Early Detection of Drug Susceptibility Test for *Mycobacterium Tuberculosis* by Slide DST Method Using Two Media- Middle Brook 7H9 Broth and Human Blood Media

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Abstract: This study was designed for early detection of drug resistance TB from direct sputum sample and also to suggest a simple method to detect multi drug resistance MTB. Samples were collected from 496 clinically suspected pulmonary tuberculosis cases. From that 207 samples were positive by smear microscopy. Culture and drug susceptibility were done by slide culture method using two media Middle brook 7H9 broth and Human blood media (HBM). From 207 samples 73 sample found to be MDR by Slide DST using Middle brook 7H9 broth and 74 sample using HBM. Sensitivity of Slide DST method was 97% with human blood media and 96% in case of Middle brook 7H9 broth. Time taken for the result was much less than conventional method. Susceptibility results were available much faster by slide DST method (10 days) compared to that by conventional DST (60 days). Cost of Slide DST method(20 Rs. for HBM & 45 Rs. for Middle brook 7H9 broth) for the detection was much lower than conventional DST. Slide DST method using these two medium could be applied as rapid diagnostic tool to detect drug resistance as it is very cheap and faster for diagnosis of drug resistance TB.

Keyword: Slide DST, Early detection, drug susceptibility, tuberculosis

1. Introduction

*Mycobacterium tuberculosis* cause the major air borne disease in human called TB. Due to drug resistance it is a major health problem in the world. In the world every year 9.4 million new cases detected, From that 1.5 million patient were died from TB according to WHO 2015(1). In 2013 22 countries reported 80% cases of TB. From 8.6 million cases 2.2 million cases occurred in India making India the world’s highest tuberculosis burden country(TB India 2014)(2). In world in 2013 there were 5% of TB cases estimated to have MDR-TB (3.5% new cases and 20.5 of previously treated TB cases)(1). Globally Drug resistance data show that an estimated 480000 people develop MDR-TB in 2013 and 210000 people died. In 2013 100 countries report XDR TB. On a average an estimated 9% of people with MDR-TB have XDR-TB. (1)

Multi drug resistance means Mycobacterium tuberculosis that are resistant to at least isoniazide and rifampicin- which are most effective anti- TB drug. MDR-TB cause by primary infection or during drug treatment (3). MDR TB patient carry strain resistant to the most anti tuberculosis drug, so it is very difficult to treat. There is chance to develop extensive drug resistance Tuberculosis from MDR TB. (XDR-TB)(4)

For control of TB ,early diagnosis is necessary. It is a mystery to diagnose mycobacterial infection (5). With the correct diagnosis and appropriate treatment disease can be cure (5). Solid egg based media like Lowenstein Jensen(LJ) and ogawa media used in most of developing countries for culture and susceptibility test due to low cost. Time taken by this method 3-8 week for culture and for susceptibility it takes further4-6 weeks. This is gold standard method, gives accurate result but take too much time.(6)

Bactec mycobacterial growth indicator tube (MIGT)- the fully automated commercial system gives rapid detection of resistance of first and second line drug within 10 days , but it require expensive equipment that are not easily available or suitable for resource limited countries & Gene expert is one of the system which give the result of susceptibility within three hours, it is rapid but require more expensive instrument (7).

In liquid media Mycobacterium can grow more rapidly than solid media, where it grow as micro colony ,based on this observation, a new , efficient , reliable and inexpensive method known as slide DST, which gives the result for detection within 10 days,(8) This method has to have the following desirable features like results as early as possible, simple training, easy interpretation, and allow start of the treatment as early as possible.(9)

Robert Koch was the first to use rapid slide culture technique using coagulated human serum, he was successful in obtaining the growth of Mycobacterium bacilli in 7 days but contamination hindered further success.(10)

For early and rapid diagnosis of drug resistance TB as well as MDRTB/XDR TB in poor setting countries slide DST may be suitable system. Accurate TB diagnosis can be made by good contamination control and sufficient equipment in smear positive cases (11). There is need for improved technologies for rapid detection of drug resistance because of the emergence of MDR-TB and XDR-TB in TB.(12). The
The objective of this study is early detection of drug resistance among TB patient and evolution of simple technique.

2. Material and Methods

A total of 495 clinical samples (sputum) were obtained from patients of suspected pulmonary TB from June 2013 to August 2015. The patient was asked to take early morning sputum. When he wakes up without brushing the teeth, asked to a deep breath and expectorates sputum in given wide mouthed sterile plastic container.

2.1 Staining procedure

Ziehl-Neelsen (ZN) staining was performed by standardize method.(13)

2.2 Slide DST by middle brook 7H9 broth

This method is used for detection of drug susceptibility of Mycobacterium tuberculosis to isoniazide (INH) and rifampicin (RIF). For identification between MTB and NMTB paranitro benzoic was used. The study adapted the method described by hemid et al. using middle brook 7H9 broth – a liquid media with oleic acid albumin dextrose catalase as growth supplement and PANTA used as decontaminant (0.1 ml/5 ml media) and anti-TB drugs. 7 ml drug containing and 7 ml drug free media were poured in two different sterile heavy MCcartnry bottle. For each sample 5 sputum smear were made direct from sputum sample on one end of autoclaved slide which was cut in half longitudinally. Slides were dipped in drug free and drug containing medium containing MCCartnry bottle. After 10 days slides were removed from bottle and dipped in sterile distilled water to remove excess blood for 5 min. After washing slides were air dried and heat fixed on hot plate for 30 min at 80 c. ZN staining was performed and examined under 100x for microcolonies. Any number of well developed colonies with cording in presence of drug was interpreted as resistance.

2.3 Slide DST by human blood media

For preparing the media, outdated but not more than 4 week old citrated human blood was used with an equal volume sterile deionized water; this blood was diluted to cause hemolysis. The media was made selective by adding polymixin-B (2,00,000 unit/l), carbenicillin (100mg/l), trimethoprim (10mg/l) and amphotericin-B (10mg/l) and adjusted the pH between 6.5 to 7.5. For each sputum sample 3 drug free media and 2 drug containing bottle 1 of 0.2µg/ml isoniazide and 1 of 1µg/ml rifampicin were prepared. For each sputum sample 5 sputum smear were made directly from sputum sample on the one end of the sterile autoclaved glass slide which was cut in half longitudinally. Slides were dipped in drug free and drug containing medium containing MCCartnry bottle. After 10 days slides were removed from bottle and dipped in sterile distilled water to remove excess blood for 5 min. After washing slides were air dried and heat fixed on hot plate for 30 min at 80 c. ZN staining was performed and examined under 100x for microcolonies. Any number of well develop colonies with cording in presence of drug was interpreted as resistance. The growth was recorded as follows

| 0  | No division of AFB as compared to Control slide |
|----|-------------------------------------------------|
| 1+ | Small clumps of up to 4 bacilli provided these were not present in control slide |
| 2+ | Large clumps of bacilli |
| 3+ | Large clumps with some cord formation |
| 4+ | Micro colonies with good cord |

Figure 1: Microcolonies of M.tuberculosis showing cord formation at 100x magnification under Bright Field Microscope.

Figure 2: Microcolonies of M.tuberculosis showing cord formation at 100x magnification under Bright Field Microscope.

3. Result

From 496 suspected pulmonary TB sample 207 samples were found positive by smear microscopy. All Smear positive samples saw culture positive by Slide DST and LJ method.

| Method for diagnosis | Total samples | No. of Positive Cases | % of positive cases |
|----------------------|---------------|-----------------------|-------------------|
| Smear microscopy      | 496           | 207                   | 41.73             |
| LJ                   | 496           | 207                   | 41.73             |
| Slide DST            | 496           | 207                   | 41.73             |

By Slide DST method 37 cases were mono resistance and 74 cases were multi drug resistance. (Table 2)
Slim et al. 2006 found 100% sensitivity and 62% specificity. Fahmida et al. 2011 found sensitivity of slide DST was 98.8% (14). Hamid sensitivity of slide DST method was 97% using HBM and in infection while under treatment (17). By this study the rate of contamination by slide DST in their study (11). Rate of study by conventional method, (18). One author found 0.18% resistance direct from sputum sample which are smear positive and introduce a simple and rapid method.

In Slide DST, 47% cases were found to have resistance against Isoniazide drug and 45% against rifampicin drug. Total 74 cases were found to have drug resistance against isoniazid and rifampicin (MDR) from 207 smear positive cases. Fahmida et al. 2011 author found 80% cases were detected as MDR TB in category 2 failure patient (14), Kamal et al. 2009 and Rahman et al. 2009 found 83% and 87% MDRTB respectively among category 2 failure patient by (15-16). The probable causes of above finding may be due to treatment with an inadequate drug regimen or re infection while under treatment (17). By this study the sensitivity of slide DST method was 97% using HBM and 96% using middlebrook 7H9 broth. Fahmida et al. 2011 found sensitivity of slide DST was 98.8% (14). Hamid Slim et al. 2006 found 100% sensitivity and 62% specificity (11).

7 bottles were contaminated in slide DST by middle brook 7H9 broth and 10 bottles were contaminated in Human blood media. rate of contamination was 1.4% and 2% in middle brook 7H9 broth and Human blood media respectively. Tortollet, et al., found 11% contamination in their study by conventional method, (18), one author found 0.18% contamination in slide DST method. Hamid et al. was found 7.4% contamination by slide DST in their study (11). Rate of contamination is lower than conventional method. Possible explanation of these authors findings are these of penicillin instead of PANTA or MAST tablet.

Drug susceptibility detection time by slide DST method in the present study was 10 days, and it correlates with other studies - 10 days by Hamid et al. 11, 8 days by Dickinson et al. 10 Whereas, detection time by conventional DST was 60.4±5.9 days (14). Accordingly, slide DST provides good opportunity for rapid identification of MDR strain. This early information is of great advantage in clinical settings to choose an appropriate drug regimen.

There are many limitation of slide DST although it is cheaper than automated culture method, it require costly media, growth and antibiotic supplement, which are not commonly available in field level laboratories in developing countries. Only microscopically positive sputum sample could be tested. As control strains cannot be used so its quality assurance is also a challenge. Slide DST is qualitative tests in which the observer confirms resistance by visualizing growth under microscope; where any well developed micro colony in presence of drug interprets as resistance to that drug. In slide DST there are no discrete colonies to count and a proportion cannot be calculated. As a result slide DST might over diagnose resistance, that is, isolates read as susceptible by reference method. Slide DST can provide good opportunity for rapid.

Slide DST can provide good opportunity for rapid detection of drug resistance TB including MDR/XDR strain from direct sample within 10 days. This method can be implemented as rapid diagnostic tool to detect drug resistance tuberculosis.

| Table 2: Result of drug resistance by Slide DST method |
| --- | --- | --- |
| Resistance pattern | Anti-TB drug | Slide DST by HBM | Slide DST by 7H9 broth |
| Mono | INH | 20 | 20 |
| RMP | 17 | 16 |
| Multi | INH+RMP | 74 | 73 |
| Susceptible | 96 | 98 |
| Total | 207 | 207 |

| Table 3: Comparison of time required for diagnosis of Drug resistance TB |
| --- | --- |
| Methods for diagnosis | Time taken for the result |
| Conventional Method | 60 days |
| Slide DST by HBM | 10 days |
| Slide DST by Middlebrook 7H9 | 10 days |

| Table 4: Comparison of cost of methods for diagnosis of Drug resistance TB |
| --- | --- |
| Methods for diagnosis | Cost (Rs.) |
| Conventional LJ | 250 |
| Slide DST by HBM | 20 |
| Slide DST by Middlebrook 7H9 | 45 |

4. Discussion

Early drug susceptibility test is important in tuberculosis patient. Because of there are chance of treatment failure patient it is more important. In India DST is done on solid media because it is low cost and it takes 2-3 month time. Present study was designed to rapid detection of drug resistances direct from sputum sample which are smear positive and introduce a simple and rapid method.

Statically analysis shows Sensitivity of Slide DST using HBM have 97% and 96% by Middlebrook 7H9. Time taken for the result was much less then proportion method. (Table 3)

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