Original Research Article

Comparative Evaluation of GeneXpert MTB/RIF Assay and Microscopy for Rapid Diagnosis of Tuberculous Meningitis in Children

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ABSTRACT

Tuberculous meningitis (TBM) is the most devastating complication of Mycobacterium tuberculosis infection. Diagnosis of TBM is challenging in young children due to the paucibacillary nature of disease. Microscopy for acid-fast bacilli in CSF is fast but has very low sensitivity whereas culture may take up to 42 days. Due to the urgency of diagnosis in suspected TBM cases, a rapid, accurate diagnostic test could have a great impact on survival. Aims of the study are to prospectively determine the diagnostic accuracy of Xpert MTB/RIF in a large consecutive series of samples from patients presenting with suspected Tuberculous meningitis. In this prospective hospital-based study, 147 children presenting with suspected tuberculous meningitis from May 2017 to February 2018 were included. Cerebrospinal fluid samples were tested by Ziehl-Neelsen smear, mycobacterial culture on Lowenstein Jensen medium, and Xpert MTB/RIF assay. Kappa statistics (K) was used to determine agreement beyond what would be expected by chance between smear microscopy and Gene Xpert assay results. Out of a total of 147 CSF samples tested, 10 (6.8%) were positive for Mycobacterium tuberculosis by Xpert MTB/RIF assay. Out of these 10 samples, only one showed AFB on ZN smear. Culture was negative for all the 147 CSF samples. The MTB/RIF test has a short turnaround time and simultaneously detects M. tuberculosis and RIF resistance in less than 2h. It could be a useful tool for rapid identification of M. tuberculosis, especially in smear-negative clinical samples.

Keywords

Drug resistance; GeneXpert; Meningitis; Tuberculosis

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Introduction

Tuberculous meningitis (TBM) is the most devastating consequence of infection with Mycobacterium tuberculosis. Approximately a third of patients die soon after presenting to hospital, and many of those surviving are left with severe neurological sequelae. Rapid TBM diagnosis and treatment is a strong prognostic indicator for reduced death and neurologic deficit. However, many patients are diagnosed late because initial signs are non-specific, and rapid and sensitive diagnostic tests are lacking.

Available TBM diagnostic tests are inadequate, for the most part due to low
bacillary loads in CSF.\(^4\) Ziehl-Neelsen (ZN) microscopy staining of cerebrospinal fluid (CSF) is the most widely applied rapid diagnostic technique; however, sensitivity for TBM rarely exceeds 20%.\(^5\) Liquid culture techniques, including the mycobacterial growth indicator tube (MGIT; Bactec) and the mycobacterial observation drug susceptibility assay (MODS) culture offer improved sensitivity over solid culture, to a sensitivity of almost 60%.\(^6\) The clinical value of culture techniques is limited to diagnostic confirmation and drug susceptibility testing, because they take 1 to 4 weeks to return a positive result, and negative results cannot be used to exclude a TBM diagnosis.

The GeneXpert MTB/RIF test (Cepheid) is a closed-cartridge based system that is easy to operate by minimally trained staff and gives results in approximately 2 hours.\(^7\) The Xpert MTB/RIF test was approved by the WHO in 2010 for the diagnosis of pulmonary TB following extensive evaluation projects in six countries led by the Foundation for Innovative New Diagnostics (FIND).\(^8\)

The test is based on a real-time heminested PCR test which detects the presence of *M. tuberculosis* complex bacilli.\(^9\) By using 5 molecular beacons which span the rpoB gene 81-bp rifampin resistance-determining region (RRDR), the test simultaneously determines susceptibility to rifampin, which can be used as a surrogate marker for multidrug resistance (MDR).\(^9\) The closed-cartridge system makes it possible for the assay to be used outside the laboratory environment, and studies assessing biosafety have suggested that the use of Xpert MTB/RIF carries a smaller biohazard risk than smear microscopy.\(^7\) The risk of cross-contamination is also reduced with the closed cartridge system.\(^7\) The test has shown sensitivity above 90% for culture positive tuberculosis, with high specificity in sputum samples. Sensitivity in individuals with HIV coinfection is over 80%.\(^10-12\) Several studies have reported successful use of the Xpert MTB/RIF test on extrapulmonary samples, with overall sensitivities of over 80% and specificity reaching 100%.\(^13-17\)

Due to the urgency of diagnosis in suspected TBM cases, a rapid, accurate diagnostic test which also is able to identify rifampin resistance could have a great impact on survival. The aim of the present study was to prospectively determine the diagnostic accuracy of Xpert MTB/RIF as compared to smear microscopy in a large consecutive series of samples from patients presenting to the hospital with suspected tuberculous meningitis.

**Materials and Methods**

In this prospective hospital-based study, 147 children presenting with suspected tuberculous meningitis from May 2017 to February 2018 were included. A clinical diagnosis of meningitis was made based on clinical findings, CSF criteria or both.\(^18\)

**Clinical criteria** of meningitis included headache, fever and neck stiffness, with or without altered consciousness.

**CSF criteria** were cell count >10 cell/mm\(^3\), protein concentration >45 mg/dL, or the CSF: blood glucose ratio <0.5; either alone or in combination.

Signs and symptoms were recorded using standardized forms, and CSF was obtained by lumbar puncture for routine examination and microbiological testing.

All samples were subjected to Ziehl-Neelsen (ZN) staining, Xpert MTB/RIF (Cepheid, Sunnyvale, US) assay and culture inoculation. All samples were processed in level II biosafety cabinet.
Ziehl-Neelsen smear

Ziehl-Neelsen smears were prepared using standard methods. The layered smear was then stained according to standard procedures. The ZN smear was meticulously examined under a 1,000 magnification before being recorded as negative. Observation of a single acid-fast bacillus was considered a positive result.\(^{(19)}\)

Solid culture on LJ medium

Sterile, uncentrifuged CSF samples were directly inoculated without decontamination. A slope of LJ medium was inoculated with 0.25 to 0.5 mL of each specimen and incubated at 37°C for eight weeks. The LJ slants were inspected weekly. Growth on the LJ slants resembling mycobacterial colonies i.e., rough, tough and buff-coloured, was subjected to ZN staining and Gram staining to confirm the presence of acid-fast bacilli and rule out contamination respectively.

Xpert MTB/RIF

The Xpert MTB/RIF test was performed using the G4 version of cartridges as per the manufacturer’s instruction (Cepheid, Sunnyvale, CA). Unprocessed samples were used directly for performing the test and no frozen samples were used in the study. The sample was mixed with the sample reagent supplied with the test (2:1 ratio) and left to stand for 15 min, as per the manufacturer’s instructions, with intermittent manual shaking. The solution was then transferred to the Xpert cartridge using a provided pipette, and the cartridge was loaded onto the Xpert machine for analysis. Results were reported as positive or negative for \(M.\) \textit{tuberculosis}. Positive results were placed in one of four categories; very low, low, medium, or high. Rifampin resistance results were reported as susceptible or resistant.\(^{(20)}\)

Analysis

Statistical analysis was performed using SPSS version 22 (IBM Inc., Armonk, NY). Kappa statistics (K) was used to determine agreement beyond what would be expected by chance between smear microscopy result and Gene Xpertmtb/rif assay results. Four levels of agreement for kappa: <0.40 (poor), 0.40-0.59 (fair), 0.60-0.80 (good), and >0.80 (excellent) were reported.

Results and Discussion

A total of 147 consecutive patients with suspected meningitis were included. Patients were mostly male (60%), with a median age of 6 years (range 0.5-16 years), presenting after a median of 14 days (range 7–30 days) of symptoms, mostly with fever (100%), headache (67%) and altered consciousness (31.3%) as chief complaints. On examination, nuchal rigidity (77.2%), lowered consciousness (52.6%), and focal neurological signs were common.

Out of a total of 147 CSF samples tested, 10 (6.8%) were positive for \(Mycobacterium\) tuberculosis by GeneXpert MTB/RIF assay. None showed rifampicin resistance. Out of these 10 samples, only one showed AFB on ZN smear. Culture was negative for all the 147 CSF samples (Figure 1).

A comparison of smear microscopy and GeneXpert was done to gauge the concordance of microscopy results and GeneXpert MTB/RIF results, the findings revealed a significant (\(p=0.003\)) variation in agreement between the two assays 93.8% with a Cohen’s kappa, 0.172(CI= -0.122 - 0.465) (Table 1).

In this study, performance of the GeneXpert MTB/RIF assay with CSF specimens of patients who were suspected as having tubercular meningitis was evaluated.
Table 1 Kappa score on reliability of smear microscopy and GeneXpert assay

| Smear microscopy | GeneXpert MTB/RIF assay | Positive N (%) | Negative N (%) | Agreement | Kappa | 95% CI | p-value |
|------------------|-------------------------|----------------|----------------|-----------|-------|--------|---------|
| Positive         | 1                       | 0              | 93.8           | 0.172     | -0.122 - 0.465 | 0.003 |
| Negative         | 9                       | 137            |                |           |       |        |         |

N: Number of cases

Fig. 1 Venn diagram of overlap in TB meningitis diagnostics

The positive rate was 6.8% by GeneXpert MTB/RIF, and this rate was higher than that (0.68%) by a direct AFB smear examination. Our finding that the ZN smear is less sensitive than the GeneXpert MTB/RIF test is reasonable because the ZN smear method requires $5 \times 10^3$ to $1 \times 10^4$ bacilli/ml of specimen to generate a positive result. The out performance of Xpert MTB/RIF detected in current work is in agreement with other researchers who established the diagnosis in a significant proportion of patients and up to 10–15% of smear-negative TB. In this study, culture was negative for all the 147 CSF samples, may be due to less number of viable \textit{M. tuberculosis} in CSF samples.

The average turnaround time (including time taken to process specimens and testing time)
of the GeneXpert MTB/RIF assay and ZN smear method was significantly shorter (2.5 and 3.5 hours) than that of the solid culture on LJ medium (42 days). A shorter turnaround time can help TB patients be diagnosed early and treated in time. Therefore, the GeneXpert MTB/RIF assay has an obvious advantage in management of TB.

In our National TB Control Program (RNTCP), CBNAAT is the preferable investigation of choice for diagnosis of pediatric tuberculosis, especially in cases of extrapulmonary TB.\(^{(24)}\)

Although GeneXpert assay is considered a breakthrough in the diagnosis of TB and extra pulmonary TB (EPTB), one of the major limitations of this technique is that it cannot distinguish between viable and non-viable microorganisms while detecting mycobacterial DNA. A study by Boyles et al., on previously treated tuberculosis cases showed false positivity of CBNAAT due to dead bacilli.\(^{(25)}\)

The GeneXpert MTB/RIF assay is a simple, rapid, and accurate test method for detecting \textit{M.tuberculosis}, is less dependent on the operator’s skills, and staff with minimal training can use the equipment. Although the GeneXpert MTB/RIF assay has these advantages, similar to other tests for \textit{M.tuberculosis}, a negative result can not exclude the diagnosis of TB, and patients with positive results can also be assessed comprehensively with results of the ZN smear test, culture, clinical symptoms, and radiographic evidence.

In conclusion, the MTB/RIF test has a short turnaround time and simultaneously detects \textit{M. tuberculosis} and RIF resistance in less than 2 hours. It could be a useful tool for rapid identification of \textit{M. tuberculosis}, especially in smear-negative clinical samples. Further studies with culture or predefined CRS (certified reference standard) with follow up are required to define the exact sensitivity and specificity of GeneXpert.

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