

## User's Manual

# Anti-Spermatozoa Antibody ELISA

Enzyme linked immunosorbent assay (ELISA) for the determination of antibodies directed against spermatozoa antigens in seminal plasma

**REF**

**IB79155**



**96**

**RUO**

**For Research Use Only – Not for Use in Diagnostic Procedure**

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## INTENDED USE

The anti-spermatozoa antibody ELISA from IBL-America is a reliable test for the determination anti-bodies directed against human spermatozoa. This test is intended for the use with seminal plasma.

**Please note:** the terms “anti-spermatozoa antibodies”, “anti-sperm antibodies” and “sperm antibodies” are equivalent. In these instructions the rather unwieldy but correct term “anti-spermatozoa antibodies” is used. **For Research Use Only – Not for Use in Diagnostic Procedure**

## DISCUSSION

Antibodies directed against spermatozoa antigens may cause infertility in women or men. The application of the Anti-Spermatozoa Antibody ELISA from IBL-America is recommended for the diagnosis of immunologically caused disorders of fertility.

Unwanted childlessness is a growing problem with which up to 20% of all couples in the reproductive age are confronted temporarily or long-term. In 20% of these cases the presence of anti-spermatozoa antibodies in the male or the female individual is detectable (Lahteenmaki A et al: Hum Reprod (1995) 10, 2824-28; Nagy ZP et al: Hum Reprod (1995) 10, 1775-80).

The definition of infertility according to the WHO (WHO Laboratory Manual for the Examination of Human Semen and Semen Cervical-Mucus Interaction, 1999) is the absence of a conception within 12 months of unprotected intercourse. The main cause of an immunological fertility disorder is the formation of antibodies directed against spermatozoa antigens.

Anti-spermatozoa antibodies exert heterogeneous effects on the ability of spermatozoa to fertilize. The inhibiting effect of anti-spermatozoa antibodies on the motility of spermatozoa by binding to their surface and by agglutinating processes is well-known (Zouari R et al: Fertil Steril (1993) 59, 606-12).

The penetration of the spermatozoa into the cervical mucus is impaired by the presence of anti-spermatozoa antibodies in the seminal plasma and/or in the cervical mucus (Eggert-Kruse W et al: Hum Reprod (1993) 8, 1025-31). Anti-spermatozoa antibodies negatively influence the capacitation and the acrosome reaction of spermatozoa and thereby impede the interaction of the spermatozoa with the oocyte (Francavilla F et al: Front Biosci (1999): 1;4:9-25; Bohring C et al.: Hum Reprod (2001) 7:113-8).

The interaction of the spermatozoon with the oocyte and the subsequent binding to and penetration of the zona pellucida may be inhibited by anti-spermatozoa antibodies. The following fusion of the oocyte and a spermatozoon may also be impaired by the presence of anti-spermatozoa antibodies (Mazumdar S et al.: Fertil Steril (1998) 70, 799-810; Kutteh WH: Hum Reprod, (1999) 14, 2426-9).

According to Crosignani et al. (Crosignani et al.: PG et al.: Hum Reprod (1998) 13, 2025-32) the rate of pregnancies in couples with anti-spermatozoa antibodies on the part of the man or the woman are 38% lower compared to the control groups. Furthermore an influence on the implantation and on the early embryological development could be confirmed. An association of anti-spermatozoa antibodies and miscarriages is discussed.

The frequency of anti-spermatozoa antibodies in infertile couples amounts to 20% (Lahteenmaki A et al.: Hum Reprod (1995) 10, 2824-28; Nagy ZP et al.: Hum Reprod (1995) 10, 1775-80).

Anti-spermatozoa antibodies may occur dissolved in the ejaculate or bound to the surface of spermatozoa. Anti-spermatozoa antibodies may be found in men and in women (Clarke GN et al.: Am J Reprod Immunol Microbiol (1985) 7, 143-7). In women anti-spermatozoa antibodies may be found in cervical mucus, oviduct liquid and follicular liquid. Men having more than 50% of their spermatozoa coated with anti-spermatozoa antibodies show a conspicuously reduced rate of fertility (Abshagen K et al.: Fertil Steril (1998) 70, 355-6).

**PRINCIPLES OF THE ASSAY METHOD**

The anti-spermatozoa antibody ELISA (**Enzyme Linked ImmunoSorbent Assay**) from IBL-America is a solid-phase sandwich enzyme-immunoassay for the determination of anti-spermatozoa antibodies in human seminal plasma.

The ELISA-plate is coated with a mix of spermatozoa proteins which are recognized by anti-spermatozoa antibodies. The samples and standards are pipetted into the wells and then incubated. During this incubation anti-spermatozoa antibodies bind to the spermatozoa proteins and are thus immobilised on the plate. After washing the enzyme conjugate, consisting of anti-human globulin antibodies covalently coupled to horseradish peroxidase, is added. After removal of the unbound conjugate by washing the horseradish peroxidase oxidizes the then added substrate TMB (3,3',5,5'-tetramethylbenzidine) yielding a colour reaction which is stopped with 0.5 N acidic solution. The extinction is measured at a wavelength of 450 nm with a microplate reader. The use of a reference measurement with a wavelength  $\geq 550$  nm is recommended.

**REAGENTS**

(sufficient for 96 determinations)

1. Microtiter strips coated with sperm antigen	96 wells
2. Sperm Antibody ELISA standard set - per vial	0.5 ml
– Standard 1 ( 31 U/ml – colourless screw cap)	
– Standard 2 ( 62 U/ml – white screw cap)	
– Standard 3 (125 U/ml – yellow screw cap)	
– Standard 4 (250 U/ml – blue screw cap)	
3. Control (green screw cap) equivalent to 70-120 U/ml	0.5 ml
4. Dilution buffer (also used as blank / zero standard / 0 U/ml )	50 ml
5. Washing solution (10x concentrated)	50 ml
6. Enzyme conjugate (ready for use)	8 ml
7. Substrate solution (solution of TMB, ready for use)	13 ml
8. Stop solution (0.5 N acidic solution)	13 ml
9. Holder for single strips	1 x
10. Adhesive Sheet	2 x

**MATERIALS REQUIRED BUT NOT INCLUDED**

1. Microplate reader with 450 nm filter, optionally with a reference filter  $\geq 550$  nm.
2. Microliter pipettes with disposable tips: 5  $\mu$ l, 50  $\mu$ l and 500  $\mu$ l.
3. Tubes for the dilution of the samples
4. Distilled or deionised water
5. Absorbent paper.
6. Please use only calibrated pipettes and instruments.

**WARNINGS AND PRECAUTIONS**

1. This kit is intended for research use only.
2. Avoid contact with the stop solution, it may cause skin irritations and burns.
3. Do not pipette reagents by mouth.
4. Please regard all samples as potentially infectious and handle them with utmost care.
5. Handling and disposal should be in accordance with the procedures defined by an appropriate national biohazard safety guideline or regulation where this exists.

## INSTRUCTIONS FOR REAGENT PREPARATION

1. The components of this kit are intended for use as an integral unit and should not be interchanged with the components of other kits.
2. All reagents and samples must be brought to room temperature before use.
3. All reagents have to be mixed without foaming.
4. Once the test procedure has been started, all steps should be continued without interruption.
5. Pipette all reagents and samples onto the bottom of the wells. Mixing or shaking after pipetting is not required.
6. Use new disposable tips for each sample.
7. Before starting the assay, all reagents to be used should be prepared and ready for immediate use, all needed strips should be secured in the holder etc. This will ensure equal time periods for each pipetting step without interruption.
8. For optimal results it is important to wash the wells thoroughly after incubation and to remove even the last water drops by hitting the plate on absorbent paper or cloth.
9. Since the kinetics of the enzymatic reaction depends on the surrounding temperature different extinctions correlating with the respective room temperature may be observed. The optimum laboratory room temperature is 20 °C – 22 °C (68 °F – 72 °F).
10. It is recommended to effect all tests in double determination in order to minimize the consequences of pipetting or handling errors.

## STORAGE INSTRUCTIONS AND SHELF LIFE INFORMATION

1. Store the reagents at 2 °C – 8 °C (36 °F – 46 °F).
2. The reagents remain stable until the expiration date of the kit.
3. The diluted washing solution is stable for 4 weeks at refrigerator temperatures (4 °C – 8 °C / 39 °F – 46 °F).
4. Put caps back on the vials immediately after use.
5. Store the microtiter strips in a dry bag with desiccants. The remaining strips must be stored in the tightly resealed bag together with the desiccants. Under these storage conditions, they are stable at least for 4 weeks after opening of the sealed bag.

## SAMPLE MATERIAL

Seminal plasma

## SAMPLE COLLECTION AND PREPARATION

Collect fresh ejaculate, centrifuge at room temperature and take the supernatant (seminal plasma). Avoid repeated freezing and thawing of the seminal plasma. Store tubes closed as they may be a danger of contamination or of alteration of concentration.

1. Handle all samples with utmost care since they may be infectious.
2. There are no known interferences with extrinsic factors or other substances.
3. Samples may be stored at different temperatures for the following time-spans:
  - Environmental temperature up to 30 °C (86 °F): up to three days
  - Refrigerator temperature (2 – 8 °C / 36 °F – 46 °F): up to one week
  - Household freezer temperature (-10 °C – -20 °C / 14 °F – -4 °F): up to one year

**ATTENTION!** There are no test methods available which may guarantee that Hepatitis B virus, Human Immunodeficiency Virus (HIV/HTLV-III/LAV), or other infectious agents are absent from the reagents in this kit. Therefore, all human products of human origin, including samples, should be considered potentially infectious.

**ASSAY PROCEDURE**

1. Warm all reagents to room temperature and mix thoroughly before use.
2. Preparation of the washing solution (10x): Dilute the concentrated washing solution (50 ml) by adding 450 ml distilled or demonized water. **Attention:** Do not use unpurified tap water!
3. Dilute seminal plasma 1:5 (1+4) with dilution buffer, for example, dilute 100 µl seminal plasma with 400 µl dilution buffer.
4. Fix the required number of coated wells or strips in the strip holder.
5. Pipette 50 µl of standards into the respective wells.
6. Pipette 50 µl of the diluted seminal plasma with new disposable tips into the respective wells.
7. Incubate for 60 min at 37 °C. **Please use the adhesive sheet provided with this kit or, better still, use a humid chamber** in order to minimize loss of liquid due to evaporation.
8. Briskly shake out the contents of the wells and then rinse the wells 3 times with 200 µl diluted washing solution.
9. Knock the residual water out of the wells by hitting them (in the holder) on absorbent paper or cloth.
10. Dispense 50 µl of the enzyme conjugate into each well.
11. Incubate for 60 min at 37 °C. **Please use the adhesive sheet provided with this kit or, better still, use a humid chamber** in order to minimize loss of liquid due to evaporation.
12. Briskly shake out the contents of the wells and then rinse the wells 5 times with 200 µl diluted washing solution.
13. Knock the residual water out of the wells by hitting them (in the holder) on absorbent paper or cloth.
14. Dispense 50 µl of substrate solution immediately after the washing to each well.
15. Incubate for 30 min at room temperature.
16. Stop the enzymatic reaction by adding 50 µl of stop solution into each well in the same sequence and time interval as dispensing the substrate.
17. Measure the extinction of the samples at 450 nm. It is recommended to carry out the measurement of the extinction within 10 minutes after stopping the reaction.

As a general rule the enzymatic reaction is linearly proportional to time and temperature. This makes interpolation possible for fixed physico-chemical conditions.

Since calibrators are assayed in each run, absorbance fluctuations do not affect the absolute results. In any case it is highly recommended to use an additional internal control if available.

**PIPETTING SCHEME FOR THE SPERM ANTIBODY ELISA FROM IBL-AMERICA**

	1	2	3	4	5	6	7	8	9	10	11	12
A	BLANK	BLANK	P	3	P	11	P	19	P	27	P	35
B	S	1	P	4	P	12	P	20	P	28	P	36
C	S	2	P	5	P	13	P	21	P	29	P	37
D	S	3	P	6	P	14	P	22	P	30	P	38
E	S	4	P	7	P	15	P	23	P	31	P	39
F	P	C	P	8	P	16	P	24	P	32	P	40
G	P	1	P	9	P	17	P	25	P	33	P	41
H	P	2	P	10	P	18	P	26	P	34	P	42

In this pipetting scheme the recommended positions for the blank (please use the dilution buffer included in this kit), standards (S1 – S4), positive control (PC) and for the samples (P1 – P42) are shown as double determinations.

## RESULTS

1. Calculate the average absorbance values for each set of reference standards, controls and unknown samples
2. The optical density of each standard value is plotted as y value (y-axis), the corresponding anti-spermatozoa antibody value is drawn in as the x-value (x-axis). The resulting calibration curve is used to determine the values of the samples. The OD values of the samples are correlated with the corresponding sperm antibody concentration values by interpolation.
3. Using the mean absorbance value for each sample determine the corresponding concentration of anti-spermatozoa antibody in U/ml from the standard curve.

## LIMITATIONS OF THE ASSAY

- At temperatures higher than 30 °C (86 °F) the samples should be transported cooled or refrigerated. The time to stop the (enzymatic colour) reaction may have to be adjusted (shortened).

## ASSAY PERFORMANCE CHARACTERISTICS

**1. Intraassay variation coefficient: 6.44% (5.69 – 7.92%)**

For the determination of the intraassay variation coefficient 6 kits from 6 different batches (produced on different days) were used. One sample (optical density about 1.0) was applied 96 times per testing procedure.

**2. Interassay variation coefficient: 7.15% (6.04 – 8.21)**











For the determination of the interassay variation coefficient one strip each of 12 kits stemming from 6 different batches (produced on different days) were used. One sample (optical density about 1.0) was applied 72 times per testing procedure.

### Manufactured for :

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## SYMBOLS USED WITH IBL-AMERICA ASSAYS

Symbol	English	Deutsch	Français	Español	Italiano
	Consult instructions for use	Gebrauchsanweisung beachten	Consulter les instructions d'utilisation	Consulte las instrucciones de uso	Consultare le istruzioni per l'uso
	European Conformity	CE-Konformitätskennzeichnung	Conformité aux normes européennes	Conformidad europea	Conformità europea
	In vitro diagnostic device	In-vitro-Diagnostikum	Usage Diagnostic in vitro	Para uso Diagnóstico in vitro	Per uso Diagnostica in vitro
	For research use only	Nur für Forschungszwecke	Seulement dans le cadre de recherches	Sólo para uso en investigación	Solo a scopo di ricerca
	Catalogue number	Katalog-Nr.	Numéro de catalogue	Número de catálogo	Numero di Catalogo
	Lot. No. / Batch code	Chargen-Nr.	Numéro de lot	Número de lote	Numero di lotto
	Contains sufficient for <n> tests/	Ausreichend für "n" Ansätze	Contenu suffisant pour "n" tests	Contenido suficiente para <n> ensayos	Contenuto sufficiente per "n" saggi
	Storage Temperature	Lagerungstemperatur	Température de conservation	Temperatura de conservación	Temperatura di conservazione
	Expiration Date	Mindesthaltbarkeitsdatum	Date limite d'utilisation	Fecha de caducidad	Data di scadenza
	Legal Manufacturer	Hersteller	Fabricant	Fabricante	Fabbricante
Distributed by	Distributor	Vertreiber	Distributeur	Distribuidor	Distributore
Content	Content	Inhalt	Conditionnement	Contenido	Contenuto
Volume/No.	Volume / No.	Volumen/Anzahl	Volume/Quantité	Volumen/Número	Volume/Quantità

Symbol	Portugues	Dansk	Svenska	Ελληνικά
	Consulte as instruções de utilização	Se brugsanvisning	Se bruksanvisningen	Εγχειρίδιο χρήστη
	Conformidade com as normas europeias	Europæisk overensstemmelse	Europeisk överensstämmelse	Ευρωπαϊκή Συμμόρφωση
	Diagnóstico in vitro	In vitro diagnostik	Diagnostik in vitro	in vitro διαγνωστικό
				
	Catálogo n.º	Katalognummer	Katalog nummer	Αριθμός καταλόγου
	No do lote	Lot nummer	Batch-nummer	Αριθμός Παρτίδος
		Indeholder tilstrækkeligt til "n" test	Innehåller tillräckligt till "n" tester	Περιεχόμενο επαρκές για «n» εξετάσεις
	Temperatura de conservação	Opbevarings-temperatur	Förvaringstemperatur	Θερμοκρασία αποθήκευσης
	Prazo de validade	Udløbsdato	Bäst före datum	Ημερομηνία λήξης
	Fabricante	Producent	Tillverkare	Κατασκευαστής
Distributed by				
Content	Conteúdo	Indhold	Innehåll	Περιεχόμενο
Volume/No.	Volume/Número	Volumen/antal	Volym/antal	Όγκος/αριθ..