Original Research Article

Truenat - A Novel Diagnostic Tool for Rapid Detection of Mycobacterium tuberculosis and Rifampicin Resistance in Pulmonary Samples

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ABSTRACT

Tuberculosis is a major public health problem and second largest cause of death among infectious diseases. Rapid diagnosis of tuberculosis and detection of drug resistance is essential for effective disease control. Control of TB is hindered by the lack of a simple and effective diagnostic tool that can be utilised in resource limited settings. Molecular techniques using Real time PCR technology have gained momentum for rapid diagnosis and detection of drug resistance. To study the prevalence of Pulmonary Tuberculosis and Rifampicin resistance in Kurnool district by using Truenat chip based micro PCR system and to understand the demographical distribution of cases attending the tertiary care hospital. This was a prospective study done during the period from March 2019 to Aug 2019. All the presumptive TB cases with chest symptoms attending Govt.General Hospital, Kurnool were included in the study. Sputum samples were processed by Truenat, a chip based nucleic acid amplification test (Molbio diagnostics pvt. ltd., Goa). The procedure included two steps: first - sputum samples were processed for separation of DNA by using True prep device. Second-Presence of MTB was detected by Truelab™ Real Time micro PCR Analyzer. All the MTB positive samples were tested for the presence of Rifampicin resistance by using another specialized chip. A total of 1825 sputum samples were processed, out of which 368 (20.16%) were positive for MTB. Rifampicin resistance was detected in 70 (19.02%). Positivity was more in males 63.23% as compared to females 36.76%. Majority of the cases were in the age group 40-60 years (46.58%), followed by 20-40 years (33.04%). Truenat is a simple method and laboratory technician with minimal training can perform the procedure. The Turn Around Time (TAT) is approximately one hour. As no special infrastructure or Airconditioning system is required to set up the device it can be used in resource limited settings for rapid diagnosis of MTB and detection of Rifampicin resistance.

KEYWORDS
Chip based assay, Trueprep AUTO Universal Cartridge Based Sample Prep Device Truenat™ MTB chip

Introduction

Tuberculosis causes the highest number of deaths globally despite the availability of potent anti-TB drugs. The global burden of TB remains enormous. As per Global TB report 2017, more than 9 million new tuberculosis cases and 1.6 million deaths occur annually worldwide. Estimated MDR/RR cases are 6 lakhs. India is the highest TB burden country. 1/4th of the global
annual new cases occur in India. Incidence is 2.7 million cases annually and mortality is 4.35 lakhs per year (1). Over 25% of patients seeking care in India’s public sector are neither diagnosed nor started on treatment (2). Tuberculosis (TB) is the second largest killer worldwide, after HIV and is the leading cause of death in HIV patients. Pulmonary TB spreads through aerosols and is highly contagious. Over 80% of TB infections are pulmonary and if left untreated, a pulmonary TB patient can infect up to 10-15 other people through close contact over the course of a year(3). Due to the highly infectious nature of pulmonary TB, it is important to diagnose and treat the disease very early. Despite the availability of highly effective treatment for decades, TB remains a major global health problem mainly because of poor case detection(4). The most common method for diagnosing pulmonary TB worldwide is sputum smear microscopy. However sensitivity of direct smear microscopy is low and estimates range from 30% to 70%. It is even lower in case of HIV-infected patients. Culture is more sensitive than microscopy and is considered the current gold standard. Culture requires specialized and controlled laboratory facility and highly skilled manpower and takes 2 to 6 weeks to provide the result. Automated methods and Molecular techniques such as polymerase chain reaction (PCR) or Real Time PCR are much more sensitive than microscopy and culture. However these tests have so far been restricted to centralized reference laboratories as they require skilled manpower and elaborate infrastructure. Also the turnaround time for results could take a few day (5,6).

Moreover Drug resistance is a major issue in the treatment of tuberculosis. Drug resistance is because of either mismanagement of TB patients- wrong diagnosis, delay in diagnosis, wrong or interrupted treatment and injudicious use of both first and second line drugs. Multiple approaches to improve diagnosis of TB are in development. Amongst these are CBNAAT (GeneXpert) and LPA, endorsed by WHO to be used in RNTCP for rapid diagnosis of MTB and detection of Rifampicin resistance (7). These require uninterrupted power supply, air conditioning system, proper infrastructure and trained personnel. Patients have to travel to nearest testing centre or samples need to be transported. This limits its use in peripheral settings and Active Case Finding program (8).

The need is for accurate, feasible, rapid, affordable, and if possible, near-point-of-case TB diagnostic tests for use in resource limited settings (9,10). The Truelab™ Real Time micro PCR System enables decentralization and near patient diagnosis of MTB by making real time PCR technology rapid, simple, robust and user friendly and offering “sample to result” capability even at resource limited settings. It is supplied by Molbio diagnostics pvt. ltd., Goa, funded by Bigtec labs, India. This test has been evaluated by the premiere institutes for its sensitivity and specificity and has been recommended in health care settings (9,11). It is extensively validated and licensed by Indian FDA. It has sensitivity of 99% and specificity of 100%. The MTB strain H37Rv from Zeptometrix was used for LOD (Limit of Detection) determination. LOD was determined to be 100 CFU/ml in sputum sample.

To know the prevalence of Tuberculosis in Kurnool district by using chip based Truelab™ Real Time micro PCR System
To understand the demographics of TB
To know the prevalence of Rifampicin resistance among New and Previously treated cases.

Materials and Methods

Sputum samples of all the Presumptive pulmonary TB cases were collected in universal container. Having supplied by the
Govt to our institute in March 2019, the Truenat device was used for this study.

Each sample was processed by Trueprep Auto MTB sample pretreatment pack which employs a combination of reagents. These reagents homogenise the sample to release the bacilli, concentrate to get better yields of bacilli and also discard potentially inhibitory substances. The pretreated sample is loaded in Trueprep AUTO Universal Cartridge Based Sample Prep kit (Figure 1) and Device (Figure 2) to enable further extraction and purification of the bacterial DNA so that it is free from PCR inhibitors. It is a light weight, portable device designed to operate at room temperature. It operates on mains and/or rechargeable battery and is fully automatic with minimal hands on time. The cartridge based extraction process is quick, reliable and efficient and does not require highly skilled personnel to carry out the extraction process. All the waste from processing of the sample is contained within the cartridge dump area thus posing no risk from potentially biohazardous material. At the end of processing the bound DNA is eluted and collected in elution chamber. The entire process takes 20 min. The elute is transferred to the Elute Collection tube (ECT).

The Truenat™ MTB chip(Figure 4) is placed on the chip tray of the Truelab™ Real Time micro PCR Analyzer (Figure 3). Six μL of the purified DNA is then dispensed using the provided micropipette and tip into the microtube containing freeze dried PCR reagents and allowed to stand for 30-60 seconds to get a clear solution. Six μL of this clear solution is then pipetted out and dispensed into the reaction well of the Truenat™ MTB chip and the test is started. A positive amplification causes the dual labelled fluorescent probe to release the fluorophore in an exponential manner which is then captured by the built-in opto-electronic sensor and displayed as amplification curve on the analyzer screen, on a real time basis during the test run. The Cycle threshold (Ct) is defined as the number of amplification cycles required for the fluorescent signal to cross the threshold. In the case of negative samples, amplification does not occur and a horizontal amplification curve is displayed on the screen during the test run.

At the end of the test run, a MTB “DETECTED” or “NOT DETECTED” result is displayed and in positive cases, quantitative value is also displayed on the screen. Based on the Ct of the internal positive control (IPC), the validity of the test run is also displayed. The IPC is a full process control that undergoes all the processes the specimen undergoes - from extraction to amplification thereby validating the test run from sample to result.

The target sequence for this kit is part of the ribonucleoside-diphosphate reductase gene, the product of which provides the precursor for DNA synthesis. The region selected is specific to the MTB complex. The used Truenat™ MTB micro PCR chip in submerged in freshly prepared 0.5% sodium hypochlorite solution for 30 minutes before disposal as per the standard medical waste disposal guidelines.

The samples which were positive for MTB were loaded into another chip designed for detection of rifampicin resistance (Truenat™ MTB/RIF micro PCR chip) and loaded into Truelab™ Real Time micro PCR Analyzer.

**Results and Discussion**

A total of 1825 sputum samples were processed during the period from March 2019 to Aug 2019. Out of these 368 (20.16%) were positive for MTB and 1457 (79.83%) were negative for MTB.

Out of 368 MTB positive cases, Rifampicin
resistance was detected in 70 (19.02%) and 298 (80.97%) cases were Rifampicin sensitive. Most of the samples were from age group 40-60 (46.58%) followed by 20-40 (33.04%). Out of the total 1825 cases, Males were 1154 (63.23%); amongst them, MTB was detected in 274 (23.74%) and Rifampicin resistance seen in 53 (19.34%). Females contributed to 671 (36.76%); amongst them MTB was detected in 94 (14.01%) and Rifampicin resistance seen in 17 (18.08%).

Out of the total 1825 cases, New cases were 1260; amongst these MTB was detected in 206 (16.34%) and Rifampicin resistance in 28 (13.59%). Previously treated cases were 565; amongst these MTB was detected in 162 (28.67%) and Rifampicin resistance in 42 (25.92%). A major WHO priority for TB diagnostics is to implement a rapid, sputum-based molecular test to replace smear microscopy at the peripheral level (i.e., designated microscopy centres and attached primary healthcare facilities) (9,11,12). RNTCP has recently developed National Strategic Plan (NSP) to work towards achieving the goal of eliminating TB by 2025(12). Drug resistant TB (DRTB) is a major impediment to achieve this goal (13). As per the Drug resistance survey conducted in India in 2014-16, the prevalence of MDR-TB was 2.84% and 11.6% among new and previously treated TB cases respectively (2).

Best method to control TB is early diagnosis and initiation of appropriate treatment. Truenat Real time quantitative micro PCR assay serves the need of the day as it is a rapid, portable and can be easily used in resource limited settings. It reduces the logistical hurdles involved with transportation of sputum samples to distant referral laboratories (12). The Turn around time is about one hour which allows same day diagnosis, same day treatment decision and initiation (13). A limitation of Truenat is, it cannot detect MDRTB and resistance to second line injectables which is of most significance in high burden countries like India. But as compared to any other molecular method, it is very effective(11). In the present study, out of the total 1825 sputum samples, Mycobacterium tuberculosis was detected in 368 samples (20.16%). Rifampicin resistance was seen in 70 (19.02%). People in the age group of 40-60 were in maximum numbers (46.58%). Males were predominantly affected (63.23%). Rifampicin resistance was more in males (19.34%) and among previously treated cases 25.92%. Similar results were observed in many studies (14) (Table 1)

<table>
<thead>
<tr>
<th>Age in years</th>
<th>No. of samples</th>
</tr>
</thead>
<tbody>
<tr>
<td>0-20</td>
<td>116 (06.35%)</td>
</tr>
<tr>
<td>20-40</td>
<td>603 (33.04%)</td>
</tr>
<tr>
<td>40-60</td>
<td>850 (46.58%)</td>
</tr>
<tr>
<td>&gt;60</td>
<td>256 (14.02%)</td>
</tr>
<tr>
<td>Total</td>
<td>1825</td>
</tr>
</tbody>
</table>

In order for wide availability as well as affordability even in peripheral settings, there is a need for make in India diagnostics to cover millions of people under risk of tuberculosis (Fig. 5 and 6).

Table.1 Age wise distribution of samples
**Fig. 1** Trueprep AUTO Universal Cartridge Based Sample Prep kit

![Fig. 1](image1)

**Fig. 2** Trueprep AUTO Universal Cartridge Based Sample Prep device

![Fig. 2](image2)

**Fig. 3** Truelab™ Real Time micro PCR Analyzer

![Fig. 3](image3)

**Fig. 4** Truenat™ MTB chip

![Fig. 4](image4)
Fig. 5

MTB positive
MTB negative

Fig. 6

Chart 1 Results obtained gender wise

Males: 1154 (63.23%)
  MTB detected: 274 (23.74%)
  Rif resistant: 53 (19.34%)

Females: 671 (36.76%)
  MTB detected: 94 (14.01%)
  Rif resistant: 17 (18.08%)
The performance of Truenat has been evaluated extensively by various researchers and has been compared with conventional culture based as well as with other molecular diagnostic methods. Nikam et al., from Hinduja Hospital and Medical research centre, Mumbai evaluated the performance of Truenat RTPCR in comparison with GeneXpert on sputum samples from Pulmonary TB cases and found a high concordance(96%) with GeneXpert(9).

In another study by Nikam et al., Truenat MTB test was found to have sensitivity of 91.1% and in- house nested PCR a sensitivity of 90.5%(11). Though culture based methods are considered gold standard in the diagnosis of tuberculosis, molecular diagnostic methods have been used extensively in the diagnosis of tuberculosis because of the rapidity of the detection of cases which helps in early diagnosis and initiation of treatment. GeneXpert developed by Cepheid company, has long been considered one of the best molecular method because of the ease of doing as it is a cartridge based method which involves simple procedure. But the cost of the instrument and cost per test has been a major drawback for its extensive utilisation in peripheral setups.

Truenat PCR system is a cartridge based test for the rapid diagnosis of tuberculosis along with the detection of rifampicin resistance which is developed in India. With the extensive availability of this economically viable diagnostic method in resource poor country like India will not only help in the early diagnosis but also in the appropriate management and infection control measures and controls the spread of this deadly disease. Truenat will improve linkage-to-care and increase life expectancy also(2,12).

Truenat MTB test is extremely useful test in resource limited health care settings such as designated microscopy centers (DMCs) and primary healthcare facilities in India. It is a rapid test, allows detection of TB in approximately one hour and can be utilized in near-care settings to provide quick and accurate diagnosis. As a portable platform it could also be utilized in ACF (Active case finding) programs, which are currently the need of the hour.

References


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