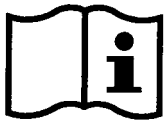
# Product information

Information about other products is available at: www.ibl-america.com



Useŕs Manual

# AMH ELISA

REF **IB79346**

Σ **96 wells**

**RUO** **For Research Use Only – Not for Use in Diagnostic Procedures**

Version 01-
06/20a/ DMC
Updated 200820

Manufactured for:
**Immuno-Biological Laboratories, Inc. (IBL-America)**
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**Please use only the valid version of the Instructions for Use provided with the kit.**

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## 1 INTENDED USE

The **IBL-America AMH ELISA** is an enzyme immunoassay for the measurement of Anti-Müllerian Hormone (AMH) in serum or plasma (EDTA or lithium heparin plasma). For research use only – Not for use in diagnostic procedures.

## 2 PRINCIPLE OF THE TEST

The IBL-America AMH ELISA is a solid phase enzyme-linked immunosorbent assay (ELISA) based on the **sandwich principle.** The microtiter wells are coated with a monoclonal (mouse) antibody directed towards a unique antigenic site of the AMH molecule. During the first incubation, the AMH in the added sample bind to the immobilized antibody. The simultaneously added enzyme conjugate, which contains an AMH antibody conjugated to horseradish peroxidase, binds to the AMH forming a sandwich complex. After a washing step to remove all unbound substances, the solid phase is incubated with the substrate solution. The colorimetric reaction is stopped by addition of stop solution, and optical density (OD) of the resulting yellow product is measured. The intensity of colour is proportional to the concentration of the analyte in the sample. A standard curve is constructed by plotting OD values against concentrations of standards, and concentrations of unknown samples are determined using this standard curve.

## 3 WARNINGS AND PRECAUTIONS

1. This kit is for research use only. Not for use in diagnostic procedures.
2. All reagents of this test kit which contain human serum or plasma have been tested and confirmed negative for HIV I/II, HBsAg and HCV by FDA approved procedures. All reagents, however, should be treated as potential biohazards in use and for disposal.
3. Before starting the assay, read the instructions completely and carefully. Use the valid version of instructions for use provided with the kit. Be sure that everything is understood.
4. The microplate contains snap-off strips. Unused wells must be stored at 2 °C to 8 °C in the sealed foil pouch and used in the frame provided.
5. Pipetting of samples and reagents must be done as quickly as possible and in the same sequence for each step.
6. Use reservoirs only for single reagents. This especially applies to the substrate reservoirs. Using a reservoir for dispensing a substrate solution that had previously been used for the conjugate solution may turn solution coloured. Do not pour reagents back into vials as reagent contamination may occur.
7. Mix the contents of the microplate wells thoroughly to ensure good test results. Do not reuse microwells.
8. Do not let wells dry during assay; add reagents immediately after completing the rinsing steps.
9. Allow the reagents to reach room temperature (20 °C to 26 °C) before starting the test. Temperature will affect the optical density readings of the assay. However, values for the samples will not be affected.
10. Never pipet by mouth and avoid contact of reagents and samples with skin and mucous membranes.
11. Do not smoke, eat, drink, or apply cosmetics in areas where samples or kit reagents are handled.
12. Wear disposable latex gloves when handling samples and reagents. Microbial contamination of reagents or samples may give false results.
13. Handling should be done in accordance with the procedures defined by an appropriate national biohazard safety guideline or regulation.
14. Do not use reagents beyond expiry date as shown on the kit labels.
15. All indicated volumes have to be performed according to the protocol. Optimal test results are only obtained when using calibrated pipettes and microtiter plate readers.
16. Do not mix or use components from kits with different lot numbers. It is advised not to exchange wells of different plates even of the same lot. The kits may have been shipped or stored under different conditions and the binding characteristics of the plates may result slightly different.
17. Avoid contact with Stop Solution containing 0.5 M $H_2SO_4$. It may cause skin irritation and burns.
18. Some reagents contain Proclin 300, BND and/or MIT as preservatives. In case of contact with eyes or skin, flush immediately with water.
19. TMB substrate has an irritant effect on skin and mucosa. In case of possible contact, wash eyes with an abundant volume of water and skin with soap and abundant water. Wash contaminated objects before reusing them. If inhaled, take the person to open air.

20. Chemicals and prepared or used reagents have to be treated as hazardous waste according to the national biohazard safety guideline or regulation.
21. For information on hazardous substances included in the kit please refer to Safety Data Sheets. Safety Data Sheets for this product are available upon request directly from IBL-America.

## 4 REAGENTS

### 4.1 Reagents provided

1. **SORB MT Microtiterwells**,12 x 8 (break apart) strips, 96 wells;
   Wells coated with anti-AMH antibody (monoclonal).
2. **CAL 0 – 5 Standard (Standard 0 - 5)**, 6 vials, 1 mL each, ready to use;
   Concentrations: 0.0 – 0.4 – 1.0 – 4.0 – 10 – 20 ng/mL
   Conversion: 1 ng/mL = 7.14 pmol/L
   The standards are calibrated against the following reference material: 1st WHO International Reference Reagent, Mullerian Inhibiting Substance/Anti-Mullerian Hormone, NIBSC code: 16/190
   Contain non-mercury preservative.
3. **CONTROL low & high Control Low & High**, 2 vials, 1 mL each, ready to use; For control values and ranges please refer to Certificate of Analysis. Contain non-mercury preservative.
4. **ENZ CONJ Enzyme Conjugate**, 1 vial, 14 mL, ready to use; Anti-AMH antibody conjugated with horseradish peroxidase; Contains non-mercury preservative.
5. **SUB TMB Substrate Solution**, 1 vial, 14 mL, ready to use; Tetramethylbenzidine (TMB).
6. **STOP SOLN Stop Solution**, 1 vial, 14 mL, ready to use; Contains 0.5 M $H_2SO_4$, Avoid contact with the stop solution. It may cause skin irritations and burns.
7. **WASH SOLN 40x Wash Solution**, 1 vial, 30 mL (40X concentrated); See “Reagent Preparation“.

**Note:** Additional Standard 0 for sample dilution is available upon request.

### 4.2 Materials required but not provided

- A calibrated microtiter plate reader (450 nm, with reference wavelength at 620 nm to 630 nm)
- Calibrated variable precision micropipettes
- Absorbent paper
- Distilled water
- Timer
- Graph paper or software for data reduction

### 4.3 Storage Conditions

When stored at 2 °C to 8 °C unopened reagents will retain reactivity until expiration date. Do not use reagents beyond this date. Opened reagents must be stored at 2 °C to 8 °C. Microtiter wells must be stored at 2 °C to 8 °C. Once the foil bag has been opened, care should be taken to close it tightly again. Opened kits retain activity for 8 weeks if stored as described above.

### 4.4 Reagent Preparation

Bring all reagents and required number of strips to room temperature (20 °C to 26 °C) prior to use.

**Wash Solution**

Add distilled water to the 40X concentrated Wash Solution. Dilute 30 mL of concentrated Wash Solution with 1170 mL distilled water to a final volume of 1200mL. The diluted Wash Solution is stable for 2 weeks at room temperature.

### 4.5 Disposal of the Kit

The disposal of the kit and all used materials/reagents must be performed according to the national regulations. Special information for this product is given in the Safety Data Sheet, section 13.

### 4.6 Damaged Test Kits

In case of any damage to the test kit or components, IBL-America must be informed in writing, at the latest one week after receiving the kit. Damaged single components should not be used for a test run. They have to be stored until a final solution has been found. After this, they should be disposed of according to the official regulations.

## 5 SAMPLE COLLECTION AND PREPARATION

Serum or plasma (EDTA or lithium heparin) can be used in this assay.
**Note**: Samples containing sodium azide should not be used in the assay.
In general, it should be avoided to use haemolytic, icteric, or lipaemic samples. For further information refer to chapter “Interfering Substances”.

### 5.1 Sample Collection

**Serum:**
Collect blood by venipuncture (e.g. Sarstedt Monovette for serum), allow to clot, and separate serum by centrifugation at room temperature. Do not centrifuge before complete clotting has occurred. Individuals receiving anticoagulant therapy may require increased clotting time.

**Plasma:**
Whole blood should be collected into centrifuge tubes containing anticoagulant (e.g. Sarstedt Monovette with the appropriate plasma preparation) and centrifuged immediately after collection.

### 5.2 Sample Storage and Preparation

Samples should be capped and may be stored for up to 7 days at 2 °C to 8 °C prior to assaying.
Samples stored for a longer time (up to 3 months) should be frozen only once at -20 °C prior to assay.
Thawed samples should be inverted several times prior to testing.

### 5.3 Sample Dilution

If in an initial assay, a sample is found to contain more analyte than the highest standard, the sample can be diluted with Standard 0 and re-assayed as described in “Assay Procedure”.
For the calculation of the concentrations this dilution factor has to be taken into account.

Example:
a) dilution 1:10: 10 µL sample + 90 µL Standard 0 (mix thoroughly)
b) dilution 1:100: 10 µL dilution a) 1:10 + 90 µL Standard 0 (mix thoroughly).

## 6 ASSAY PROCEDURE

### 6.1 General Remarks

- All reagents and samples must be allowed to come to room temperature before use. All reagents must be mixed without foaming.
- Once the test has been started, all steps should be completed without interruption.
- Use new disposal plastic pipette tips for each standard, control or sample in order to avoid cross contamination.
- Optical density is a function of the incubation time and temperature. Before starting the assay, it is recommended that all reagents are ready, caps removed, all needed wells secured in holder, etc. This will ensure equal elapsed time for each pipetting step without interruption.
- As a general rule the enzymatic reaction is linearly proportional to time and temperature.

### 6.2 Test Procedure

Each run must include a standard curve.

1. Secure the desired number of Microtiter wells in the frame holder.
2. Dispense **25 µL** of each **Standard, Control** and **sample** with new disposable tips into appropriate wells.
3. Dispense **100 µL Enzyme Conjugate** into each well.
   Thoroughly mix for 10 seconds. It is important to have a complete mixing in this step.
4. Incubate for **60 minutes** at room temperature.
5. Rinse the wells **3 times** with **400 µL** diluted Wash Solution per well, if a plate washer is used.
   - OR -
   Briskly shake out the contents of the wells.
   Rinse the wells **3 times** with **300 µL** diluted Wash Solution per well for manual washing. Strike the wells sharply on absorbent paper to remove residual droplets.
   **Important note:**
   The sensitivity and precision of this assay is markedly influenced by the correct performance of the washing procedure!
6. Add **100 µL** of **Substrate Solution** to each well.
7. Incubate for **15 minutes** at room temperature.
8. Stop the enzymatic reaction by adding **50 µL** of **Stop Solution** to each well.
9. Determine the optical density of the solution in each well at **450 nm (reading) and at 620 nm to 630 nm (background subtraction, recommended)** with a microtiter plate reader.
   It is recommended that the wells be read **within 10 minutes** after adding the Stop Solution.

### 6.3 Calculation of Results

1. Calculate the average optical density (OD) values for each set of standards, controls and samples.
2. Using graph paper, construct a standard curve by plotting the mean OD obtained from each standard against its concentration with OD value on the vertical (Y) axis and concentration on the horizontal (X) axis.
3. Using the mean OD value for each sample determine the corresponding concentration from the standard curve.
4. Automated method: The results in the Instructions for Use have been calculated automatically using a 4-Parameter curve fit. (4-Parameter Rodbard or 4-Parameter Marquardt are the preferred methods.) Other data reduction functions may give slightly different results.
5. The concentration of the samples can be read directly from this standard curve. Samples with concentrations higher than that of the highest standard have to be further diluted or reported as > 20 ng/mL. For the calculation of the concentrations this dilution factor has to be taken into account.

## 7 QUALITY CONTROL

Good laboratory practice requires that controls be run with each calibration curve. A statistically significant number of controls should be assayed to establish mean values and acceptable ranges to assure proper performance. It is recommended to use control samples according to state and federal regulations. The use of control samples is advised to assure the day to day validity of results. Use controls at both normal and pathological levels. The controls and the corresponding results of the QC-Laboratory are stated in the QC certificate added to the kit. The values and ranges stated on the QC sheet always refer to the current kit lot and should be used for direct comparison of the results.
It is also recommended to make use of national or international Quality Assessment programs in order to ensure the accuracy of the results. Employ appropriate statistical methods for analysing control values and trends. If the results of the assay do not fit to the established acceptable ranges of control materials results should be considered invalid. In this case, please check the following technical areas: Pipetting and timing devices; photometer, expiration dates of reagents, storage and incubation conditions, aspiration and washing methods. After checking the above mentioned items without finding any error contact your distributor or IBL-America directly.

## 8 PERFORMANCE CHARACTERISTICS

### 8.1 Assay Dynamic Range

The range of the assay is between 0.062 ng/mL – 20.0 ng/mL.

### 8.2 Specificity of Antibodies (Cross-Reactivity)

The following substances were tested for cross-reactivity of the assay:

| Substance | Added conc. (ng/mL) | Mean cross-reactivity (%) |
|---|---|---|
| AMH | 0.40 – 10 | 100 |
| Inhibin A | 2.0 – 2000 | 0.03 |
| Actvin AB | 2.0 – 2000 | 0.27 |
| LH | 2.0 – 2000 | 0.18 |
| FSH | 2.0 – 2000 | 0.26 |
| HCG | 2.0 – 2000 | 0.12 |
| TSH | 2.0 – 2000 | 0.39 |
| TGF-β1 | 2.0 – 2000 | 0.18 |
| TGF-β2 | 2.0 – 2000 | 0.18 |
| Prolactin | 2.0 – 2000 | 0.38 |

No substantial cross-reactivity of the assay to structurally related substances is detected.

### 8.3 Sensitivity

The analytical sensitivity of the IBL-America ELISA was calculated by adding 2 standard deviations to the mean of 20 replicate analyses of the Standard 0 and was found to be 0.044 ng/mL.

The Limit of Blank (LoB) is 0.044 ng/mL.
The Limit of Detection (LoD) is 0.052 ng/mL.
The Limit of Quantification (LoQ) is 0.062 ng/mL.

### 8.4 Reproducibility

#### 8.4.1 Intra-Assay

The within-assay variability was determined by measuring each sample 10 times per run (n = 10):

| Sample | n | Mean (ng/mL) | CV (%) |
|---|---|---|---|
| 1 | 10 | 0.22 | 7.3 |
| 2 | 10 | 0.67 | 3.1 |
| 3 | 10 | 5.76 | 2.4 |
| 4 | 10 | 15.88 | 2.6 |

#### 8.4.2 Inter-Assay

The between-assay variability was determined by measuring each sample 10 times per run for 3 days (n = 30):

| Sample | n | Mean (ng/mL) | CV (%) |
|---|---|---|---|
| 1 | 30 | 0.23 | 6.0 |
| 2 | 30 | 0.69 | 5.4 |
| 3 | 30 | 5.73 | 3.1 |
| 4 | 30 | 15.90 | 4.1 |

### 8.5 Recovery

Samples have been spiked by adding AMH solutions with known concentrations.
The recovery (%) was calculated by multiplying the ratio of measured and expected values with 100.

| | | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|---|---|---|---|---|---|
| **Concentration (ng/mL)** | | 0.20 | 0.70 | 5.85 | 15.89 |
| **Average Recovery (%)** | | 98.8 | 97.5 | 97.0 | 98.5 |
| **Range of Recovery (%)** | from | 94.2 | 95.4 | 94.1 | 95.9 |
| | to | 101.8 | 100.3 | 101.1 | 100.8 |

### 8.6 Linearity

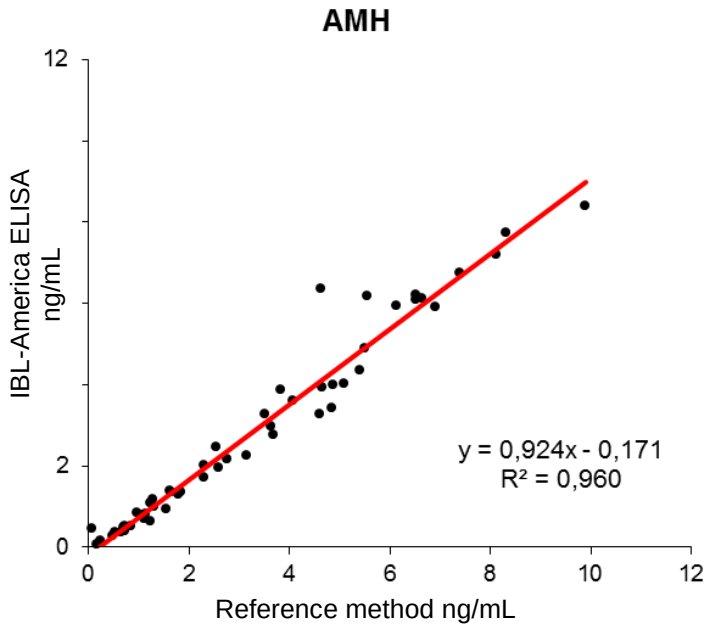
Samples were measured undiluted and in serial dilutions with standard 0. The recovery (%) was calculated by multiplying the ratio of expected and measured values with 100.

| | | Sample 1 | Sample 2 | Sample 3 | Sample 4 |
|---|---|---|---|---|---|
| **Concentration (ng/mL)** | | 3.01 | 5.99 | 9.75 | 16.29 |
| **Average Recovery (%)** | | 105.2 | 106.8 | 95.2 | 110.3 |
| **Range of Recovery (%)** | from | 94.7 | 102.4 | 94.5 | 106.6 |
| | to | 111.9 | 110.9 | 95.9 | 113.8 |

### 8.7 Comparison Studies

A comparison of the IBL-America AMH ELISA (IB79346) (y) and a reference (competing) method (x) using samples gave the following correlation:

n = 52
r = 0.980



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## 9 LIMITATIONS OF USE

Reliable and reproducible results will be obtained when the assay procedure is performed with a complete understanding of the package insert instruction and with adherence to good laboratory practice.
Any improper handling of samples or modification of this test might influence the results.

### 9.1 Interfering Substances

Haemoglobin (up to 4 mg/mL), Bilirubin (up to 0.5 mg/mL) and Triglyceride (up to 7.5 mg/mL) have no influence on the assay results.

### 9.2 Drug Interferences

Until today no substances (drugs) are known to us, which have a substantial influence to the measurement of AMH in a sample.

### 9.3 High-Dose-Hook Effect

Hook effect was not observed in this test up to a concentration of 400 ng/mL of AMH.

## 10 LEGAL ASPECTS

### 10.1 Reliability of Results

The test must be performed exactly as per the manufacturer's instructions for use. Moreover the user must strictly adhere to the rules of GLP (Good Laboratory Practice) or other applicable national standards and/or laws. This is especially relevant for the use of control reagents. It is important to always include, within the test procedure, a sufficient number of controls for validating the accuracy and precision of the test. The test results are valid only if all controls are within the specified ranges and if all other test parameters are also within the given assay specifications. In case of any doubt or concern please contact IBL-America.

### 10.2 Liability

Any modification of the test kit and/or exchange or mixture of any components of different lots from one test kit to another could negatively affect the intended results and validity of the overall test. Such modification and/or exchanges invalidate any claim for replacement. Regardless, in the event of any claim, the manufacturer's liability is not to exceed the value of the test kit. Any damage caused to the test kit during transportation is not subject to the liability of the manufacturer.

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## SYMBOLS USED WITH IBL-AMERICA ASSAYS

| Symbol | English | Deutsch | Francais | Espanol | Italiano |
|---|---|---|---|---|---|
| CE | European Conformity | CE-Konfirmitäts-kennzeichnung | Conforme aux normes européennes | Conformidad europea | Conformità europea |
|  | Consult instructions for use | Gebrauchsanweisung beachten | Consulter les instructions d'utilisation | Consulte las Instrucciones | Consultare le istruzioni per l'uso |
| IVD | In vitro diagnostic device | In-vitro-Diagnostikum | Ussage Diagnostic in vitro | Diagnóstico in vitro | Per uso Diagnostica in vitro |
| RUO | For research use only | Nur für Forschungszwecke | Seulement dans le cadre de recherches | Sólo para uso en investigación | Solo a scopo di ricerca |
| REF | Catalogue number | Katalog-Nr. | Référence | Número de catálogo | No. di Cat. |
| LOT | Lot. No. / Batch code | Chargen-Nr. | No. de lot | Número de lote | Lotto no |
|  | Contains sufficient for <n> tests/ | Ausreichend für "n" Ansätze | Contenu suffisant pour "n" tests | Contenido suficiente para <n> ensayos | Contenuto sufficiente per "n" saggi |
|  | Note warnings and precautions | Warnhinweise und Vorsichtsmaßnahmen beachten | Avertissements et mesures de précaution font attention | Tiene en cuenta advertencias y precauciones | Annoti avvisi e le precauzioni |
|  | Storage Temperature | Lagerungstemperatur | Temperature de conservation | Temperatura de conservacion | Temperatura di conservazione |
|  | Expiration Date | Mindesthaltbarkeits-datum | Date limite d'utilisation | Fecha de caducidad | Data di scadenza |
|  | Legal Manufacturer | Hersteller | Fabricant | Fabricante | Fabbricante |
| *Distributed by* | Distributor | Vertreiber | Distributeur | Distribuidor | Distributtore |