Melioidosis by aminoglycoside susceptible *Burkholderia pseudomallei*: First case in Bangladesh

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Abstract

*Burkholderia pseudomallei* is the etiological agent of melioidosis. It is a gram-negative bacillus present in environment and intrinsically resistant to many antibiotics including aminoglycosides. However, recently aminoglycoside susceptible *B. pseudomallei* has been isolated from melioidosis cases and reported from some countries of the world. But, such aminoglycoside susceptible *B. pseudomallei* has never been detected in Bangladesh either from melioidosis cases or from environment. All the *B. pseudomallei* isolated so far in Bangladesh were resistant to gentamicin and other aminoglycosides. Here, we describe a disseminated case of melioidosis caused by aminoglycoside susceptible *B. pseudomallei* in a 55 years old Bengali male patient. This is the first case of melioidosis due to aminoglycoside susceptible *B. pseudomallei* in Bangladesh.

Introduction

*Burkholderia pseudomallei*, a gram-negative bacillus, causing melioidosis is intrinsically resistant to aminoglycosides, macrolides and colistin [1-2]. This intrinsic resistance to aminoglycosides and colistin is used as identification criterion for *B. pseudomallei*. Also, these antibiotics are used to prepare selective media for isolation of *B. pseudomallei*. However, recently aminoglycoside susceptible *B. pseudomallei* strains have been detected in some regions of Malaysia, Thailand and Australia [3-6]. But, such aminoglycoside susceptible *B. pseudomallei* has so far not been detected in Bangladesh. All the *B. pseudomallei* isolated and reported from Bangladesh were resistant to aminoglycosides [7]. But, here we describe a disseminated case of melioidosis caused by aminoglycoside susceptible *B. pseudomallei* in a 55 years old Bengali male patient.

Case Summary

A 55 years old smoker, non-diabetic male presented with a 3 month history of high grade fever, non-productive cough and weight loss. About a year back, he developed intermittent to high grade fever, cough and loss of appetite and was diagnosed as a smear negative pulmonary tuberculosis case and treated accordingly in a local hospital. Initially, his symptoms improved, but about 3 months back he gradually developed high grade fever again, the highest recorded temperature being 104°F. With deteriorating symptoms he was admitted to Medicine unit of Dhaka Medical College Hospital. Four days after admission he developed pain and swelling on the left elbow. About 1 month following admission, the patient developed sudden, severe headache along with a single episode of vomiting followed by restlessness and disorientation. He had 5 episodes of seizures in 2 days with loss of consciousness (Glasgow Coma Scale was 3/15).

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On general examination, patient was mildly anemic, dyspneic and the temperature was 103°F. Respiratory system examination revealed features suggestive of consolidation. The left elbow joint was erythematous, swollen (4cm×4cm), tender without any discharge or regional lymphadenopathy. Liver was just palpable. Blood analysis yielded hemoglobin 9.6 g/dl, ESR 70 mm at first hour, total white cell count 6.8×10^9/L, platelets 70×10^9/L, SGPT 80 U/L and SGOT 173 U/L. HbsAg and anti HBc IgM was positive. Chest radiograph showed patchy and in-homogenous opacities in both upper, mid and right lower zone of both lungs (Fig-1). Blood, sputum, urine and cerebrospinal fluid (CSF) culture yielded no growth. Sputum for acid fast bacilli (AFB) and Mycobacterium tuberculosis by Ziehl–Neelsen stain and MTB/RIF–GeneXpert test was negative respectively. Ultrasonography of whole abdomen revealed mild hepatosplenomegaly and grossly enlarged prostate. Magnetic resonance imaging (MRI) of brain exhibited features suggestive of venous infarct due to suspected venous thrombosis involving superior sagittal sinus.

Pus was aspirated from the left elbow joint swelling using a sterile syringe. Gram stain of the pus showed gram-negative bacilli arranged in bipolar ‘safety pin’ pattern. Culture of the pus yielded growth of gram-negative and oxidase positive bacilli in Blood and MacConkey agar media, but no growth was detected on modified Ashdown’s selective media containing gentamicin 5µg/ml. The isolate was identified as B. pseudomallei by colony morphology and biochemical tests [8]. The isolate was further confirmed by monoclonal antibody based latex agglutination test for B. pseudomallei (Melioidosis Research Center, Khon Kaen, Thailand). Polymerase Chain Reaction (PCR) and Loop Mediated Isothermal Amplification based assay (LAMP) using B. pseudomallei specific primers (Table-1) were also performed for further confirmation of the isolate. Both PCR and LAMP tests confirmed the isolate as B. pseudomallei (Fig-2 and Fig-3). The isolate was sensitive to ceftazidime, meropenem, amoxicillin+clavulanic acid, piperacillin+tazobactem and aminoglycosides namely gentamicin, amikacin and netilmicin (Table-2) and resistant to trimethoprim-sulphamethoxazole (TMP-SMX) and colistin. Since aminoglycoside susceptible B. pseudomallei was never been detected in Bangladesh further enquiry was made to track the possible source of the organism. On enquiry, it was found that he had been a construction worker in Malacca, Malaysia for the past 10 years where he developed high grade intermittent fever, cough and loss of appetite about a year ago and then he returned to Bangladesh. Based on the above, the patient was finally diagnosed as a case of disseminated melioidosis by aminoglycoside susceptible B. pseudomallei. Probably, the patient could have acquired the infection while in Malaysia because such aminoglycoside susceptible B. pseudomallei strains are prevalent there. The patient was successfully treated with standard antibiotic regimen for melioidosis and discharged with improved general condition.

Fig.1: Chest X-Ray P/A view showing patchy opacities in upper, middle and lower zone of lungs
Table 1: Primers targeting TTS1 gene used in conventional PCR [9] and in-house LAMP [10]

<table>
<thead>
<tr>
<th>Primers for PCR</th>
<th>Primer sequence</th>
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<tbody>
<tr>
<td>Forward primer</td>
<td>5’-CGTCTCTTACTGTCGAGCAATCG-3’</td>
</tr>
<tr>
<td>Reverse primer</td>
<td>5’-CGTGACACCGGTCAGTACTC-3’</td>
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<thead>
<tr>
<th>Primers for LAMP</th>
<th>Primer sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Forward Outer Primer (F3-2)</td>
<td>TCGATTGCTTCGACAGCG</td>
</tr>
<tr>
<td>Forward Inner Primer (FIP-2)</td>
<td>TTGCCTCACGATCAAACGATCCAAATTTCGCTCAG</td>
</tr>
<tr>
<td>Backward Outer Primer (B3-2)</td>
<td>AATCCCTGCCCGAATTG</td>
</tr>
<tr>
<td>Backward Inner Primer (BIP-2)</td>
<td>CTCGTATGCGCAGGATCAGGCGCGACG</td>
</tr>
<tr>
<td>Forward Loop Primer (LF)</td>
<td>ATACCGTCAATGTCCGACCCG</td>
</tr>
<tr>
<td>Backward Loop Primer (LB)</td>
<td>CGGCATCGATAACACAG</td>
</tr>
</tbody>
</table>

Table 2: Results of disc diffusion and MIC tests of isolated B. pseudomallei

<table>
<thead>
<tr>
<th>Antimicrobial agents</th>
<th>Disk diffusion test</th>
<th>MIC test (µg/ml)</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Zone of inhibition of test organism (mm)</td>
<td>Sensitive zone diameter breakpoints (mm)</td>
<td>MIC of test organism (µg/ml)</td>
</tr>
<tr>
<td>Gentamicin&lt;sup&gt;a&lt;/sup&gt;</td>
<td>22</td>
<td>≥15</td>
<td>1</td>
</tr>
<tr>
<td>Amikacin&lt;sup&gt;a&lt;/sup&gt;</td>
<td>19</td>
<td>≥17</td>
<td>2</td>
</tr>
<tr>
<td>Netilmicin&lt;sup&gt;a&lt;/sup&gt;</td>
<td>20</td>
<td>≥15</td>
<td>Not done</td>
</tr>
</tbody>
</table>

Note: a=Zone diameter interpretive standards of CLSI for P. aeruginosa by disk diffusion method [11].
Discussion

In Bangladesh, the first case of melioidosis was reported in 1988 [12]. Since then, about 54 human melioidosis cases from different districts of Bangladesh have been recorded till 2018 [13]. In addition, *B. pseudomallei* was isolated from soil samples of Gazipur district [14], rendering Bangladesh as a ‘definite’ melioidosis endemic country.

*B. pseudomallei* is known to be intrinsically resistant to aminoglycosides [2]. Resistance to aminoglycosides and colistin is an important identification criterion for *B. pseudomallei*. All *B. pseudomallei* that have been isolated in Bangladesh till now were resistant to amikacin and gentamicin by both Kirby-Bauer disk diffusion and MIC methods [7]. However, aminoglycoside sensitive *B. pseudomallei* have been isolated in certain regions of Southeast Asia and Australia [3-6]. There have been reports of rare cases of melioidosis due to aminoglycoside susceptible strains (0.1%) in Thailand [5] and in Australia [6]. On the contrary, 86% of *B. pseudomallei* isolates in Sarawak, Malaysian Borneo were found sensitive to Gentamicin [3]. However, in Kedah, Malaysia, about 21% and 6% of isolates were susceptible to gentamicin and amikacin respectively [4].

Resistance to aminoglycosides in *B. pseudomallei* is due to *amrAB-oprA*-mediated efflux [5,15]. Studies comprising of genome sequencing have revealed that a mutation or absence of *amr*-8 transcripts that encodes for the multidrug efflux system accounts for the susceptibility to aminoglycosides as well as macrolides in the aforementioned isolates [3-4]. Apparently, our patient with disseminated melioidosis was residing in Malacca, Malaysia for the past 10 years. So, it could be reasonably assumed that this particular strain could be acquired in Malaysia where aminoglycoside susceptible strains had previously been detected. However, we should not dismiss the possibility of presence of such aminoglycoside susceptible strains indigenously in Bangladesh. True origin of our aminoglycoside susceptible isolate could be determined if we could do sequence analysis or multilocus sequence typing.

In a resource limited country like ours, culture of *B. pseudomallei* constitutes the diagnostic gold standard. Intrinsic resistance of *B. pseudomallei* to aminoglycosides is used for the development of selective media for its isolation from samples like sputum or soil that contain other organisms. This particular case demonstrates that the use of gentamicin incorporated selective media might fail to detect such susceptible strains and would undermine the true extent of its presence in environment and clinical samples.

Author contributions

SF did the experiments and wrote the manuscript; MSAJ supervised the work and contributed in writing the manuscript; AA collected the clinical data and involved in the management of the patient; JAH contributed in writing and editing the manuscript.

Conflict of interest

The authors hereby, declare that no conflict of interest exists.

Ethical statement

Written consent was obtained from the patient for publication of the case.

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References


