**Mycobacterium tuberculosis** End-Point PCR Kit

**Product Insert**

**Product# EP42100**

**Intended Use**

Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is designed for the detection of *M. tuberculosis* DNA based on the use of end-point PCR technology. This kit is designed for research use only and not for use in diagnostic procedures.

**Background Information**

*Mycobacterium tuberculosis* is a pathogenic bacterial species belonging to the genus Mycobacterium, and is the causative agent of tuberculosis. Tuberculosis (TB) is a multifaceted disease and challenging public health problem in both industrialized and developing countries. According to the WHO, 8.8 million active cases of TB are diagnosed each year and of these, almost 2 million die. Once thought to be under control or even close to extinction, TB infection levels are rising and the threat is compounded by new, virulent and drug-resistant strains. Although most cases occur in the developing world (22 countries accounting for 80% of all global cases), increasing population mobility combined with facility of transmission means that no country is immune from the resurgence of TB. TB control programs are currently facing a number of constraints. Worldwide, fewer than 25% of all tuberculosis cases are detected. Of utmost concern is the absence of a timely and accurate test for the diagnosis of mycobacterial disease. Early diagnosis is crucial for the prevention of further spread of the disease.

**Product Description**

Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is designed for the detection of *M. tuberculosis* specific DNA based on the use of end-point PCR technology. This kit is designed for research use only and not for use in diagnostic procedures. The kit includes Master Mix and primers for the specific amplification of a 307 nucleotide region of the *M. tuberculosis* genome, as well as a positive control and a negative control to confirm the integrity of the kit reagents. In addition, the kit contains loading dye and a DNA ladder to facilitate analysis of the results.

The detection of *Mycobacterium tuberculosis* specific DNA is based on end-point PCR technology, utilizing polymerase chain reaction (PCR) for the amplification of specific *M. tuberculosis* DNA sequences. For analysis of the PCR data, the PCR reaction is loaded on an agarose DNA gel along with the provided DNA ladder for qualitative analysis.

Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit was developed and validated to be used with the following PCR instruments:

- Qiagen Rotor-Gene Q
- BioRad CFX96 Touch™ Real-Time PCR Detection System

**Kit Components**

<table>
<thead>
<tr>
<th>Component</th>
<th>Product # EP42100 (24 preps)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDx 2X PCR Master Mix</td>
<td>350 µL</td>
</tr>
<tr>
<td>M. tuberculosis Primer Mix</td>
<td>70 µL</td>
</tr>
<tr>
<td>M. tuberculosis Positive Control</td>
<td>50 µL</td>
</tr>
<tr>
<td>Nuclease-Free Water</td>
<td>1.25 mL</td>
</tr>
<tr>
<td>Loading Dye</td>
<td>100 µL</td>
</tr>
<tr>
<td>DNA Ladder</td>
<td>100 µL</td>
</tr>
<tr>
<td>Product Insert</td>
<td>1</td>
</tr>
</tbody>
</table>
Storage Conditions and Product Stability
- All kit components should be stored at -20°C upon arrival.
- Repeated thawing and freezing (> 2 x) of the Master Mix and Positive Control should be avoided, as this may affect the performance of the assay. If the reagents are to be used only intermittently, they should be frozen in aliquots.
- All reagents can be stored for 1 year at -20°C without showing any reduction in performance.

Customer-Supplied Reagents and Equipment
- Appropriate End-point PCR Instrument
- DNA Purification Kit
  - The kit is compatible with all DNA purification kits that yield high quality, inhibitor-free DNA
  - Recommended Purification Kit: Norgen’s Sputum DNA Isolation Kit (Cat. 46200) or Urine DNA Isolation Kit (Slurry Format) (Cat. 48800)
- Disposable powder-free gloves
- Benchtop microcentrifuge
- Micropipettors
- Sterile pipette tips with filters
- PCR tubes
- Vortex mixer
- Agarose gel electrophoresis apparatus
- UV transilluminator with suitable gel documentation system
- PCR reaction preparation station (Optional)

Quality Control
In accordance with Norgen’s ISO 9001 and ISO 13485-certified Quality Management System, each lot of Norgen’s Mycobacterium tuberculosis End-Point PCR Kit is tested against predetermined specifications to ensure consistent product quality.

Warnings and Precautions
- Norgen’s Mycobacterium tuberculosis End-Point PCR Kit is intended for research purposes only. It is not intended for diagnostic use.
- Follow universal precautions. All specimens should be considered as potentially infectious and handled accordingly.
- Ensure that a suitable lab coat, disposable gloves and protective goggles are worn when handling specimens and kit reagents.
- Use sterile pipette tips with filters. Use proper pipetting techniques and maintain the same pipetting pattern throughout the procedure to ensure optimal and reproducible values.
- As contamination of specimens or reagents can produce erroneous results, it is essential to use aseptic techniques. Pipette and handle reagents carefully to avoid mixing of the samples.
- Do not use supplies and equipment across the dedicated areas of i) specimen extraction, ii) reaction set-up and iii) amplification/detection. No cross-movement should be allowed between the different areas. Personal protective equipment, such as laboratory coats and disposable gloves, should be area specific.
- Store and extract positive material (specimens, controls and amplicons) separately from all other reagents and add it to the reaction mix in a spatially separated facility.
Dispose of unused kit reagents and specimens according to local, provincial or federal regulations.

Do not substitute or mix reagents from different kit lots or from other manufacturers. Do not use components of the kit that have been stored for more than 1 year.

The presence of PCR inhibitors may cause false negative or invalid results.

Potential mutations within the target regions of the M. tuberculosis genome covered by the primers in this kit may result in failure to detect the presence of the pathogen.

Good laboratory practice is essential for the proper performance of this kit. Ensure that the purity of the kit and reactions is maintained at all times, and closely monitor all reagents for contamination. Do not use any reagents that appear to be contaminated.

Ensure that appropriate specimen collection, transport, storage and processing techniques are followed for optimal performance of this test.

Instructions for Use

A. Sample Preparation

Purified DNA is the starting material for Norgen’s Mycobacterium tuberculosis End-Point PCR Kit. The quality of the DNA template will have a major impact on the performance of the M. tuberculosis detection test. The user must ensure that the method used for DNA purification is compatible with end-point PCR. We recommend the use of Norgen’s Sputum DNA Isolation Kit (Cat. 46200) or Urine DNA Isolation Kit (Slurry Format) (Cat. 48800). Norgen’s Sputum DNA Isolation Kit or Urine DNA Isolation Kit (Slurry Format) has been fully validated with Norgen’s Mycobacterium tuberculosis End-Point PCR Kit.

If using a different spin column based sample preparation procedure that includes ethanol-based wash buffers is used, a column drying step consisting of centrifugation for 3 minutes at 20,000 x g (~14,000 RPM), using a new collection tube, is highly recommended prior to the elution of the DNA. This will help to prevent the carry-over of any ethanol into the purified DNA, as ethanol is known to be a strong inhibitor of PCR. Ensure that any traces of ethanol from the sample preparation steps are eliminated prior to the elution of the DNA.

B. PCR Assay Preparation

Notes:
• Before use, suitable amounts of all PCR components should be completely thawed at room temperature, mixed by gentle vortexing or by pipetting, and centrifuged briefly.
• Work quickly on ice.
• The amount of MDx 2X PCR Master Mix provided is enough for up to 34 PCR reactions (24 sample PCR, 4 positive control PCR and 4 no template control PCR).
• For every PCR run, one reaction containing M. tuberculosis Positive Control and one reaction as no template control must be included for proper interpretation of results.
• The recommended minimum number of DNA samples tested per PCR run is 6.
• To avoid any contamination while preparing the PCR assay, follow the order outlined in Tables 1, 2 and 3 below to prepare the Negative Control, Detection Assay and Positive Control:
  1. Prepare the PCR Negative Control (Table 1)
  2. Prepare the PCR M. tuberculosis Assay (Table 2)
  3. Prepare the PCR Positive Control (Table 3)
To further avoid contamination, add the components to the PCR tubes in the order shown in the tables below (ie: 1) Nuclease-free water; 2) Master Mix; 3) Primer Mix; and 4) the Sample DNA or Positive Control).

1. For each PCR set, prepare one no template control PCR as shown in Table 1 below:

<table>
<thead>
<tr>
<th>PCR Components</th>
<th>Volume Per PCR Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nuclease-Free Water</td>
<td>8 µL</td>
</tr>
<tr>
<td>MDx 2X PCR Master Mix</td>
<td>10 µL</td>
</tr>
<tr>
<td>M. tuberculosis Primer Mix</td>
<td>2 µL</td>
</tr>
<tr>
<td>Total Volume</td>
<td>20 µL</td>
</tr>
</tbody>
</table>

2. Prepare the PCR reaction for sample detection as shown in Table 2 below. The recommended amount of sample DNA to be used is 2.5 µL. However, a volume between 1 and 5 µL of sample DNA may be used as template. Adjust the final volume of the PCR reaction to 20 µL using the Nuclease-Free Water provided.

<table>
<thead>
<tr>
<th>PCR Components</th>
<th>Volume Per PCR Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nuclease-Free Water</td>
<td>5.5 µL</td>
</tr>
<tr>
<td>MDx 2X PCR Master Mix</td>
<td>10 µL</td>
</tr>
<tr>
<td>M. tuberculosis Primer Mix</td>
<td>2 µL</td>
</tr>
<tr>
<td>Sample DNA</td>
<td>2.5 µL</td>
</tr>
<tr>
<td>Total Volume</td>
<td>20 µL</td>
</tr>
</tbody>
</table>

3. For each PCR set, prepare one positive control PCR as shown in Table 3 below:

<table>
<thead>
<tr>
<th>PCR Components</th>
<th>Volume Per PCR Reaction</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDx 2X PCR Master Mix</td>
<td>10 µL</td>
</tr>
<tr>
<td>M. tuberculosis Primer Mix</td>
<td>2 µL</td>
</tr>
<tr>
<td>M. tuberculosis Positive Control (PosC)</td>
<td>8 µL</td>
</tr>
<tr>
<td>Total Volume</td>
<td>20 µL</td>
</tr>
</tbody>
</table>
C. *M. tuberculosis* PCR Assay Programming

1. Program the thermocycler according to the program shown in Table 4 below.
2. Run one step PCR.

<table>
<thead>
<tr>
<th>PCR Cycle</th>
<th>Step</th>
<th>Temperature</th>
<th>Duration</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cycle 1</td>
<td>Step 1</td>
<td>95°C</td>
<td>3 min</td>
</tr>
<tr>
<td>Cycle 2 (40x)</td>
<td>Step 1</td>
<td>94°C</td>
<td>15 sec</td>
</tr>
<tr>
<td></td>
<td>Step 2</td>
<td>60°C</td>
<td>30 sec</td>
</tr>
<tr>
<td></td>
<td>Step 3</td>
<td>72°C</td>
<td>45 sec</td>
</tr>
<tr>
<td>Cycle 3</td>
<td>Step 1</td>
<td>72°C</td>
<td>5 min</td>
</tr>
<tr>
<td>Cycle 4</td>
<td>Step 1</td>
<td>4°C</td>
<td>∞</td>
</tr>
</tbody>
</table>

D. *M. tuberculosis* PCR Assay Interpretation

- For the analysis of the PCR data, the entire 20 µL PCR reaction should be loaded on a 1X TAE 1.4 % Agarose DNA gel along with 10 µL of Norgen’s DNA Ladder (provided).
- The PCR products should be resolved on the 1X TAE, 1.4 % Agarose gel at 150V for 30 minutes (Gel running time will vary depending on an electrophoresis apparatus).

<table>
<thead>
<tr>
<th>Input Type</th>
<th>Target amplification</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><em>M. tuberculosis</em> Target Band (307 bp)</td>
<td>PCR control Band (150 bp)</td>
</tr>
<tr>
<td>Positive Control</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>Positive Control</td>
<td>Yes</td>
<td>No</td>
</tr>
<tr>
<td>Negative Control</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Sample</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>Sample</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>Sample</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>Sample</td>
<td>Yes</td>
<td>No</td>
</tr>
</tbody>
</table>

For results obtained that are not covered in Table 5 above, please refer to the Frequently Asked Questions.
**E. *M. tuberculosis* PCR Assay Specificity**

The specificity of Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is first and foremost ensured by the selection of the *M. tuberculosis* -specific primers, as well as the selection of stringent reaction conditions. The *M. tuberculosis* primers were checked for possible homologies to all microorganisms in GenBank published sequences by sequence comparison analysis.

**Frequently Asked Questions**

1. **How many samples should be included per PCR run?**
   - Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is designed to test 24 samples. For every 6 samples, a non-template control (Nuclease Free Water) and a Positive Control must be included. It is preferable to pool and test 6 samples at a time.

2. **How should it be interpreted if in negative control the *M. tuberculosis* PCR control and the *M. tuberculosis* target showed amplification in a sample?**
   - The assay has to be repeated. It could happen when there is carryover contamination.

3. **How should it be interpreted if only the *M. tuberculosis* target was amplified in a sample?**
   - The sample tested should be considered as *M. tuberculosis* positive. At high *M. tuberculosis* input, the *M. tuberculosis* amplicon will be predominant and thus the *M. tuberculosis* PCR control may not amplify as they compete for PCR resources.
Related Products | Product #
---|---
*Mycobacterium tuberculosis* Primers and Control Set | EP42110
*Mycobacterium tuberculosis* SYBR Green RT-PCR Kit | SG42100
*Mycobacterium tuberculosis* TaqMan RT-PCR Kit | TM42100
*Mycobacterium tuberculosis* TaqMan Probe/Primer and Control Set - 100 Reactions | TM42110
Sputum DNA Isolation Kit | 46200
Urine DNA Isolation Kit (Slurry Format) | 48800

**Technical Support**
Contact our Technical Support Team between the hours of 8:30 and 5:30 (Eastern Standard Time) at (905) 227-8848 or Toll Free at 1-866-667-4362.

Technical support can also be obtained from our website (www.norgenbiotek.com) or through email at techsupport@norgenbiotek.com.

**Product Use Restriction**
Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is designed for the detection of *M. tuberculosis* specific DNA based on the use of end-point PCR technology. This kit is designed for research use only and not for use in diagnostic procedures.

Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit is intended for use by professional users such as technicians and biologists experienced and trained in molecular biological techniques including PCR.

Good laboratory practice is essential for the proper performance of this kit. Ensure that the purity of the kit and reactions is maintained at all times, and closely monitor all reagents for contamination. Do not use any reagents that appear to be contaminated.

Ensure that appropriate specimen collection, transport, storage and processing techniques are followed for optimal performance of this test.

The presence of PCR inhibitors may cause false negative or invalid results.

Potential mutations within the target regions of the *M. tuberculosis* genome covered by the primers in this kit may result in failure to detect the presence of the pathogen.

The respective user is liable for any and all damages resulting from application of Norgen’s *Mycobacterium tuberculosis* End-Point PCR Kit for use deviating from the intended use as specified in the user manual.

All products sold by Norgen Biotek are subjected to extensive quality control procedures and are warranted to perform as described when used correctly. Any problems should be reported immediately. The kit contents are for laboratory use only, and they must be stored in the laboratory and must not be used for purposes other than intended. The kit contents are unfit for consumption.

Norgen Biotek Corp.
3430 Schmon Parkway, Thorold, ON Canada L2V 4Y6
Phone: (905) 227-8848
Fax: (905) 227-1061
Toll Free in North America: 1-866-667-4362

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