

QuantiFERON[®]-TB Gold

**The Whole Blood IFN-gamma Test
Measuring Responses to
ESAT-6, CFP-10 & TB7.7 Peptide Antigens**

PACKAGE INSERT

For *In Vitro* Diagnostic Use



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1. INTENDED USE

QuantIFERON®-TB Gold In-Tube (QFT®) is an *in vitro* diagnostic test using a peptide cocktail simulating ESAT-6, CFP-10 and TB7.7(p4) proteins to stimulate cells in heparinised whole blood. Detection of interferon- γ (IFN- γ) by Enzyme-Linked Immunosorbent Assay (ELISA) is used to identify *in vitro* responses to these peptide antigens that are associated with *Mycobacterium tuberculosis* infection.

QFT is an indirect test for *M. tuberculosis* infection (including disease) and is intended for use in conjunction with risk assessment, radiography and other medical and diagnostic evaluations.

2. SUMMARY AND EXPLANATION OF THE TEST

Tuberculosis is a communicable disease caused by infection with *M. tuberculosis* complex organisms (*M. tuberculosis*, *M. bovis*, *M. africanum*), which typically spreads to new hosts via airborne droplet nuclei from patients with respiratory tuberculosis disease. A newly infected individual can become ill from tuberculosis within weeks to months, but most infected individuals remain well. Latent tuberculosis infection (LTBI), a non-communicable asymptomatic condition, persists in some, who might develop tuberculosis disease months or years later. The main purpose of diagnosing LTBI is to consider medical treatment for preventing tuberculosis disease. Until recently the tuberculin skin test (TST) was the only available method for diagnosing LTBI. Cutaneous sensitivity to tuberculin develops from 2 to 10 weeks after infection. However, some infected individuals, including those with a wide range of conditions hindering immune functions, but also others without these conditions, do not respond to tuberculin. Conversely, some individuals who are unlikely to have *M. tuberculosis* infection exhibit sensitivity to tuberculin and have positive TST results after vaccination with bacille Calmette-Guérin (BCG), infection with mycobacteria other than *M. tuberculosis* complex, or undetermined other factors.

LTBI must be distinguished from tuberculosis disease, a reportable condition which usually involves the lungs and lower respiratory tract, although other organ systems may also be affected. Tuberculosis disease is diagnosed from historical, physical, radiological, histological, and mycobacteriological findings.

The QFT test is a test for Cell Mediated Immune (CMI) responses to peptide antigens that simulate mycobacterial proteins. These proteins, ESAT-6, CFP-10 and TB7.7(p4), are absent from all BCG strains and from most non-tuberculosis mycobacteria with the exception of *M. kansasii*, *M. szulgai* and *M. marinum*.¹ Individuals infected with *M. tuberculosis* complex organisms usually have lymphocytes in their blood that recognise these and other mycobacterial antigens. This recognition process involves the generation and secretion of the cytokine, IFN- γ . The detection and subsequent quantification of IFN- γ forms the basis of this test.

The antigens used in QFT are a peptide cocktail simulating the proteins ESAT-6, CFP-10 and TB7.7(p4). Numerous studies have demonstrated that these peptide antigens stimulate IFN- γ responses in T-cells from individuals infected with *M. tuberculosis* but generally not from uninfected or BCG vaccinated persons without disease or risk for LTBI.¹⁻³² However, medical treatments or conditions that impair immune functionality can potentially reduce IFN- γ responses. Patients with certain other mycobacterial infections might also be responsive to ESAT-6, CFP-10 and TB7.7(p4) as the genes encoding these proteins are present in *M. kansasii*, *M. szulgai* and *M. marinum*.^{1,23} The QFT test is both a test for LTBI and a helpful aid for diagnosing *M. tuberculosis* complex infection in sick patients. A positive result supports the diagnosis of tuberculosis disease; however, infections by other mycobacteria (e.g., *M. kansasii*) could also lead to positive results. Other medical and diagnostic evaluations are necessary to confirm or exclude tuberculosis disease.

Principles of the Assay

The QFT system uses specialised blood collection tubes, which are used to collect whole blood. Incubation of the blood occurs in the tubes for 16 to 24 hours, after which, plasma is harvested and tested for the presence of IFN- γ produced in response to the peptide antigens.

The QFT test is performed in two stages. First, whole blood is collected into each of the QuantiFERON[®]-TB Gold blood collection tubes, which include a Nil Control tube, TB Antigen tube, and a Mitogen Control tube.

The Mitogen tube is used with the QFT test as a control for correct blood handling and incubation and serves as a control especially where there is doubt as to the individual's immune status and thus ability to respond in the test.

The tubes should be incubated at 37°C as soon as possible, and within 16 hours of collection. Following a 16 to 24 hour incubation period, the tubes are centrifuged, the plasma is removed and the amount of IFN- γ (IU/mL) measured by ELISA.

A test is considered positive for an IFN- γ response to the TB Antigen tube that is significantly above the Nil IFN- γ IU/mL value. The Mitogen-stimulated plasma sample serves as an IFN- γ positive control for each specimen tested. A low response to Mitogen (<0.5 IU/mL) indicates an indeterminate result when a blood sample also has a negative response to the TB antigens. This pattern may occur with insufficient lymphocytes, reduced lymphocyte activity due to improper specimen handling, incorrect filling/mixing of the Mitogen tube, or inability of the patient's lymphocytes to generate IFN- γ . The Nil sample adjusts for background, heterophile antibody effects, or non-specific IFN- γ in blood samples. The IFN- γ level of the Nil tube is subtracted from the IFN- γ level for the TB Antigen tube and Mitogen tube.

Time Required for Performing Assay

The time required to perform the QFT assay is estimated below; the time of testing multiple samples when batched is also indicated:

37°C Incubation of blood tubes: 16 to 24 hours

ELISA: Approx. 3 hours for one ELISA plate

- <1 hour labor
- Add 10-15 minutes for each extra plate

3. REAGENTS AND STORAGE

Blood Collection Tubes

Catalogue Number: T0590 0301

- | | |
|---|-------------|
| 1. Nil Control (Grey cap with white ring) | 100 x tubes |
| 2. TB Antigen (Red cap with white ring) | 100 x tubes |
| 3. Mitogen Control (Purple cap with white ring) | 100 x tubes |

High Altitude (HA) Blood Collection Tubes (for use above 1,020 metres)

Catalogue Number: T0590 0505 (refer Section 5)

- | | |
|--|-------------|
| 4. Nil Control (Grey cap with yellow ring) | 100 x tubes |
| 5. TB Antigen (Red cap with yellow ring) | 100 x tubes |
| 6. Mitogen Control (Purple cap with yellow ring) | 100 x tubes |

Catalogue Number: T0597 0403

- | | |
|---|-------------|
| 7. Dispenser Pack (4 x 25 Single Patient Packs) | 300 x tubes |
|---|-------------|

ELISA Components

<i>ELISA Components</i>	Catalogue No: 0594-0201	Catalogue No: 0594-0501
	2-plate kit	Reference Lab Pack
1. Microplate strips coated with murine anti-human IFN- γ monoclonal antibody	2 x 96 well plates	20 x 96 well plates
2. Human IFN- γ Standard, lyophilized (<i>contains recombinant human IFN-γ, bovine casein, 0.01% w/v Thimerosal</i>)	1 x vial (8 IU/mL when reconstituted)	10 x vials (8 IU/mL when reconstituted)
3. Green Diluent (<i>contains bovine casein, normal mouse serum, 0.01% w/v Thimerosal</i>)	1 x 30mL	10 x 30mL
4. Conjugate 100X Concentrate, lyophilized (<i>murine anti-human IFN-γ HRP, contains 0.01% w/v Thimerosal</i>)	1 x 0.3mL (when reconstituted)	10 x 0.3mL (when reconstituted)
5. Wash Buffer 20X Concentrate (<i>pH 7.2, contains 0.01% w/v Thimerosal</i>)	1 x 100mL	10 x 100mL
6. Enzyme Substrate Solution (<i>contains H₂O₂, 3,3',5,5' Tetramethylbenzidine</i>)	1 x 30mL	10 x 30mL
7. Enzyme Stopping Solution (<i>contains 0.5M H₂SO₄</i>)	1 x 15mL	10 x 15mL

Materials Required (but not provided)

- 37°C incubator. CO₂ not required.
- Calibrated variable-volume pipettes for delivery of 10µL to 1000µL with disposable tips.
- Calibrated multichannel pipette capable of delivering 50µL and 100µL with disposable tips.
- Microplate shaker.
- Deionised or distilled water - 2L.
- Microplate washer (automated washer recommended).
- Microplate reader fitted with 450nm filter and 620nm to 650nm reference filter.

Storage Instructions

Blood Collection Tubes

- Store blood collection tubes at 4°C to 25°C.

Kit Reagents

- Store kit refrigerated at 2°C to 8°C.
- Always protect Enzyme Substrate Solution from direct sunlight.

Reconstituted and Unused Reagents

For instructions on how to reconstitute the reagents, please see Section 6 (page 11).

- The reconstituted Kit Standard may be kept for up to 3 months if stored at 2°C to 8°C.
 - *Note the date the Kit Standard was reconstituted.*
- Once reconstituted, unused Conjugate 100X concentrate must be returned to storage at 2°C to 8°C and must also be used within 3 months.
 - *Note the date the Conjugate was reconstituted.*
- Working strength Conjugate must be used within 6 hours of preparation.
- Working strength Wash Buffer may be stored at room temperature for up to 2 weeks.

4. WARNINGS AND PRECAUTIONS

Warnings

- A negative QFT result does not preclude the possibility of *M. tuberculosis* infection or tuberculosis disease: false-negative results can be due to stage of infection (e.g., specimen obtained prior to the development of cellular immune response), co-morbid conditions which affect immune functions, incorrect handling of the blood collection tubes following venipuncture, incorrect performance of the assay, or other immunological variables.
- A positive QFT result should not be the sole or definitive basis for determining infection with *M. tuberculosis*. Incorrect performance of the assay may cause false positive responses.
- A positive QFT result should be followed by further medical evaluation and diagnostic evaluation for active tuberculosis disease (e.g., AFB smear and culture, chest x-ray).
- While ESAT-6, CFP-10 and TB7.7(p4) are absent from all BCG strains and from most known non-tuberculous mycobacteria, it is possible that a positive QFT result may be due to infection by *M. kansasii*, *M. szulgai* or *M. marinum*. If such infections are suspected, alternative tests should be investigated.

Precautions

- **For *in vitro* diagnostic use.**
- **Harmful: Enzyme Substrate Solution** contains 3,3',5,5' Tetramethylbenzidine that is harmful by ingestion, inhalation and skin contact. Skin and eye irritant. Mutagen. Use eye protection, wear gloves and handle as a potential carcinogen.
- **Harmful: Enzyme Stopping Solution** contains H₂SO₄ that is harmful by ingestion, eye contact, skin contact, and inhalation. Use eye protection, wear gloves and normal laboratory protective clothing. If the stopping solution contacts the skin or eyes, flush with copious quantities of water and seek medical attention.
- **Harmful: IFN- γ Standard and Conjugate 100X Concentrate** may be discomforting if ingested and may cause skin irritation. Wear gloves and normal laboratory protective clothing.
- **Handle human blood as if potentially infectious.** Observe relevant blood handling guidelines.
- **Thimerosal** is used as a preservative in some reagents. It may be toxic upon ingestion, inhalation or skin contact.
- **Green Diluent** contains normal mouse serum and casein, which may trigger allergic responses; avoid contact with skin.
- Deviations from the directions for use in the Package Insert may yield erroneous results. Please read the instructions carefully before use.
- Do not use kit if any reagent bottle shows signs of damage or leakage prior to use.
- Do not mix or use ELISA reagents from other QFT kit batches.
- Discard unused reagents and biological samples in accordance with Local, State, and Federal regulations.
- Do not use the blood collection tubes or ELISA kit after the expiry date.
- Ensure that laboratory equipment such as plate washers and readers have been calibrated/validated for use.

5. SPECIMEN COLLECTION AND HANDLING

QFT uses the following collection tubes:

1. Nil Control (Grey cap with white ring) (use between sea-level and 810m)
2. TB Antigen (Red cap with white ring) (use between sea-level and 810m)
3. Mitogen Control (Purple cap with white ring) (use between sea-level and 810m)
4. Nil Control (Grey cap with yellow ring) (use between 1020m and 1875m)
5. TB Antigen (Red cap with yellow ring) (use between 1020m and 1875m)
6. Mitogen Control - (Purple cap with yellow ring) (use between 1020m and 1875m)

Antigens have been dried onto the inner wall of the blood collection tubes so it is essential that the contents of the tubes be thoroughly mixed with the blood. The tubes must be transferred to a 37°C incubator as soon as possible and within 16 hours of collection.

The following procedures should be followed for optimal results:

1. For each subject collect 1mL of blood by venipuncture directly into each of the QFT blood collection tubes. This procedure should be performed by a trained phlebotomist.
 - Standard QFT blood collection tubes should be used up to an altitude of 810 metres. High Altitude (HA) QFT blood collection tubes should be used at altitudes between 1020 and 1875 metres.
 - *If using QFT blood collection tubes outside these altitude ranges, or if low blood draw volume does occur, blood can be collected using a syringe and 1mL transferred to each of the three tubes. For safety reasons, this is best performed by removing the syringe needle, **ensuring appropriate safety procedures**, removing the caps from the three QFT tubes and adding 1mL of blood to each (to the black mark on the side of the tube label). Replace the caps securely and mix as described below.*
 - As 1mL tubes draw blood relatively slowly, keep the tube on the needle for 2-3 seconds once the tube appears to have completed filling, to ensure that the correct volume is drawn.

The black mark on the side of the tubes indicates the 1mL fill volume. QFT blood collection tubes have been validated for volumes ranging from 0.8mL to 1.2mL. If the level of blood in any tube is not close to the indicator line, it is recommended to obtain another blood sample.

- If a “butterfly needle” is being used to collect blood, a “purge” tube should be used to ensure that the tubing is filled with blood prior to the QFT tubes being used.
2. Immediately after filling tubes, shake them ten (10) times just firmly enough to ensure the entire inner surface of the tube is coated with blood, to solubilize antigens on tube walls.
 - Tubes should be between 17°C - 25°C at the time of blood filling.
 - Over-energetic shaking may cause gel disruption and could lead to aberrant results.
 3. Label tubes appropriately.
 4. The tubes must be transferred to a 37°C ± 1°C incubator as soon as possible, and within 16 hours of collection. Prior to incubation, maintain the tubes at room temperature (22°C ± 5°C). Do not refrigerate or freeze the blood samples.

6. DIRECTIONS FOR USE

Stage One – Incubation of Blood and Harvesting of Plasma

Materials Provided

QFT blood collection tubes (Refer to Section 3).

Materials Required (but not provided)

Refer to Section 3.

Procedure

1. If the blood is not incubated immediately after collection, **re-mixing of the tubes by inverting 10 times must be performed immediately prior to incubation.**
2. Incubate the tubes **UPRIGHT** at 37°C for 16 to 24 hours. The incubator does not require CO₂ or humidification.
3. Blood collection tubes may be held between 4°C and 27°C for up to 3 days prior to centrifugation.
4. After incubation of the tubes at 37°C, harvesting of plasma is facilitated by centrifuging tubes for 15 minutes at 2000 to 3000 RCF (g). The gel plug will separate the cells from the plasma. If this does not occur, the tubes should be centrifuged again at a higher speed.
 - It is possible to harvest the plasma without centrifugation, however, additional care is required to remove the plasma without disturbing the cells.
5. **After centrifugation, avoid pipetting up and down or mixing plasma by any means prior to harvesting. At all times, take care not to disturb material on the surface of the gel.**
 - Plasma samples should only be harvested **using a pipette.**
 - Plasma samples can be loaded directly from centrifuged blood collection tubes into the QFT ELISA plate, including when automated ELISA workstations are used.
 - Plasma samples can be stored for up to 28 days at 2°C to 8°C or, if harvested, below -20°C for extended periods.

Stage Two - Human IFN- γ ELISA

Materials Provided

QFT ELISA kit (Refer to Section 3).

Materials Required (but not provided)

Refer to Section 3.

Procedure

1. All plasma samples and reagents, except for Conjugate 100X Concentrate, must be brought to room temperature ($22^{\circ}\text{C} \pm 5^{\circ}\text{C}$) before use. Allow at least 60 minutes for equilibration.
2. Remove strips that are not required from the frame, reseal in the foil pouch, and return to the refrigerator for storage until required.

Allow at least one strip for the QFT Standards and sufficient strips for the number of subjects being tested (refer to Figure 2, page 15 for recommended plate layout). After use, retain frame and lid for use with remaining strips.

3. Reconstitute the freeze dried Kit Standard with the volume of deionised or distilled water indicated on the label of the Standard vial. Mix gently to minimise frothing and ensure complete solubilisation. Reconstitution of the Standard to the stated volume will produce a solution with a concentration of 8.0 IU/mL.

Note: The reconstitution volume of the Kit Standard will differ between batches.

Use the reconstituted Kit Standard to produce a 1 in 4 dilution series of IFN- γ in Green Diluent (GD) – refer to Figure 1. S1 (Standard 1) contains 4 IU/mL, S2 (Standard 2) contains 1 IU/mL, S3 (Standard 3) contains 0.25 IU/mL, and S4 (Standard 4) contains 0 IU/mL (GD alone). The standards should be assayed at least in duplicate.

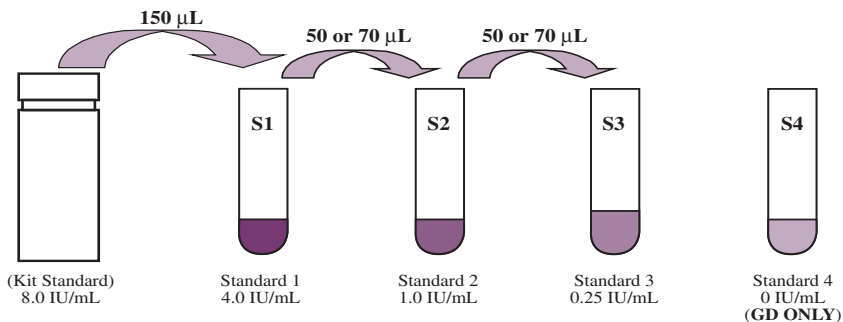
RECOMMENDED PROCEDURE FOR DUPLICATE STANDARDS

- Label 4 tubes "S1", "S2", "S3", "S4".
- Add **150µL** of GD to S1, S2, S3, S4.
- Add **150µL** of the Kit Standard to S1 and mix thoroughly.
- Transfer **50µL** from S1 to S2 and mix thoroughly.
- Transfer **50µL** from S2 to S3 and mix thoroughly.
- GD alone** serves as the zero standard (S4).

RECOMMENDED PROCEDURE FOR TRIPPLICATE STANDARDS

- Label 4 tubes "S1", "S2", "S3", "S4".
- Add **150µL** of GD to S1.
- Add **210µL** of GD to S2, S3, S4.
- Add **150µL** of the Kit Standard to S1 and mix thoroughly.
- Transfer **70µL** from S1 to S2 and mix thoroughly.
- Transfer **70µL** from S2 to S3 and mix thoroughly.
- GD alone** serves as the zero standard (S4).

FIGURE 1. Preparation of Standard Curve



- Prepare fresh dilutions of the Kit Standard for each ELISA session.
4. Reconstitute freeze dried Conjugate 100X Concentrate with 0.3mL of deionised or distilled water. Mix gently to minimize frothing and ensure complete solubilisation of the Conjugate.
- Working Strength conjugate is prepared by diluting the required amount of reconstituted Conjugate 100X Concentrate in Green Diluent as set out in Table 1 - Conjugate Preparation.

TABLE 1. Conjugate Preparation

NUMBER OF STRIPS	VOLUME OF CONJUGATE 100X CONCENTRATE	VOLUME OF GREEN DILUENT
2	10 μ L	1.0mL
3	15 μ L	1.5mL
4	20 μ L	2.0mL
5	25 μ L	2.5mL
6	30 μ L	3.0mL
7	35 μ L	3.5mL
8	40 μ L	4.0mL
9	45 μ L	4.5mL
10	50 μ L	5.0mL
11	55 μ L	5.5mL
12	60 μ L	6.0mL

- Mix thoroughly but gently to avoid frothing.
 - Return any unused Conjugate 100X Concentrate to 2°C to 8°C immediately after use.
 - Use only Green Diluent.
5. For plasma samples harvested from blood collection tubes and subsequently frozen or stored for more than 24 hours prior to assay, thoroughly mix before addition to the ELISA well.
 - If plasma samples are to be added directly from the centrifuged QFT tubes, any mixing of the plasma should be avoided.
 6. Add 50 μ L of freshly prepared Working Strength conjugate to the required ELISA wells using a multichannel pipette.
 7. Add 50 μ L of test plasma samples to appropriate wells using a multichannel pipette (Refer to recommended plate layout on page 15 – Figure 2). Finally, add 50 μ L each of the Standards 1 to 4.

**FIGURE 2. Recommended Sample Layout for Nil, TB Antigen & Mitogen Tubes
(28 tests per plate)**

Row	1	2	3	4	5	6	7	8	9	10	11	12
A	1N	1A	1M	S1	S1	S1	13N	13A	13M	21N	21A	21M
B	2N	2A	2M	S2	S2	S2	14N	14A	14M	22N	22A	22M
C	3N	3A	3M	S3	S3	S3	15N	15A	15M	23N	23A	23M
D	4N	4A	4M	S4	S4	S4	16N	16A	16M	24N	24A	24M
E	5N	5A	5M	9N	9A	9M	17N	17A	17M	25N	25A	25M
F	6N	6A	6M	10N	10A	10M	18N	18A	18M	26N	26A	26M
G	7N	7A	7M	11N	11A	11M	19N	19A	19M	27N	27A	27M
H	8N	8A	8M	12N	12A	12M	20N	20A	20M	28N	28A	28M

- S1 (Standard 1), S2 (Standard 2), S3 (Standard 3), S4 (Standard 4).
 - 1N (Sample 1. Nil Control plasma); 1A (Sample 1. TB Antigen plasma); 1M (Sample 1. Mitogen Control plasma).
8. Mix the conjugate and plasma samples/standards thoroughly using a microplate shaker for 1 minute.
 9. Cover each plate with a lid and incubate at room temperature ($22^{\circ}\text{C} \pm 5^{\circ}\text{C}$) for 120 ± 5 minutes.
 - Plates should not be exposed to direct sunlight during incubation.
 10. During the incubation, dilute one part Wash Buffer 20X Concentrate with 19 parts deionised or distilled water and mix thoroughly. Sufficient Wash Buffer 20X Concentrate has been provided to prepare 2L of Working Strength wash buffer.

Wash wells with **400 μL** of Working Strength wash buffer for at least 6 cycles. An automated plate washer is recommended.

- Thorough washing is very important to the performance of the assay. Ensure each well is **completely filled** with wash buffer to the top of the well for each wash cycle. A soak period of at least 5 seconds between each cycle is recommended.
- Standard laboratory disinfectant should be added to the effluent reservoir, and established procedures followed for the decontamination of potentially infectious material.

11. Tap plates face down on absorbent towel to remove residual wash buffer. Add 100 μ L of Enzyme Substrate Solution to each well and mix thoroughly using a microplate shaker.
12. Cover each plate with a lid and incubate at room temperature ($22^{\circ}\text{C} \pm 5^{\circ}\text{C}$) for 30 minutes.
 - Plates should not be exposed to direct sunlight during incubation.
13. Following the 30 minute incubation, add 50 μ L of Enzyme Stopping Solution to each well and mix.
 - Enzyme Stopping Solution should be added to wells in the same order and at approximately the same speed as the substrate in step 11.
14. Measure the Optical Density (OD) of each well within 5 minutes of stopping the reaction using a microplate reader fitted with a 450nm filter and with a 620nm to 650nm reference filter. OD values are used to calculate results.

7. CALCULATIONS AND TEST INTERPRETATION

QFT Analysis Software, used to analyse raw data and calculate results, is available from Cellestis (check that the most current version of the software is used).

The software performs a Quality Control assessment of the assay, generates a standard curve and provides a test result for each subject, as detailed in the Interpretation of Results section.

As an alternative to using the QFT Analysis Software, results can be determined according to the following method:

Generation of Standard Curve

(if QFT Analysis Software is not used)

Determine the mean OD values of the Kit Standard replicates on each plate.

Construct a $\log_{(e)}\text{-}\log_{(e)}$ standard curve by plotting the $\log_{(e)}$ of the mean OD (y-axis) against the $\log_{(e)}$ of the IFN- γ concentration of the standards in IU/mL (x-axis), omitting the zero standard from these calculations. Calculate the line of best fit for the standard curve by regression analysis.

Use the standard curve to determine the IFN- γ concentration (IU/mL) for each of the test plasma samples, using the OD value of each sample.

These calculations can be performed using software packages available with microplate readers, and standard spreadsheet or statistical software (such as Microsoft Excel). It is recommended that these packages be used to calculate the regression analysis, the coefficient of variation (%CV) for the standards, and the correlation coefficient (r) of the standard curve.

Quality Control of the Test

The accuracy of test results is dependent on the generation of an accurate standard curve. Therefore, results derived from the standards must be examined before test sample results can be interpreted.

For the ELISA to be valid:

- **The mean OD value for Standard 1 must be ≥ 0.600 .**
- **The %CV for Standard 1 and Standard 2 replicate OD values must be $\leq 15\%$.**
- **Replicate OD values for Standard 3 and Standard 4 must not vary by more than 0.040 optical density units from their mean.**
- **The correlation coefficient (r) calculated from the mean absorbance values of the standards must be ≥ 0.98 .**

The QFT Analysis Software calculates and reports these quality control parameters.

If the above criteria are not met the run is invalid and must be repeated.

- **The mean OD value for the Zero Standard (Green Diluent) should be ≤ 0.150 . If the mean OD value is > 0.150 the plate washing procedure should be investigated.**

Interpretation of Results

QFT results are interpreted using the following criteria:

NOTE: Diagnosing or excluding tuberculosis disease, and assessing the probability of LTBI, requires a combination of epidemiological, historical, medical, and diagnostic findings that should be taken into account when interpreting QFT results.

Nil [IU/mL]	TB Antigen minus Nil [IU/mL]	Mitogen minus Nil [IU/mL] ¹	QFT Result	Report/Interpretation
≤ 8.0	< 0.35	≥ 0.5	Negative	<i>M. tuberculosis</i> infection NOT likely
	≥ 0.35 and < 25% of Nil value	≥ 0.5		
	≥ 0.35 and ≥ 25% of Nil value	Any	Positive²	<i>M. tuberculosis</i> infection likely
	< 0.35	< 0.5	Indeterminate³	Results are indeterminate for TB Antigen responsiveness
≥ 0.35 and < 25% of Nil value	< 0.5			
> 8.0 ⁴	Any	Any		

¹ Responses to the Mitogen positive control (and occasionally TB Antigen) can be commonly outside the range of the microplate reader. This has no impact on test results.

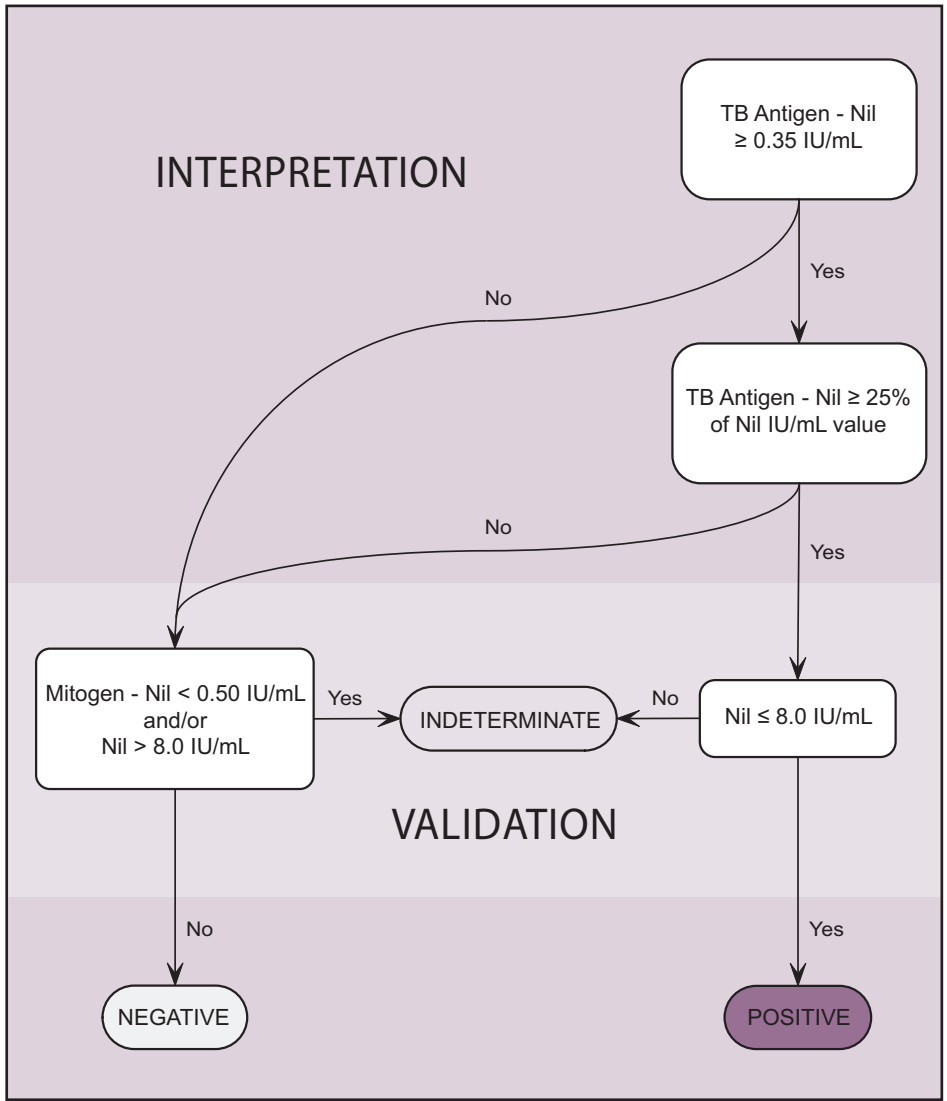
² Where *M. tuberculosis* infection is not suspected, initially positive results can be confirmed by retesting the original plasma samples in duplicate in the QFT ELISA. If repeat testing of one or both replicates is positive, the individual should be considered test positive.

³ Refer to Trouble Shooting section for possible causes.

⁴ In clinical studies, less than 0.25% of subjects had IFN-γ levels of > 8.0 IU/mL for the Nil Control.

The magnitude of the measured IFN-γ level cannot be correlated to stage or degree of infection, level of immune responsiveness, or likelihood for progression to active disease.

FIGURE 3. Interpretation Flow Diagram



8. LIMITATIONS

Results from QFT testing must be used in conjunction with each individual's epidemiological history, current medical status, and other diagnostic evaluations.

Individuals with Nil values greater than 8 IU/mL are classed as "indeterminate" because a 25% higher response to the TB Antigens may be outside the assay measurement range.

Unreliable or indeterminate results may occur due to:

- Deviations from the procedure described in the Package Insert,
- Excessive levels of circulating IFN- γ or presence of heterophile antibodies,
- Longer than 16 hours from blood specimen drawing to incubation at 37°C.

9. PERFORMANCE CHARACTERISTICS

Clinical Studies

As there is no definitive standard for latent tuberculosis infection (LTBI), an estimate of sensitivity and specificity for QFT cannot be practically evaluated. Specificity of QFT was approximated by evaluating false positive rates in the persons with low risk (no known risk factors) of tuberculosis infection. Sensitivity was approximated by evaluating groups of patients with culture-confirmed active TB disease.

Specificity

In a U.S. study involving 866 volunteers, blood was drawn for QFT when a TST was placed. Demographic information and risk factors for TB were determined using a standard survey at the time of testing. Of 432 volunteers with no known risk factors for *M. tuberculosis* infection, QFT and TST results were available for 391. None were BCG vaccinated. A second specificity study was performed with QFT in low risk individuals in Japan, approximately 90% who had received BCG vaccination. Results from both the specificity studies are shown in Table 2.

TABLE 2. QFT specificity: Results for persons with no reported risk for *M. tuberculosis* infection.

STUDY	BCG Status % Vaccinated	Total tested	No. QFT Indeterminate	No. QFT Positive / No. Valid Tests	QFT Specificity (95% CI)	No. TST Positive / No. Tested	TST* Specificity (95% CI)
USA (unpublished)	0%	391	1	3 / 390	99.2% (97.6-99.8)	6 / 391	98.5% (96.5-99.4)
Japan (unpublished)	~90%	190	4	3 / 186	98.4% (95-99.6)	–	–
TOTAL		581	5 / 584 (0.9%)	6 / 576	99.0%	–	–

*Using a 10mm TST cut off. TST specificity estimate is 99.1% if using a 15mm cut off.

Sensitivity for active TB

TB-suspects from Australia and Japan who were subsequently confirmed to have *M. tuberculosis* infection by culture were tested to evaluate sensitivity of QFT. While there is no definitive standard test for latent tuberculosis infection (LTBI), a suitable surrogate is microbiological culture of *M. tuberculosis* since patients with disease are by definition infected. The patients had received less than 8 days of treatment prior to collecting blood for QFT testing.

Table 3 summarizes findings from the two groups of *M. tuberculosis* culture positive patients. Overall sensitivity of QFT for active TB disease was 89% (48/54).

TABLE 3. QFT: Subjects with culture-confirmed *M. tuberculosis* infection.

STUDY	Disease Confirmed By	No. QFT Positive / No. Valid Tests	QFT Sensitivity (95% CI)
Japan TB Patients Validation study	Culture	24 / 27	89% (72-96%)
Australian TB Patients Validation Study	Pulmonary	7 / 10	70% (40-89%)
	Extra-Pulmonary	17 / 17	100% (82-100%)
TOTAL		48 / 54	89% (78-95%)

Diagnosis of LTBI

A number of studies have been published which demonstrate the performance of QFT in various populations at risk of LTBI. The principle findings of some selected studies are shown in Table 4.

TABLE 4. Selected published studies on QFT in populations at risk of LTBI.

STUDY	Total tested	Outcomes and Findings
Indian HCW (Pai <i>et al</i> 2005) ²⁶	726	Setting of very high TB rates. 40% QFT-Gold IT positive cf 41% TST positive at 10mm. High concordance with TST, no effect of BCG on either test. Both tests related to risk factors of age and period of work in healthcare.
Danish HIV (Brock <i>et al</i> 2006) ⁵	590	Overall prevalence of LTBI by QFT-Gold IT was 4.6% (27/590) in HIV+ persons. Positive results were associated with TB risks. Two QFT-Gold IT positive subjects progressed to active TB within one year. Indeterminate responses (n=20, 3.4%) were significantly associated with a CD4 count <100 / μ L.
Hospitalized Children (Dogra <i>et al</i> 2006) ¹⁰	105	Children in whom TB was suspected or had a history of TB contact were tested with QFT-Gold IT and TST. 10.5% QFT-Gold IT positive cf 9.5% TST positive at 10mm. Agreement between tests was 95.2% overall and 100% in non-BCG vaccinated.
German Contacts (Diel <i>et al</i> 2006) ⁹	309	Close contacts of 15 different index cases were tested. 51% were BCG vaccinated, 27% foreign born. 70% of BCG vaccinates and 18% of non-vaccinated were TST positive (5mm), whereas 9% and 11% were QFT-Gold IT positive, respectively. QFT-Gold IT was associated with TB risk. TST was only associated with BCG vaccination.

Many more publications describe the performance of the less-sensitive liquid antigen version of QuantiFERON®-TB Gold (the precursor to QFT) and the QFT test. These studies include use of the test(s) in contacts of active TB cases^{9,11,19,25}, children^{6-10,25,28}, HIV positive^{2,5,20}, healthcare workers^{13,26,32}, immune suppressed^{3,4,22,23,27,30,31}, as well as TB suspects^{7,8,10,18} and low risk individuals¹⁵.

Repeatability and effect of TST on subsequent QFT testing

As part of the U.S. specificity study, a subset of the volunteers were retested between 4 and 5 weeks after the original QFT test and TST. QFT results for 260 recruits were available at both time points and the level of agreement was 99.6% (259/260). A prior TST did not induce positive QFT responses.

10. TECHNICAL INFORMATION

Indeterminate Results

Indeterminate results should be uncommon and may be related to the immune status of the individual being tested, but may also be related to a number of technical factors:

- Longer than 16 hours from blood draw to incubation at 37°C,
- Storage of blood outside the recommended temperature range (22°C ± 5°C),
- Insufficient mixing of blood collection tubes,
- Incomplete washing of the ELISA plate.

If technical issues are suspected with the collection or handling of blood samples, repeat the entire QFT test with a new blood specimen. Repeating the ELISA testing of stimulated plasmas can be performed if inadequate washing or other procedural deviation with the ELISA test is suspected. Indeterminate tests that result from low Mitogen or high Nil values would not be expected to change on repeat unless there was an error with the ELISA testing. Indeterminate results should be reported as such. Physicians may choose to redraw a specimen or perform other procedures as appropriate.

Clotted Plasma Samples

Should fibrin clots occur with long term storage of plasma samples, centrifuge samples to sediment clotted material and facilitate pipetting of plasma.

ELISA Trouble Shooting

Non-specific Colour Development

POSSIBLE CAUSE	SOLUTION
Incomplete washing of the plate.	Wash the plate at least 6 times with 400 μ L/well of wash buffer. More than 6 washing cycles may be required depending on the washer being used. A soak time of at least 5 seconds between cycles should be used.
Cross-contamination of ELISA wells.	Take care pipetting and mixing sample to minimise risk.
Kit / Components have expired.	Ensure kit is used within the expiry date. Ensure reconstituted Standard and Conjugate 100X Concentrate are used within three months of the reconstitution date.
Enzyme Substrate Solution is contaminated.	Discard substrate if blue colouration exists. Ensure clean reagent reservoirs are used.
Mixing of plasma in centrifuge tubes before harvesting	Ensure that plasma samples are carefully harvested from above gel without pipetting up and down and taking care not to disturb material on the surface of the gel.

Low Optical Density Readings for Standards

POSSIBLE CAUSE	SOLUTION
Standard dilution error.	Ensure dilutions of the Kit Standard are prepared correctly as per the Package Insert.
Pipetting error.	Ensure pipettes are calibrated and used according to manufacturer's instructions.
Incubation temperature too low.	Incubation of the ELISA should be performed at Room Temperature, 17°C to 27°C.
Incubation time too short.	Incubation of the plate with the conjugate, standards and samples should be for 120 ± 5 minutes. The Enzyme Substrate Solution is incubated on the plate for 30 minutes.
Incorrect plate reader filter used.	Plate should be read at 450nm with a reference filter of between 620 and 650nm.
Reagents are too cold.	All reagents, with the exception of the Conjugate 100X Concentrate, must be brought to room temperature prior to commencing the assay. This takes approximately one hour.
Kit / Components have expired.	Ensure kit is used within the expiry date. Ensure reconstituted Standard and Conjugate 100X Concentrate are used within three months of the reconstitution date.

High Background

POSSIBLE CAUSE	SOLUTION
Incomplete washing of the plate.	Wash the plate at least 6 times with 400µL/well of wash buffer. More than 6 washing cycles may be required depending on the washer being used. A soak time of at least 5 seconds between cycles should be used.
Incubation temperature too high.	Incubation of the ELISA should be performed at Room Temperature, 17°C to 27°C.
Kit / Components have expired.	Ensure kit is used within the expiry date. Ensure reconstituted Standard and Conjugate 100X Concentrate are used within three months of the reconstitution date.
Enzyme Substrate Solution is contaminated.	Discard substrate if blue colouration exists. Ensure clean reagent reservoirs are used.

Non-linear Standard Curve and Duplicate Variability

POSSIBLE CAUSE	SOLUTION
Incomplete washing of the plate.	Wash the plate at least 6 times with 400µL/well of wash buffer. More than 6 washing cycles may be required depending on the washer being used. A soak time of at least 5 seconds between cycles should be used.
Standard dilution error.	Ensure dilutions of the Standard are prepared correctly as per the Package Insert.
Poor mixing.	Mix reagents thoroughly by inversion or gentle vortexing prior to their addition to the plate.
Inconsistent pipetting technique or interruption during assay set-up.	Sample and standard addition should be performed in a continuous manner. All reagents should be prepared prior to commencing the assay.

Assay procedure video and solution to most technical problems can be found on the Product Information and Technical Guide CD-ROM available free of charge from Cellestis or via your distributor.

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12. TECHNICAL SERVICE

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13. ABBREVIATED TEST PROCEDURE

STAGE 1 – BLOOD INCUBATION

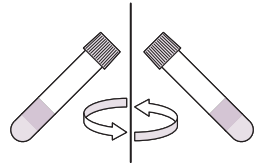
1. Collect patient blood into blood collection tubes and mix by shaking them ten (10) times just firmly enough to ensure that the entire inner surface of the tube has been coated with blood, to solubilize antigens on tube walls.



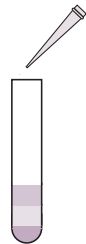
2. Incubate tubes **upright** at 37°C for 16 to 24 hours.



3. Following incubation, centrifuge tubes for 15 minutes at 2000 to 3000g RCF (g) to separate the plasma and the red cells.



4. After centrifugation, harvest plasma samples by pipette. Avoid pipetting up and down or mixing plasma by any means prior to harvesting. At all times, take care not to disturb material on the surface of the gel.



STAGE 2 – IFN- γ ELISA

1. Equilibrate ELISA components, with the exception of the Conjugate 100X Concentrate, to room temperature for at least 60 minutes.

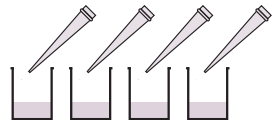


2. Reconstitute the Kit Standard to 8.0 IU/mL with distilled or deionised water. Prepare four (4) standard dilutions.

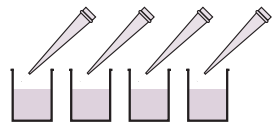


3. Reconstitute freeze-dried Conjugate 100X Concentrate with distilled or deionised water.

4. Prepare working strength conjugate in Green Diluent and add 50 μ L to all wells.



5. Add 50 μ L of test plasma samples and 50 μ L of standards to appropriate wells. Mix using shaker.

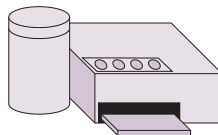


6. Incubate for 120 minutes at room temperature.

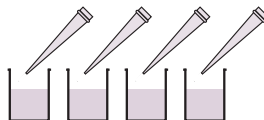


STAGE 2 – IFN- γ ELISA (Continued)

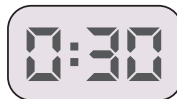
7. Wash wells at least 6 times with 400 μ L/well of wash buffer.



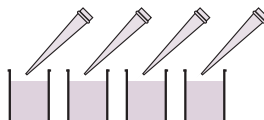
8. Add 100 μ L Enzyme Substrate Solution to wells. Mix using shaker.



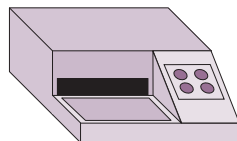
9. Incubate for 30 minutes at room temperature.



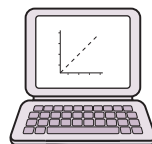
10. Add 50 μ L Stop Solution to all wells. Mix using shaker.



11. Read results at 450nm with a 620 to 650nm reference filter.



12. Analyse Results.



14. SIGNIFICANT CHANGES

Significant changes in Edition CA05990301D of the QFT Package Insert are summarised in the table below:

Section	Page	Change(s)
5. Specimen Collection and Handling	10	Amend tube shaking procedure
6. Directions for Use	11	Amendment to handling procedures for tubes containing blood
6. Directions for Use	14	Amendment to handling procedures for plasma samples
10. Technical Information	24	Addition: 'Mixing of plasma in centrifuge tubes before harvesting'
12. Technical Service	30	New email address for Technical Service



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