterile plastic test-tubes with round bottom at 37 °C for 1 hour. Adjust the sperm concentration to 20x10⁶ sp/ml with the isotonic pH buffered salt solution (pH = 7.4 - 7.5).
- 3 Mix 50 µl of control serum with 50 µl of the washed motile donor sperm in a free well on the microtiter plate or Eppendorf tube. Let incubate for 60 minutes at 37 °C.
- 4 On a micro slide, place :
 - 10 µl of the sperm-serum mixture
 - 10 µl of SpermMar Test IgG Latex Particles
 - 10 µl of SpermMar Test IgG Antiserum
- 5 Mix the sample and the Latex particles 5 times with the edge of a cover glass.
- 6 Mix the Antiserum with the Sample-Latex mixture.
- 7 The cover glass is put on the mixture and the mixture is observed under a light microscope using a 400x or 600x magnification (phase contrast or dark field illumination may also be used to facilitate reading).
- 8 Read the results after 2-3 minutes. Observe for latex particles attached to motile sperm. Count 100 spermatozoa to determine the percentage reactive sperm. If no attachment of particles to sperm is observed, read again after 10 minutes.

Note: Keep the preparation in a damp chamber (e.g. a Petri dish containing a moistened piece of filter paper).

CONTROL - CONTROL + IVD

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MATERIAL INCLUDED

Catalogue number

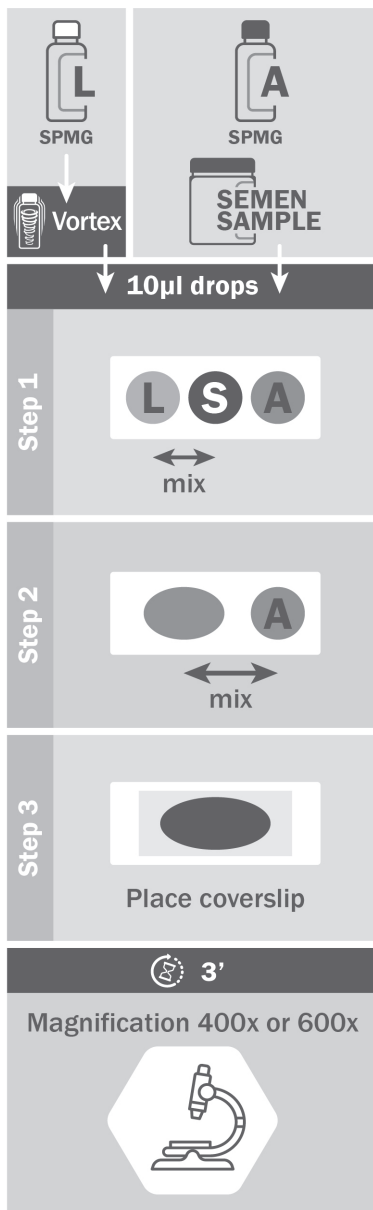
SpermMar Test IgG	
SPMG_S	SpermMar Test IgG Single kit 50 tests
SPMG_C	SpermMar Test IgG Complete kit 50 tests
SpermMar Test IgG - Positive & Negative Control	
SPMG_P	1 vial with 2.5 ml of positive control serum for the SpermMar Test IgG
SPMG_N	1 vial with 2.5 ml of negative control serum for the SpermMar Test IgG

CUSTOMER-TECHNICAL SUPPORT

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Graphic presentation of the protocol:



INTERPRETATION OF THE RESULTS

SpermMar Test IgG

When the test is properly performed, the absence of sperm antibodies will be shown by freely moving spermatozoa not covered by latex particles. The latex particles themselves will form growing agglutinates thus proving the reactivity of the reagents. In the presence of sperm antibodies however, the spermatozoa will be partially covered by latex particles. In some cases the spermatozoa might even be immobilized by the massive amount of adherent latex particles.

In the direct SpermMar Test IgG, the diagnosis of immunological infertility is suspected when 10-39% of the motile spermatozoa are covered by latex particles; if 40% or more of the spermatozoa are covered, immunological infertility is highly probable. Additional tests should confirm the diagnosis. Whenever a positive result is obtained it is recommended to perform the SpermMar Test IgA (FertiPro NV) as well.

In the indirect SpermMar Test IgG, the occurrence of 40% or more reaction between the coated latex particles and motile spermatozoa is generally accepted as the lower limit of significant antibody binding.

SpermMar Test IgG Positive and Negative Control

- The SpermMar Test IgG Positive Control should yield 80% or more of the motile spermatozoa covered with latex particles.
- The SpermMar Test IgG Negative Control should yield less than 40% spermatozoa covered with latex particles.

LIMITATIONS OF THE METHOD

The direct SpermMar Test IgG can only be performed if motile spermatozoa are present in the semen. Samples with very low sperm concentration and/or poor motility cannot be evaluated, since 100 motile spermatozoa must be assessed following incubation with the reagents. Immotile cells should not be counted. When this cannot be achieved, it is suggested to perform the indirect SpermMar Test IgG on blood serum.

Positive and Negative controls can only be applied in an indirect SpermMar Test, and the donor semen must contain motile spermatozoa negative for IgG.

PERFORMANCE CHARACTERISTICS

Direct SpermMar Test IgG

When the direct SpermMar Test IgG is compared to the direct Immunobead Test, a good correlation could be found between both tests (4-7). A positive correlation was also found between the direct SpermMar Test IgG and the flow cytometry measurement (8-10).

Indirect SpermMar Test IgG

When the indirect SpermMar Test IgG is compared to the indirect Immunobead Test, a good correlation could be found between both tests (4, 5, 11). Furthermore, a good correlation between the indirect SpermMar Test IgG and the Tray Agglutination Test was found (4, 12, 13).

Repeatability and reproducibility

Repeatability and reproducibility were assessed using samples with different severities of immunological reaction. The CV_{intra} and CV_{inter} of the SpermMar Test IgG is 3.5% and 3.23% respectively, which is well below 15%, indicating an acceptable repeatability and reproducibility for the SpermMar Test IgG.

STORAGE/DISPOSAL

- SpermMar Test IgG is intended for 50 individual (in)direct SpermMar IgG tests that can be performed spread over the shelf life. The SpermMar Test IgG Positive and Negative Control are intended for 25 individual tests spread over the shelf life. After each individual test, all used reagents and materials should be discarded. Close reagent bottles well after each use and store at 2-8 °C. Even after opening, the SpermMar Test IgG reagents and controls are stable for 18 months from the date of manufacturing.
- Do not use after expiry date.
- Do not freeze.
- Stable for transport or short term storage at elevated temperatures (up to 5 days at 37 °C).
- The reagents need to be disposed in accordance with local regulations for disposal of medical devices taking into account that the device contains human and/or animal derived substances.

WARNINGS AND PRECAUTIONS

All human, organic material should be considered potentially infectious. Handle all specimens as if capable of transmitting HIV or hepatitis. Always wear protective clothing when handling specimens.

SpermMar Test IgG latex particles contain 0.1% Bovine Serum Albumin of US origin, which is certified by an EDQM Certificate of Suitability. Furthermore, the product meets European requirements for treated technical blood products.

SpermMar Test IgG latex particles are coated with human IgG, which are biotechnologically manufactured, therefore an infection with hepatitis, HIV 1/2 or other infectious diseases can be considered impossible.

SpermMar Test IgG Antiserum does contain rabbit antiserum to human IgG. Contamination is prevented by the addition of sodium azide as a preservative (< 1g/l).

The SpermMar Test IgG Positive and Negative Controls contain human serum, which has been tested for HIV, Hepatitis B and Hepatitis C. However, the user should always wear protective clothing when handling the control sera.

Any serious incident (as defined in the European In Vitro Diagnostic Medical Device Regulation 2017/746) that has occurred should be reported to FertiPro NV and, if applicable, to the competent authority of the EU Member State in which the user and/or patient is established.

BIBLIOGRAPHY

- Cui D, Han G, Shang Y, Liu C, Xia L, Li L, et al. Antisperm antibodies in infertile men and their effect on semen parameters: a systematic review and meta-analysis. Clin Chim Acta. 2015;444:29-36.
- Lombardo F, Gandini L, Dondero F, Lenzi A. Antisperm immunity in natural and assisted reproduction. Hum Reprod Update. 2001;7(5):450-6.
- Francavilla F, Santucci R, Barbonetti A, Francavilla S. Naturally-occurring antisperm antibodies in men: interference with fertility and clinical implications. An update. Front Biosci. 2007;12:2890-911.
- Andreou E, Mahmoud A, Vermeulen L, Schoonjans F, Comhaire F. Comparison of different methods for the investigation of antisperm antibodies on spermatozoa, in seminal plasma and in serum. Hum Reprod. 1995;10(1):125-31.
- Khoo D, Feigenbaum SL, McClure RD. Screening assays for immunologic infertility: a comparison study. Am J Reprod Immunol. 1991;26(1):11-6.
- Hellstrom WJ, Samuels SJ, Waits AB, Overstreet JW. A comparison of the usefulness of SpermMar and immunobead tests for the detection of antisperm antibodies. Fertil Steril. 1989;52(6):1027-31.
- Marconi M, Nowotny A, Pantke P, Diemer T, Weidner W. Antisperm antibodies detected by mixed agglutination reaction and immunobead test are not associated with chronic inflammation and infection of the seminal tract. Andrologia. 2008;40(4):227-34.
- Rasanen M, Agrawal YP, Saarikoski S. Seminal fluid antisperm antibodies measured by direct flow cytometry do not correlate with those measured by indirect flow cytometry, the indirect immunobead test, and the indirect mixed antiglobulin reaction. Fertil Steril. 1996;65(1):170-5.
- Rasanen M, Lahteenmaki A, Saarikoski S, Agrawal YP. Comparison of flow cytometric measurement of seminal antisperm antibodies with the mixed antiglobulin reaction and the serum tray agglutination test. Fertil Steril. 1994;61(1):143-50.
- Nikolaeva MA, Kulakov VI, Ter-Avanesov GV, Terekhina LN, Pshenichnikova TJ, Sukhikh GT. Detection of antisperm antibodies on the surface of spermatozoa using flow cytometry: preliminary study. Fertil Steril. 1993;59(3):639-44.
- Kay DJ, Boettcher B. Comparison of the SpermMar test with currently accepted procedures for detecting human sperm antibodies. Reprod Fertil Dev. 1992;4(2):175-81.
- Hinting A, Vermeulen L, Comhaire F. The indirect mixed antiglobulin reaction test using a commercially available kit for the detection of antisperm antibodies in serum. Fertil Steril. 1988;49(6):1039-44.
- Comhaire FH, Hinting A, Vermeulen L, Schoonjans F, Goethals I. Evaluation of the direct and indirect mixed antiglobulin reaction with latex particles for the diagnosis of immunological infertility. Int J Androl. 1988;11(1):37-44.

SYMBOLS GLOSSARY

Symbols as defined in ISO 15223	
	Catalogue number
	Batch code
	Consult instructions for use
	Manufacturer
	In Vitro Diagnostics
	Temperature limit
	Use-by date
	Caution
	Negative control
	Positive control
	Contains biological material of animal origin
	Contains human blood or plasma derivatives

Symbol as defined in IVDR 2017/746



CE marking by Notified Body 2797



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