SYMBOL KEYS

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

Manufactured by:

Zephyr Biomedicals

A Division of Tulip Diagnostics (P) Ltd.

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0624/VER-02

FIAcheck™

TT3

Fluorescence Immunoassay for Quantitative determination of TT3 (Total Triiodothyronine) in Human Serum and Plasma

DEVICE

FOR IN VITRO DIAGNOSTIC USE ONLY

Store at 4°C to 30°C

INTENDED USE

FIAcheck™ TT3 Fluorescence Immunoassay is intended for the in-vitro quantitative measurement of TT3 (Total Triiodothyronine) in human serum and plasma.

INTRODUCTION

The hormones thyroxine (T4) and triiodothyronine (T3) circulate in the bloodstream, mostly bound to the plasma protein, thyroxine binding globulin (TBG). The concentration of T3 is much less than that of T4, but its metabolic potency is much greater. T3 determination is an important factor in the diagnosis of thyroid disease. Its measurement has uncovered a variant of hyperthyroidism in thyrotoxic patients with elevated T3 values and normal T4 values. T3 determination is also useful in monitoring both patients under treatment for hyperthyroidism and patients who have discontinued anti-thyroid drug therapy. In addition to hyperthyroidism, T3 levels are elevated in women who are pregnant, and in women receiving oral contraceptives or estrogen treatment.

PRINCIPLE OF THE TEST

FIAcheck™ TT3 is based on principle of agglutination of antibodies/anti-sera with respective antigen in immuno-chromatographic format using fluorophores as signal generators. The **FIAcheck™ TT3** test device is coated with immobilized T3-BSA full antigen complex on the test line, sheep anti chicken IgY in control line and a mixture of T3 sheep monoclonal antibody and Chicken IgY labeled with fluorescent microspheres on the binding pad.

TT3 in sample binds to the T3 sheep monoclonal antibody labeled with fluorescent microspheres in the binding pad.

The fluorescent microspheres of unconjugated T3 binds to T3-BSA in the test area, forming the test band whose fluorescence intensity was negatively correlated with the concentration of T3 in the sample. 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™ TT3** test device in a sealed pouch with desiccant and special tip.
- QR Code card for calibration.
- Sample Diluent. Ready to use.
- Empty vials for sample dilution.

Materials required but not provided

- Precision pipettes: 100-1000µl
- Disposable pipette tips
- Disposable Gloves
- **FIAcheck™** Analyzer (Time Resolved Fluorescence Immunoassay Analyzer)
- Digital Thermometer
- Stopwatch

STORAGE AND STABILITY

1. **FIAcheck™ TT3** 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.
4. Once opened, the sample diluent can be stored between 4°C to 30°C for remaining duration of shelf life.

SAMPLE COLLECTION

1. Only Human Serum/Plasma sample should be used. Other bodily fluids and samples may not give accurate result.
2. Plasma can be anti-coagulated with Heparin and Sodium citrate or Tri sodium citrate under aseptic conditions.
3. The test should be performed within 4 hours after the sample collection at room temperature.

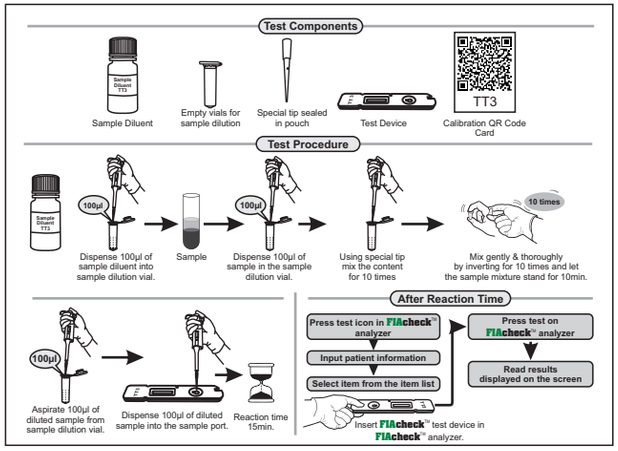
4. Avoid grossly hemolytic, lipemic or turbid samples.
5. Preferably use fresh samples. However, samples can be stored for 3 days at 2°C to 8°C, and if more than 3 days, they should be stored at -20°C.
6. The sample should be recovered to room temperature (18°C to 30°C) before testing. Avoid repeated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

PRECAUTIONS

1. Only for In vitro diagnostic use.
2. Bring all reagents and samples to room temperature before use.
3. After the test device is removed from the sealed pouch, it should be used immediately or within 30 minutes of opening the pouch.
4. Do not reuse the tested **FIAcheck**™ device. Do not use sample dilution vial for more than one sample.
5. Do not use damaged **FIAcheck**™ test device or pouch.
6. All samples should be considered potentially infectious and discarded appropriately as per Standard Bio-Safety guidelines.
7. Do not use kit after the expiry date.
8. Do not mix components of one kit with another.
9. Always use new tip for each sample and reagent.
10. Scan QR code card specific to the lot you are using.
11. Ambient temperature of testing environment directly impacts the accuracy of results. Ideal working temperature is 18°C to 30°C.
12. It is highly recommended to mix the sample diluent and sample mixture thoroughly by gently inverting the vial 10 times. (Refer pictorial presentation.)
13. It is not recommended to use the sample diluent and sample mixture beyond specified time.
14. The **FIAcheck**™ test device should be read immediately after the specified reaction time. Delay in reading might affect the accuracy of results.
15. 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**™ **TT3** kit, scan the QR code card provided with the kit.
2. Dispense **100** µl of sample diluent into the empty sample dilution vial.
3. Dispense 100 µl of sample in sample dilution vial.
4. Using the special tip mix the content for 10 times.
   Note: It is required to use the special tip which is sealed in sealed pouch.
5. Close the lid of the sample dilution vial, label with sample identity and mix the content of the vial by gently inverting it for 10 times, let it stand for 10 minutes.
6. Remove **FIAcheck**™ **TT3** test device from sealed pouch and place it horizontally on a clean table, label the device with sample identity.
7. Dispense 100 µl of the above mixture at the sample port in the **FIAcheck**™ **TT3** test device.
8. Incubate at room temperature (18°C to 30°C) for **15 minutes**.
9. After 15 minutes, insert the test device immediately into the **FIAcheck**™ analyzer and read results.



Expected Range
Reference Range: 1.3 - 2.7nmol/L
It is recommended that each laboratory establish its own reference range for the population it serves.

PERFORMANCE CHARACTERISTICS

1. Measuring Range: 0.5-10nmol/L.
2. Lower Detection Limit: ≤0.5nmol/L.
3. Upper Detection Limit: ≥10nmol/L.
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**™ **TT3** was evaluated against commercially available licensed kit with 100 random clinical samples and **FIAcheck**™ **TT3** has demonstrated 100% clinical correlation with the commercially available licensed kit.

TT3 Levels	No. of samples	FIAcheck™ TT3	EIA TT3
Normal	80	80	80
Low	8	8	8
High	12	12	12

9. In an external Study, **FIAcheck**™ **TT3** has been evaluated by a NABL accredited lab against their reference method. In this evaluation **FIAcheck**™ **TT3** 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 and plasma.

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

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