

SpermMar Test IgG

A qualitative Latex Test for detection of Sperm Antibodies

Preservative: Sodium azide 0.09%. Store at 2° to 8° C - Do Not Freeze For in vitro diagnostic use only. Reagent for professional use only.

INTENDED USE

The SpermMar Test IgG is a diagnostic kit for detecting antisperm antibodies of the IgG class in human semen or serum. The direct SpermMar Test IgG can be performed on untreated human semen provided it contains motile spermatozoa, the indirect SpermMar Test IgG can be used on serum. The presence of antisperm antibodies can interfere with sperm function and zona binding and the acrosome reaction.

GENERAL INFORMATION

The presence of sperm antibodies reacting with antigens on the spermatozoa is considered as typical and specific for immunological infertility (2, 4, 11). These antibodies are found in approximately 8% of infertile men (13). Sperm antibodies belong to two immunological classes; IgA and IgG antibodies. There are some data indicating IgA to be more elimically important they IgG antibodies.

IgA to be more clinically important than IgG antibodies. However, IgA antibodies rarely occur without IgG antibodies. Therefore, testing for IgG antibodies is sufficient for routine screening (6.7.14)

The direct SpermMar Test IqG is performed by mixing fresh The direct SpermMar Test IgG is performed by mixing fresh, untreated semen with latex particles that have been coated with human IgG. To this mixture a monospecific antihuman IgG antiserum is added. The formation of agglutinates between particles and motile spermatozoa indicates the presence of IgG antibodies on the spermatozoa (1,5,9,10). In the Indirect SpermMar Test IgG washed motile donor spermatozoa are incubated with diluted and de-complemented noting the properties of the properties of

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 IqG.

PRODUCT ORDER CODES

SpermMar Test IgG single kit – 50 tests SpermMar Test IgG complete kit – 50 tests

MATERIALS INCLUDED WITH THE TEST

- » 1 vial containing 0.7 ml SpermMar Test IgG Latex Particles
- » 1 vial containing 0.7 ml SpermMar Test lgG Antiserur
- » Micro Slides 76 x 26 mm*
- » Cover-glasses 24 x 40 mm*
 » Microcapillary pipettes calibrated at 10 microlitres*

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

MATERIALS NOT INCLUDED WITH THE TEST

- » Light microscope (with 400x to 600x magnification
- Light microscope (with 400x to 600x magninication), bright field, dark field or phase contrast)
 EBSS medium without added protein for the indirect SpermMar Test IgG (e.g. Sigma-Aldrich E2888)
 Non spermicidal condom
 Microtiter plate (e.g. Kima 650 101)

DIRECTIONS FOR USE



We recommend to watch our demonstration video (download via link on our website or scan barcode).

SPECIMEN COLLECTION & PREPARATION

Semen collection by masturbation is preferred. Where particular circumstances discourage collection by masturbation, specific plastic condoms are available from FertiPro for semen collection. Ordinary condoms should not be used for semen collection because they may interfere with the motility and viability of the spermatozoa, Ideally, semen should be nined within 1 hour after eiaculation

REAGENT PREPARATION

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

DIRECT SPERMMAR TEST IGG

- Allow the reagents and specimens to adjust to room
- temperature.
 2. On a micro slide place :

- On a micro slide place:

 10 microlitres of fresh untreated semen

 10 microlitres of SpermMar Test IgG Latex Particles

 10 microlitres of SpermMar Test IgG Antiserum
 This can be done by means of the provided 10 microlitres
 capillary pipettes (complete test).

 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 betw the thumb and the middle finger, gently squeeze the bulb to expel the liquid from the pipette.
- Mix the sample and the Latex reagent 5 times with the edge of a cover glass.
 Mix the Antiserum with the Latex reagent and sample mixture.
- The cover glass is put on the mixture and the mixture is observed under a light microscope using a 400x or a 600x magnification (phase contrast or dark field illumi nation may facilitate reading of the slides). Read the result after 2-3 minutes. Observe for latex par-
- ticles attached to motile sperm. Count 100 sperm to determine the percentage reactive sperm

If no attachment of beads to sperm is observed, read again after 10 minutes

Note: Keep the preparation in a damp chamber (e.g. a Petri dish stened niece of filter naner

INDIRECT SPERMMAR TEST IGG

- Allow all reagents and specimens to adjust to room
- temperature. 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. Adjust the pH (by adding 0.1N NaOH or HCl) of the EBSS
- to 7.4 7.5.
- 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 has to be done in 5 ml glass or sterile plastic test-tubes with round bottom at 37°C for 45 minutes Adjust the sperm concentration to 20x10⁶ sp/ml with EBSS medium (pH = 7.4 7.5)

 5. Serially dilute the inactivated serum specimen 1/16 with FBSS medium (pH = 7.4 7.5) in a titre plate
- EBSS medium (pH = 7.4 7.5) in a titre plate 6. Mix 50 microlitres of the (1/16) diluted, inactivated serum
- specimen (step 5) with 50 microlitres of the washed motile donor sperm (step 4) in a free well on the titre plate Incubate for 60 minutes at 37°C.
- On a micro slide place :

- what a mice since piece of the sperm-serum mixture

 10 microlitres of the sperm-serum mixture

 10 microlitres of SpermMar IgG Latex Particles

 10 microlitres of SpermMar IgG Antiserum

 Mix the sample and the Latex reagent 5 times with the edge
- of a cover glass. Mix the Antiserum with the Latex reagent and sample mixture. 10. 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
- magnification (phase contrast or ank field illumination m also be used to facilitate reading)

 11. Read the results after 2-3 minutes. Observe for latex particles attached to motile sperm. Count 100 spermatoz 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). To prevent evaporation during incubation, always cover with Parafilm.

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 IqG, the diagnosis of immunological infertility is suspected when 10-39% of the motile spermatozoa are 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. In the indirect SpermMar Test IgA, the occurrence of 40% or more reaction between the coated latex particles and motils exemptoza is engagedly accepted. latex particles and motile spermatozoa is generally accepted as the lower limit of significant activity.

LIMITATIONS OF THE METHOD

The direct SpermMar Test IgG can only be performed if motile spermatozoa are present in the semen. Samples with poor motility may yield false negative results in those cases it is suggested to perform the indirect SpermMar Test IgG.

PERFORMANCE CHARACTERISTICS

DIRECT SPERMMAR TEST IGG

Several hundreds of semen samples were tested with the direct MAR-Test (mixed antiglobulin reaction based on red blood cells) and with the SpermMar Test IgG. The results were similar in 97% of the cases the MAR-Test based on red blood cells was negative while the SpermMar Test IgG detected antibody coated spermatozoa, though in relatively small numbers (<40%), thus proving the higher sensitivity of the SpermMar Test IgG (10,16).

INDIRECT SPERMMAR TEST IGG

Using the value of 40% reaction between motile spermatozoa and coated Latex particles as the lower limit of significant activity, the indirect SpermMar Test IgG was found positive in some cases with negative results of the Tray Agglutination

in some cases with negative results of the Tray Agglutination Test, or other currently accepted procedures.

The SpermMar Test 1g6 was proven easier to use and more sensitive (15). A false negative indirect SpermMar Test 1g6 in comparison with the Tray Agglutination Test occurred in cases with 1gM in serum, the clinical significance of which is doubtful. It is recommended to confirm a positive result of the indirect SpermMar Test 1g6 by additional tests for the detection of agglutinating activity (Tray Agglutination Test) and of evolutions are set and the ATD-Belease. Test) and of cytotoxic activity, such as the ATP-Release cytotoxicity test. The latter tests will also assess the type of nunological effect exerted by the antisperm antibo

REAGENT STORAGE

SpermMar Test IgG reagents are stable for 18 months from the date of manufacturing. SpermMar Test IgG reagents must be stored at 2° to 8°C when not in use. DO NOT FREEZE. Suitable for transport or short term storage at elevated temperatures (up to 5 days at 37°C).

WARNINGS AND PRECAUTIONS

All human, organic material should be considered potentially 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 contains 0.1% Bovine Serum Albumin of US origin.

SpermMar Test IgG latex particles are coated with human IgG. all materials used have been tested by their original.

IgG, all materials used have been tested by their original manufacturer for Hepatitis B, Hepatitis C and HIV.

SpermMar Test IgG **Positive and Negative** Controls

Controls for use with the Indirect Test for Determination of Sperm Antibodies (SpermMar IgG)

Preservative: Sodium azide 0.09%. Store at 2° to 8° C - Do Not Freeze. For in vitro diagnostic use only. Reagent for professional use only.

INTENDED USE

The SpermMar Test IgG Positive and SpermMar Test IgG Negative Control are designed to be used as quality control with the SpermMar Test IaG

GENERAL INFORMATION

The presence of sperm antibodies reacting with antigens on the spermatozoa is considered as typical and specific for immunological infertility (2,4,11). Sperm antibodies belong to two immunological classes; IgA and IgG antibodies. There are some data indicating IgA to be more clinically important the Lot of the data and carried to the control of the than IgG antibodies. However, IgA antibodies rarely occur without IgG antibodies. Therefore, testing for IgG antibodies is sufficient as a routine screening method (6,7,14).

In the Indirect SpermMar Test IgG washed motile donor spermatozoa are incubated with diluted, de-complemented spermatozoa are incupated with diluted, 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 Control contains ready-local patients of the services of the

use patient serum with antisperm antibodies levels higher than 80%. The SpermMar Test IgG Negative Control contains ready-to-use patient serum with antisperm antibodies levels

PRODUCT CODES AND KIT CONTENTS

1 vial with 2.5 ml of positive control serum for the SpermMar Test IgG 1 vial with 2.5 ml of negative control serum for the SpermMar Test IgG

MATERIAL INCLUDED WITH THE TEST

» 1 vial with 2.5ml de-complemented patient serum diluted

A certificate of analysis and MSDS are available on request

MATERIALS NOT INCLUDED WITH THE TEST

- Microscope slides
- Cover glasses
- » Light microscope (with 400x to 600x magnification, bright
- field, dark field or phase contrast)

 EBSS medium (e.g. Sigma-Aldrich E2888)
- Microtiter plate (e.g. Kima 650 101)

DIRECTIONS FOR USE

REAGENT PREPARATION

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

SPECIMEN COLLECTION AND PREPARATION

The donor semen should be collected by masturbation or by other methods recommended by the physician.

Proferentially semen should be examined within 1 hour Preferentially, semen should be exam after ejaculation.

PROCEDURE

- Allow all reagents and specimens to adjust to room
- Wash the motile donor spermatozoa by letting them swim up in the pH adjusted EBSS medium (pH = 7.4 - 7.5). Swim up has to be done in 5 ml glass or sterile plastic test-tubes with round bottom at 37°C for 45 minutes. Adjust the sperm concentration to 20x106 sp/ml with
- Adjust the sperm concentration to 20x10° sp/ml with medium (pH = 7.4 7.5).

 Mix 50 microlitres of control serum with 50 microlitres of the washed motile donor sperm in a free well on the microtiter plate. Let incubate for 60 minutes at 37°C. On a micro slide place:

 10 microlitres of the sperm-serum mixture
- » 10 microlitres of SpermMar Test IgG Latex Particles » 10 microlitres of SpermMar Test IgG Antiserum
- 5. Mix the sample and the Latex reagent 5 times with the edge of a cover glass.

 6. Also mix the Antiserum with the Latex reagent and
- Also mix the Antiserum with the Latex reagent and sample mixture.

 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). Read the results after 2-3 minutes. Observe for latex particles attached to motile sperm. Count 100 spermatora to determine the necentage reactive sperm I for
- tozoa to determine the percentage reactive sperm. If no attachment of particles to sperm is observed, read again

Note: Keep the preparation in a damp chamber (e.g. a Petri dish containing a moistened piece of filter paper). To prevaporation during incubation, always cover with Parafilm

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.

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

LIMITATIONS OF THE PROCEDURE

The indirect SpermMar Test IgG can only be performed if motile spermatozoa are present in the semen

REAGENT STORAGE

When stored properly, SpermMar IgG control sera are stable for 18 months from the date of manufacturing. SpermMar IgG control sera must be stored at 2° to $8^{\circ}C$ when not in use.

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

Although SpermMar Test IgG Positive and Negative Controls have been tested for HIV and hepatitis the user should always othing when handling the control sera.

BIBLIOGRAPHY

TECHNICAL SUPPORT



FertiPro N.V. Industriepark Noord 32, 8730 Beernem, Belgium Tel +32 (0)50 79 18 05 Fax +32 (0)50 79 17 99





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