Case Report

Melioidosis presenting with periorbital cellulitis and eyelid abscess in Sarawak, Malaysian Borneo - A Case Report

Chee Yik Chang1


DOI: http://doi.org/10.4038/sljid.v9i2.8259

Abstract

Melioidosis is an infectious disease caused by the Gram-negative bacillus *Burkholderia pseudomallei* and presents with a broad spectrum of clinical manifestations and organ involvement. Ocular manifestation in melioidosis is rare. We present a case of periorbital cellulitis and eyelid abscess caused by *B. pseudomallei* in a diabetic patient successfully treated with antibiotic and surgical drainage.

Keywords: Melioidosis, *Burkholderia pseudomallei*, Periorbital cellulitis, Eyelid abscess

Introduction

Melioidosis is caused by *Burkholderia pseudomallei*, a Gram negative bacillus found in soil and water. It is an endemic disease in northern Australia and Southeast Asian countries, especially Thailand and Malaysia. Despite the availability of effective antimicrobial therapy in the treatment of melioidosis, the mortality rate remains high.1 Melioidosis is difficult to diagnose because of its diverse clinical presentation and can affect almost any organ. The manifestations of melioidosis are highly variable, ranging from pneumonia, septic arthritis, liver or splenic abscesses, pericardium and parapharyngeal space abscesses, skin abscesses, pan ophthalmitis to fatal septicemia.2 Ocular involvement in melioidosis is rare and the prevalence was estimated at 0.49-1.02%.3 A 23-year retrospective review of 16 cases of ocular involvement in melioidosis in Thailand revealed orbital cellulitis as the most common manifestation (7 cases) followed by endophthalmitis (4 cases), preseptal cellulitis (2 cases), pan ophthalmitis (2 cases), and pan uveitis (1 case).4 Melioidosis can also present with lid abscess in isolation without orbital cellulitis.4

Case Presentation

A 58-year-old male farmer presented in May 2018 to Kapit Hospital, Sarawak with fever and right eye swelling of 1 week duration. His past medical history included bacteraemic melioidosis which was diagnosed in 2016. At that time, he received
intravenous ceftazidime as intensive phase therapy, followed by eradication therapy with oral trimethoprim-sulfamethoxazole. He was also diagnosed with type 2 diabetes mellitus during that hospitalization and started on oral anti-diabetic medications. However, he had defaulted on his medications for 1 year.

Upon arrival to the hospital, his general condition was stable and the vital signs were normal. The random blood sugar was 31.2 mmol/L. There was periorbital swelling of the right eye with swelling and erythema of the upper and lower eyelids. The conjunctiva appeared normal. The visual acuity of the right eye was 6/15 while vision of the left eye was reduced to finger counting. The remainder of systemic examination was unremarkable.

Haematological analysis showed a haemoglobin of 15.6 g/dL, white blood cell count of 7.4 x 10^3/μL and platelet count of 162 x 10^3/μL. His renal and liver function tests were within normal limits. The chest radiograph and abdominal ultrasonography did not reveal any abnormality. Our initial working diagnosis was right periorbital cellulitis. He was started on intravenous ampicillin-sulbactam 1.5 gram 8-hourly. However, persistent fever was observed over the next 3 days. At this point, melioidosis was suspected in view of multiple risk factors including diabetes mellitus, occupational exposure and previous history of melioidosis. The antibiotic regime was escalated to intravenous ceftazidime 2 gram 8-hourly to treat for melioidosis.

He was also referred to the ophthalmologist for an opinion. Computed tomography (CT) of the orbit showed right periorbital cellulitis (Fig. 1).

Detailed ophthalmic examination showed mild non-proliferative diabetic retinopathy in both eyes and increased cup-to-disc ratio of the left eye. As a result, floater only vitrectomy of the left eye was done by the ophthalmology team.

On day 8 of hospitalization, he developed an abscess over the right upper eyelid. Incision and drainage of the abscess was performed in which copious amounts of pus was drained and sent for microbiological analysis. The cultures of blood and pus from the eyelid abscess yielded B. pseudomallei, grown on modified Ashdown selective culture medium. The organism was susceptible to ceftazidime, amoxicillin-clavulanic acid and trimethoprim-sulfamethoxazole. The identification of B. pseudomallei was confirmed by a positive PCR assay specific for the detection of B. pseudomallei.
He completed a 2-week course of intravenous ceftazidime. The fever completely settled on day 10 of hospitalization and the right eye swelling progressively reduced. His blood glucose control was good while on insulin therapy. He completed eradication therapy consisting of trimethoprim-sulfamethoxazole and doxycycline for 20 weeks. During a follow-up review at the medical clinic, he was well and showed no signs of disease recurrence. The right eye swelling had totally resolved, and visual acuity returned to baseline levels.

**Timeline of clinical progression**

<table>
<thead>
<tr>
<th>Time</th>
<th>Events</th>
<th>Actions/treatments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day 1</td>
<td>Admission for fever and right eye swelling for 1 week; treated as right periorbital cellulitis</td>
<td>IV ampicillin-sulbactam 1.5 gram 8-hourly</td>
</tr>
<tr>
<td>Day 3</td>
<td>Persistent fever and suspected melioidosis</td>
<td>IV ceftazidime 2 gram 8-hourly</td>
</tr>
<tr>
<td>Day 4</td>
<td>CT orbit showed right periorbital cellulitis</td>
<td>Co-managed by ophthalmology team</td>
</tr>
<tr>
<td>Day 5</td>
<td>Blood culture grew <em>B. pseudomallei</em></td>
<td></td>
</tr>
<tr>
<td>Day 8</td>
<td>Appearance of right eyelid abscess</td>
<td>Incision and drainage (pus culture grew <em>B. pseudomallei</em>)</td>
</tr>
<tr>
<td>Day 10</td>
<td>Resolution of fever and reducing right eye swelling</td>
<td>Antibiotic continued</td>
</tr>
<tr>
<td>Day 14</td>
<td>Completed IV ceftazidime for 14 days</td>
<td>Discharged home well with trimethoprim-sulfamethoxazole and doxycycline</td>
</tr>
</tbody>
</table>

**Discussion**

Melioidosis can present with a wide spectrum of clinical presentations. Pneumonia was the principal presentation of melioidosis in approximately half of the cases. Less common presentations include genitourinary infection, skin infection, bacteremia without evident focus, septic arthritis or osteomyelitis and neurological melioidosis. Ocular manifestation in melioidosis is rare. There have been several case reports previously on corneal ulcers, orbital cellulitis and endophthalmitis caused by *B. Pseudomallei*. Melioidosis presenting with periorbital cellulitis and eyelid abscess has not been reported previously.

Periorbital cellulitis, also known as preseptal cellulitis is an infection of the eyelid and superficial periorbital soft tissues without the involvement of the globe and orbit. Patients with periorbital cellulitis generally do not require surgical intervention except in cases of eyelid abscess where drainage of abscess is recommended in addition to antibiotic treatment. Culture of the material is important to confirm the diagnosis and as a guide to appropriate antibiotic therapy.

There are 3 modes of acquisition of *B. pseudomallei* which could result in melioidosis infection, namely inhalation, ingestion, and inoculation. The patient described in this case was a farmer who had constant exposure to soil or water in which this organism is found. We postulated that the most probable mode of acquisition was through direct inoculation into the affected eye causing primary ocular melioidosis and later, blood
stream infection. This is consistent with the observation that most cases of ocular melioidosis are associated with *B. pseudomallei* bacteraemia, as reported by Yaisawang *et al.*

Early diagnosis and prompt treatment are crucial for a favourable outcome. Periorbital cellulitis, if not treated promptly, can extend posteriorly into the orbit causing orbital cellulitis, subperiosteal abscess or orbital abscess. These would potentially lead to significant visual and central nervous system complications. In this case, our patient achieved resolution of periorbital cellulitis and eyelid abscess without any long term ophthalmic complication following early diagnosis of melioidosis, effective antibiotic therapy and surgical drainage.

**Conclusion**

Diagnosis of melioidosis should be considered in patients who present with periorbital cellulitis or eyelid abscess in areas where melioidosis is endemic. Surgical drainage of the abscess is an important part of management besides standard antibiotic treatment.

**Acknowledgements**

I would like to thank the Director General of Health Malaysia for his permission to publish this article. The author thanks the Director of Kapit Hospital, Dr Hii King Ching for her guidance and support, Dr Tan Li Mun, Ophthalmologist at Sibu Hospital for her expert opinion, and all staff of Kapit and Sibu Hospitals involved in the care of this patient.

**Conflict of interest:** The author declares that there are no conflicts of interest regarding the publication of this paper.

**Consent for publication:** Written informed consent for publication of the clinical details and clinical images was obtained from the patient.

**References**