RESEARCH ARTICLE

Antimicrobial susceptibility pattern of clinical isolates of *Burkholderia pseudomallei* in Bangladesh

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Abstract

**Background:** Melioidosis an infectious disease, caused by a Gram negative bacterium called *Burkholderia pseudomallei*, is endemic in Bangladesh. This organism is sensitive to limited number of antimicrobial agents and need prolonged treatment. There is no comprehensive data on the antimicrobial susceptibility profile of *B. pseudomallei* isolated in Bangladesh over last several years. The present study aimed to determine the antimicrobial susceptibility pattern of *B. pseudomallei* isolated in a tertiary care hospital of Dhaka city from 2009 to 2015.

**Methods:** All *B. pseudomallei* isolated from melioidosis patients over a period of 7 years (2009–2015) in the Department of Microbiology of a 725-bed tertiary care referral hospital in Dhaka city, Bangladesh were included in the study. *B. pseudomallei* was identified by Gram stain, culture, specific biochemical tests, serology and PCR using specific primers constructed from 16s rRNA region of *B. pseudomallei*. Antimicrobial susceptibility to specific agents was determined by disk diffusion and minimum inhibitory concentration methods.

**Results:** A total of 20 isolates of *B. pseudomallei* which were isolated from patients coming from different geographic locations of Bangladesh were included in the study. All the isolates were uniformly sensitive (100%) to ceftazidime, imipenem, piperacillin–tazobactam, amoxicillin–clavulanic acid and tetracycline by both disk diffusion and MIC methods. Two strains were resistant to trimethoprim–sulfamethoxazole by disk diffusion method but were sensitive by MIC method. The MIC50 and MIC90 values of the above antimicrobial agents were almost similar. All the isolates were resistant to amikacin by both MIC and disk diffusion methods.

**Conclusion:** The results of the study suggest that *B. pseudomallei* prevalent in Bangladesh were still susceptible to all recommended antimicrobial agents used for the treatment of melioidosis. However, regular monitoring is needed to detect any emergence of resistance and shifting of MIC50 and MIC90 values.

**Keywords:** *Burkholderia pseudomallei*, Antimicrobial susceptibility, Bangladesh, Minimum inhibitory concentration

Background

*Burkholderia pseudomallei* is a Gram negative saprophytic bacterium that causes melioidosis. It is an endemic disease of public health and clinical importance in many countries of the world including Bangladesh [1–3]. Several cases have been reported from Bangladesh since 1988 [3, 4]. Recently, we have found that the organism is present in the soil samples of Bangladesh and about 22.6–30.8% of people residing in melioidosis endemic districts of Bangladesh were sero-positive for anti-*B. pseudomallei* IgG antibody [5].

Melioidosis may be manifested as localized or disseminated infection involving multiple organs of the body. The disease is usually fatal if left untreated. Treatment of melioidosis requires prolonged courses of antibiotics. The current treatment regime has an intensive and maintenance phase. In the intensive phase, ceftazidime is usually the drug of choice. However, carbapenem (imipenem/meropenem) is recommended in severe infection and treatment failure cases. This is followed by the
maintenance phase which consists of oral administration of TMP–SMX or TMP–SMX and doxycycline or amoxicillin–clavulanic acid [6–9]. But recently, there are reports of resistant B. pseudomallei to ceftazidime, TMP–SMX, amoxicillin–clavulanic acid from different melioidosis endemic countries of the world including neighboring India [9–15].

Currently, no comprehensive information is available on the antimicrobial susceptibility pattern of B. pseudomallei isolated from patients in Bangladesh. Therefore, the present study was undertaken to determine the antibiotic susceptibility pattern of B. pseudomallei isolated from patients during the period of 2009–2015 in our hospital by disc diffusion and minimum inhibitory concentration (MIC) methods.

Methods

Study organisms

All B. pseudomallei isolated from melioidosis patients over a period of 7 years (January 2009 to December 2015) in the Department of Microbiology of a 725-bed tertiary care referral hospital in Dhaka city, Bangladesh were included in the study. B. pseudomallei was identified by typical colony morphology, Gram staining (bipolar staining), motility, biochemical tests, arabinose assimilation and resistance to colistin and aminoglycoside [16]. Monoclonal antibody based latex agglutination test (Melioidosis Research Center, Khon Kaen, Thailand) was performed for the final identification and confirmation of B. pseudomallei. Serologically confirmed B. pseudomallei isolates were further confirmed by PCR using specific primers constructed from 16s rRNA region of B. pseudomallei [17].

Determination of MIC

Minimum inhibitory concentrations of all antibiotics against B. pseudomallei were determined by agar dilution method [18]. Briefly, Muller–Hinton agar plates containing different dilutions of specific antimicrobial agent were spot inoculated with $10^4$ organisms. Each spot was 5–8 mm in diameter. All reading was taken between 18 and 24 h of incubation at 37 °C in aerobic condition. The MIC represented the lowest concentration of antimicrobial agents at which complete inhibition of growth of the organism occurred. The MIC (µg/ml) interpretation for susceptible (S), intermediate (I), and resistant (R) for amoxicillin–clavulanic acid, ceftazidime, imipenem, tetracycline and TMP–SMX was carried out following the CLSI approved guideline M45-A2 [19]. As reported earlier, the MIC breakpoint of P. aeruginosa for piperacillin–tazobactam, amikacin and gentamicin as mentioned in CLSI guideline was used to interpret the MIC of B. pseudomallei [14, 20]. The detail MIC interpretive values used in the study are shown in Table 1.

Disc diffusion method

The isolates were tested for susceptibility to ceftazidime (30 µg), imipenem (10 µg) piperacillin–tazobactam (30 µg), amoxicillin–clavulanic acid (30 µg), tetracycline, TMP–SMX (1.25/23.75 µg), gentamicin (10 µg) and amikacin (30 µg) by disc diffusion technique [21]. Briefly, a 0.5 McFarland suspension of each bacterial isolate was made in normal saline and inoculated onto Mueller–Hinton agar (Oxoid Ltd, UK) to prepare a uniform lawn. The disks were applied at a specific distance from each other and the zone of inhibition around the antibiotic disc was

| Antimicrobial agents | Disk diffusion test | MIC (µg/ml) |
|----------------------|---------------------|-------------|
|                      | Disk content (µg)   | Zone diameter (mm)   | R | I | S | S | I | R |
| Ceftazidime<sup>a,b</sup> | 30                   | ≤14          | 15–16 | ≥18 | ≤8 | 16 | ≥32 |
| Imipenem<sup>b</sup> | 10                   | ≤15          | 16–18 | ≥19 | ≤4 | 8  | ≥16 |
| Amox–Clav<sup>a,d</sup> | 20/10               | ≤13          | 14–17 | ≥18 | ≤8/4 | 16/8 | ≥32/16 |
| Pip–Tazo<sup>c</sup> | 100/10              | ≤14          | 15–20 | ≥21 | ≤16/4 | 32/4–64/4 | ≥128/4 |
| TMP–SMX<sup>a,d</sup> | 1.25/23.75          | ≤10          | 11–15 | ≥16 | ≤2/38 | – | ≥4/76 |
| Tetracycline<sup>c</sup> | 30                  | ≤11          | 12–14 | ≥15 | ≤4  | 8  | ≥16 |
| Amikacin<sup>c</sup> | 30                  | ≤14          | 15–16 | ≥17 | ≤16 | 32 | ≥64 |
| Gentamicin<sup>c</sup> | 10                  | ≤12          | 13–14 | ≥15 | ≤4  | 8  | ≥16 |

<sup>a</sup> MIC interpretive standards defined in M45-A2, CLSI guideline for B. pseudomallei [19]<br><sup>b</sup> Zone diameter interpretive standards of CLSI for P. aeruginosa [19]<br><sup>c</sup> MIC interpretive standard of CLSI for P. aeruginosa [20]<br><sup>d</sup> Zone diameter interpretive standards of CLSI for Enterobacteriaceae by disc diffusion method [20]
measured after 18 h of incubation of plates at 37 °C. Since interpretative standards for zone sizes of tested antibiotics are not available for *B. pseudomallei*, we used the Clinical and Laboratory Standards Institute [20] threshold zone sizes for members of the family *Enterobacteriaceae* and *Pseudomonas aeruginosa* as reported earlier [14]. The zone of inhibition was interpreted as sensitive, intermediate and resistant according to CLSI guideline. The zone sizes used for interpretation are shown in Table 1.

Each isolate was tested in duplicate by disk diffusion and MIC methods. All disks and antibiotics were obtained from Oxoid Ltd., Basingstoke, Hampshire, UK and Sigma, USA. Potency of the disks and antimicrobial agents were standardized using the reference strain *E. coli* ATCC 25922.

**Discussion**

Mortality in patients with *B. pseudomallei* infection is high and early diagnosis and specific antimicrobial therapy is necessary to reduce the fatal outcome [21-23]. The antibiotic of choice for melioidosis is ceftazidime, a third generation cephalosporin. Imipenem is considered as alternatives to ceftazidime. Since relapse rates are high, initial parenteral treatment followed by maintenance therapy with oral TMP–SMX (8–12 & 40–62 mg/kg/day) or doxycycline (4 mg/kg/day) or amoxicillin–clavulanic acid (20/5 mg/kg 8 h) for 12–20 weeks are recommended [6-8]. However, *B. pseudomallei* resistant to ceftazidime, tetracycline/doxycycline and TMP–SMX, has been recently reported from Malaysia, Thailand and Australia. About 0.6% *B. pseudomallei* in Malaysia was found to be resistant to imipenem, ceftazidime, doxycycline and amoxicillin–clavulanic acid while the rate was 10% for TMP–SMX [9]. The rate was around 0.6% in Thailand [14]. About 4% *B. pseudomallei* in Australia was reported as resistant to imipenem, ceftazidime, doxycycline and TMP–SMX [12].

In Bangladesh there was no comprehensive data available on the antibiotic susceptibility pattern of *B. pseudomallei*. Bangladesh is a melioidosis endemic country and several cases have been documented since 1988 and, recently we have reported its presence in the soil from a district where several cases of melioidosis were diagnosed [3-5]. Organisms included in the present study were isolated from patients mainly coming from different areas of north and north-eastern part of Bangladesh during the period of 2009–2015. The result showed that all the isolates were uniformly sensitive to imipenem, ceftazidime, piperacillin–tazobactam, tetracycline, amoxicillin–clavulanic acid and TMP–SMX. The absence of resistance to these antibiotics in our isolates might be due to the fact that only a few numbers of organisms were tested.

| Antimicrobial agents | Disk diffusion test | MIC test |
|----------------------|---------------------|----------|
|                      | Disk content (µg)   | Sensitive N (%) | Zone diameter range (mm) | Sensitive N (%) | MIC50 (µg/ml) | MIC90 (µg/ml) |
| Ceftazidime          | 30                  | 20 (100)      | 24–32                  | 20 (100)      | 2           | 2           |
| Imipenem             | 10                  | 20 (100)      | 33–40                  | 20 (100)      | 2           | 2           |
| Amox–Clav            | 20/10               | 20 (100)      | 19–28                  | 20 (100)      | 8/4         | 8/4         |
| Pip–Tazo             | 100/10              | 20 (100)      | 35–39                  | 20 (100)      | 2/4         | 2/4         |
| TMP–SMX              | 1.25/23.75          | 18 (90)       | 15–28                  | 20 (100)      | 1/19        | 2/38        |
| Tetracycline         | 30                  | 20 (100)      | 17–31                  | 20 (100)      | 0.5         | 1           |
| Amikacin             | 30                  | 0 (0)         | 10–15                  | 0 (0)         | 64          | 64          |
| Gentamicin           | 10                  | 0 (0)         | 0                      | ND            | ND          | ND          |

**Table 2 Results of disk diffusion and MIC tests of *B. pseudomallei* isolated during 2009–2015 (N = 20)**
Therefore, in our settings, these antimicrobial agents can empirically be used to treat suspected melioidosis patients prior to the availability of culture and sensitivity results. Comparing the results of the disk diffusion test with that of MIC it appeared that the CLSI recommended breakpoint of zone diameter for *P. aeruginosa*, and *Enterobacteriaceae* could be used to interpret the zone of inhibition by *B. pseudomallei* in disk diffusion test where facilities for MIC test is lacking or till the zone diameter of *B. pseudomallei* for these agents are standardized. MIC and disk diffusion tests showed concordant results for all antimicrobials except for TMP–SMX. Only, two isolates were resistant to TMP–SMX by disk diffusion method (zone diameter 15 mm) whereas both were found sensitive to TMP–SMX by MIC method (≤0.19 and 2/38 µg/ml). Since MIC method is considered as the gold standard for determining the susceptibility of *B. pseudomallei* to TMP–SMX [19], therefore, all our 20 isolates were actually sensitive to TMP–SMX. The interpretation of susceptibility of *B. pseudomallei* to TMP–SMX by disk diffusion test is sometimes affected by slight hazy growth within the zone of inhibition and blurring of margin. Therefore, the doubtful result of disk diffusion test for TMP–SMX should be confirmed by MIC method.

*Burkholderia pseudomallei* is intrinsically resistant to aminoglycosides like gentamicin and amikacin [24]. These antibiotics are used as diagnostic agents for identification of *B. pseudomallei* in culture. Also, most selective media incorporate gentamicin in the medium as a selective agent to isolate *B. pseudomallei* from soil and other samples. Though *B. pseudomallei* is said to be resistant to aminoglycosides, there have been reports of aminoglycoside sensitive isolates (0.1%) from Thailand [25, 26], and from Australia [27]. Recently it has been reported that 86% of *B. pseudomallei* isolates in Sarawak, Malaysian Borneo were sensitive to gentamicin [28]. The strains belonged to sequence type 881 and 997. In view of the above, we have determined the susceptibility of our strains to amikacin, an aminoglycoside. All the 20 isolates were resistant to amikacin by both disk diffusion (zone diameter range 10–15 mm) and MIC method (MIC 64–512 µg/ml). It may be mentioned that all these strains were also resistant to gentamicin where no zone of inhibition was observed by disk diffusion method. Therefore, it is assumed that amikacin and gentamicin may be used safely as a diagnostic disk and in selective media for isolation of the organism from soil and other contaminated samples.

**Conclusion**

The study, though with limited number of isolates, revealed that the prevalent *B. pseudomallei* in Bangladesh were susceptible to recommended antimicrobial agents used for both intensive and maintenance phase therapy. Continuous monitoring of antimicrobial susceptibility should be maintained to detect emergence of resistant organism.

**Abbreviations**

TMP–SMX: trimethoprim–sulfamethoxazole; Pip–Tazo: piperacillin–tazobactam; Amox–Clav: amoxicillin/clavulanic acid; PCR: polymerase chain reaction; MIC: minimum inhibitory concentration; IgG: immunoglobulin G; CLSI: Clinical and Laboratory Standards Institute.

**Authors’ contributions**

SD performed the culture, identification and antimicrobial susceptibility tests. SH performed the antimicrobial susceptibility tests, literature search and manuscript writing. MRH was involved in storage and identification of organisms and primary data analysis. JAH was involved in study design, analysis of the data and manuscript writing. All authors read and approved the final manuscript.

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**Acknowledgements**

Not applicable.

**Competing interests**

The authors declare that they have no competing interests.

**Availability of data and materials**

All relevant data of this study are within the paper.

**Funding**

The work was done with the departmental funds.

**Publisher’s Note**

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**Received: 1 November 2016 Accepted: 13 July 2017**

**20 July 2017**

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