SYMBOL KEYS

Temperature Limitation	Consult Instructions for use	Date of Manufacture	Do not reuse
Manufacturer	IVD *In vitro* Diagnostic Medical Device	This side up	Use by
Contains sufficient for <n> tests	REF Catalogue Number	LOT Batch Number / Lot Number	DEVICE Device

Manufactured by:

Zephyr Biomedicals

A Division of Tulip Diagnostics (P) Ltd.

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0123/VER-02

FIAcheck™

AMH

Fluorescence Immunoassay for Quantitative determination of Anti Mullerian Hormone (AMH) in Human Serum

DEVICE

FOR IN VITRO DIAGNOSTIC USE ONLY

Store at 4°C to 30°C

INTENDED USE

FIAcheck™ AMH Fluorescence Immunoassay is intended for the in-vitro quantitative measurement of Anti Mullerian Hormone (AMH) in human serum.

INTRODUCTION

Anti-Mullerian hormone is a glycoprotein hormone structurally related to inhibin and activin from the transforming growth factor beta superfamily, whose key roles are in growth differentiation and folliculogenesis. AMH expression is critical to sex differentiation at a specific time during fetal development and appears to be tightly regulated by nuclear receptor SF1, transcription GATA factors, sex-reversal gene DAX1, and follicle-stimulating hormone (FSH). AMH is activated by SOX9 in the Sertoli cells of the male fetus thereby arresting the development of fallopian tubes, uterus, and upper vagina. AMH is also a product of granulosa cells of the prenatal and small antral follicles in women. As such, AMH is only present in the ovary until menopause. AMH level is also lower and even below the detection limit in women with premature ovarian failure of any cause, including after cancer chemotherapy, etc.

PRINCIPLE OF THE TEST

FIAcheck™ AMH is based on principle of agglutination of antibodies/anti-sera with respective antigen in immuno-chromatographic format using fluorophores as signal generators. The **FIAcheck™ AMH** test device is coated with immobilized AMH mouse monoclonal antibody 1 on the test line, sheep anti chicken IgY in control line and a mixture of AMH mouse monoclonal antibody 2 and Chicken IgY labeled with fluorescent microspheres on the binding pad.

AMH in sample binds to the AMH mouse monoclonal antibody 2 labeled with fluorescent microspheres in the binding pad.

The fluorescent labeled Ag-Ab complex moves forward due to capillary action and is captured by the immobilized AMH mouse monoclonal antibody 1 forming a double antibody sandwich and produces the test line. Chicken IgY labelled with fluorescent microspheres binds with sheep anti chicken IgY to produce the control line. When the **FIAcheck™** test device is inserted in the **FIAcheck™** analyzer, it scans both the test line and control line. The ratio of the two fluorescence values is used to calculate the concentration of the analyte present in the sample.

MATERIALS AND COMPONENTS

Materials provided with the test kits:

- **FIAcheck™ AMH** test device in a sealed pouch with desiccant.
- QR Code card for calibration.

Materials required but not provided

- Precision pipettes: 100µl
- Disposable pipette tips
- Disposable Gloves
- **FIAcheck™** Analyzer (Time Resolved Fluorescence Immunoassay Analyzer)
- **FIAcheck™** Incubator
- Digital Thermometer
- Stopwatch

STORAGE AND STABILITY

1. **FIAcheck™ AMH** kit is stable at 4°C to 30°C upto expiry date printed on the label. DO NOT FREEZE
2. **FIAcheck™** Test device should be used within 30 minutes once the foil pouch is opened.
3. If the colour of the desiccant has changed from blue to pink or colourless at the time of opening the pouch, kindly discard the device and use another device.

SAMPLE COLLECTION

1. Only Human Serum sample should be used. Other bodily fluids and samples may not give accurate result.
2. The test should be performed within 4 hours after the sample collection at room temperature.

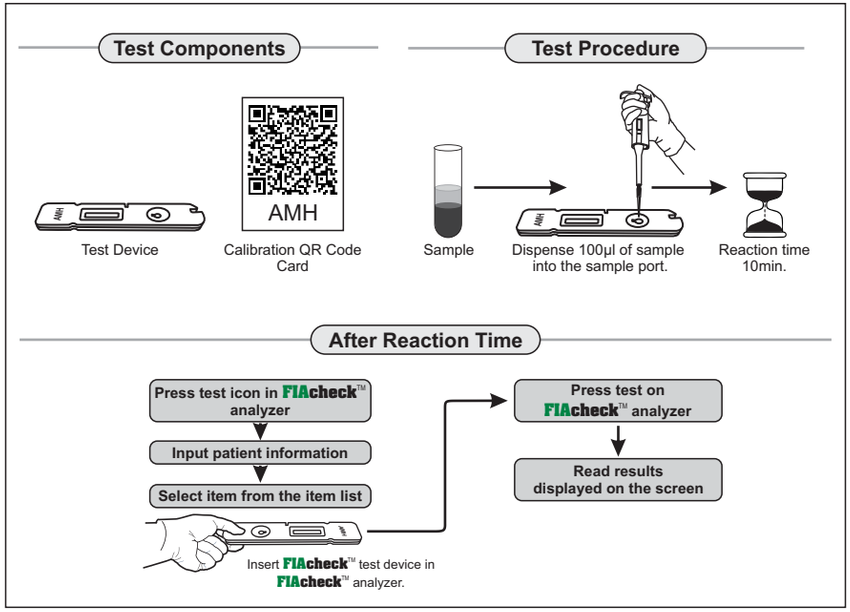
3. Avoid grossly hemolytic, lipemic or turbid samples.
4. Preferably use fresh samples. However, samples can be stored for 3 days at 2°C to 8°C, and if more than 3 days, they should be stored at -20°C.
5. The sample should be recovered to room temperature (18°C to 30°C) before testing. Avoid repeated freezing and thawing of samples as it can affect the test values.
6. Samples containing precipitate or particulate matter should be clarified by centrifugation prior to use.
7. Samples should be free from particulate matter and microbial contamination.

PRECAUTIONS

1. Only for In vitro diagnostic use.
2. After the test device is removed from the sealed pouch, it should be used immediately or within 30 minutes of opening the pouch.
3. Do not reuse the tested **FIAcheck**™ device.
4. Do not use damaged **FIAcheck**™ test device or pouch.
5. All samples should be considered potentially infectious and discarded appropriately as per Standard Bio-Safety guidelines.
6. Do not use kit after the expiry date.
7. Do not mix components of one kit with another.
8. Always use new tip for each sample.
9. Scan QR code card specific to the lot you are using.
10. Ambient temperature of testing environment directly impacts the accuracy of results. For incubation **FIAcheck**™ incubator is recommended, refer calibration QR code card for recommended incubation temperature.
11. The **FIAcheck**™ test device should be read immediately after the specified reaction time. Delay in reading might affect the accuracy of results.
12. The **FIAcheck**™ test device should be used only in conjunction with **FIAcheck**™ analyzer for accurate and reliable results.

TEST PROCEDURE

1. To calibrate the **FIAcheck**™ **AMH** kit, scan the QR code card provided with the kit.
2. Remove **FIAcheck**™ **AMH** test device from sealed pouch and place it horizontally on a clean table, label the device with sample identity.
3. Dispense 100 µl of sample onto the sample port of **FIAcheck**™ **AMH** test device.
4. Incubate in **FIAcheck**™ incubator for **10 minutes** refer calibration QR code card for recommended incubation temperature.
5. After 10 minutes, insert the test device immediately into the **FIAcheck**™ analyzer and read results.



Expected Range

The following reference interval was obtained after statistical analysis of the confidence interval for the tests of the content of AMH in serum samples of healthy people:
Male : 1.46-11.6ng/mL (95%) ;
Female 20-24 years old : 1.66-9.49ng/mL (95%);
Female 25-29 years old : 1.18-9.16ng/mL (95%);
Female 30-34 years old : 0.672-7.55ng/mL (95%);
Female 35-39 years old : 0.777-5.24ng/mL (95%);
Female 40-44 years old : 0.097-2.96ng/mL (95%);
Female 45-50 years old : 0.046-2.06ng/mL (95%).
It is recommended that each laboratory establish its own reference range for the population it serves.

PERFORMANCE CHARACTERISTICS

1. Measuring Range: 0.1-50ng/mL.
2. Lower Detection Limit: ≤0.1ng/mL.
3. Upper Detection Limit: ≥50ng/mL.
4. Accuracy: Based on comparison experiments, the relative standard deviation of ≤15%, and the correlation coefficient of r≥0.990 was observed.
5. Within-Run Precision: ≤15%.
6. Between-Run Precision: ≤15%.
7. Hook Test: No hook effect with high concentration sample. Hook test was conducted with reference material exceeding the upper limit of measuring range, and the detection result was greater than the upper limit of detection.
8. In an internal Study, **FIAcheck**™ **AMH** was evaluated against commercially available licensed kit with 100 random clinical samples and **FIAcheck**™ **AMH** has demonstrated 100% clinical correlation with the commercially available licensed kit.

AMH Levels	No. of samples	FIAcheck™ AMH	Qualisa AMH
Normal	80	80	80
Low	8	8	8
High	12	12	12

9. In an external Study, **FIAcheck**™ **AMH** has been evaluated by a NABL accredited lab against their reference method. In this evaluation **FIAcheck**™ **AMH** has demonstrated 100% correlation with the reference method.
*Data file: Zephyr Biomedicals (A Division of Tulip Diagnostics (P) Ltd).

LIMITATIONS OF THE ASSAY

1. As with all diagnostic tests, a definite clinical diagnosis should not be based on the results of a single test, but should only be made by the physician after all clinical and laboratory findings have been evaluated.
2. This kit is only for human serum.

WARRANTY

This product is designed to perform as described on the label and package insert. The manufacturer disclaims any implied warranty of use and sale for any other purpose. In the event of performance changes or product malfunction, please contact manufacturer.

BIBLIOGRAPHY

1. Xiong Ziwei, Hu Jian, Chen Xinyu, et al. The value of anti-Mullerian hormone in the diagnosis of polycystic ovary syndrome. Molecular Imaging Journal, 2015, 38(2): 80-83.
2. Wu Xueqing, Kong Rui, Tian Li, et al. Expert consensus on ovarian hyporesponsiveness. Reproductive and Contraception,2015,2: 71-79.
3. Tan Jiaqi, Chen Xiaoli, Li Yu, et al. Study on the value of anti-Mullerian hormone in predicting ovarian response. Journal of Practical Obstetrics and Gynecology, 2015, 31(8): 583-586.