

# Brunei International Medical Journal

OFFICIAL PUBLICATION OF  
THE MINISTRY OF HEALTH  
AND  
UNIVERSITI BRUNEI DARUSSALAM

Volume 18

22 April 2022 (20 Ramadan 1443H )

## DACRYOADENITIS WITH MULTIPLE UNILATERAL OCULAR ABSCESES IN DISSEMINATED MELIOIDOSIS.

SYLVES P<sup>1</sup>, SEAH RB<sup>2</sup>, CAROLINE B<sup>2</sup>, YAP JY<sup>1</sup>, SHUAIBAH AG<sup>1</sup>, HANIDA H<sup>3</sup>.

<sup>1</sup>Department of Ophthalmology, Faculty of Medicine and Health Sciences Universiti Malaysia Sabah, Jalan UMS, 88400 Kota Kinabalu, Sabah, Malaysia.

<sup>2</sup>Department of Ophthalmology, Queen Elizabeth Hospital, 88586 Kota Kinabalu, Sabah, Ministry of Health Malaysia.

<sup>3</sup>Oculoplastic and Reconstructive Unit, Department of Ophthalmology, Queen Elizabeth Hospital, 88586 Kota Kinabalu, Sabah, Ministry of Health Malaysia.

### ABSTRACT

Melioidosis is caused by the gram-negative bacterium *Burkholderia pseudomallei* and has a wide range of manifestations in many organ systems. Ocular melioidosis is a rare manifestation of *Burkholderia pseudomallei* that commonly presents with orbital cellulitis rather than dacryoadenitis and multiple unilateral ocular abscesses, which are uncommon. The risk of blindness in ocular melioidosis is high because this bacterium is resistant to many antibiotics, requires a longer period of treatment with antibiotics and the diagnosis is not always straightforward. Here we describe a different presentation of ocular melioidosis manifested with left preseptal abscess, orbital cellulitis with abscess and dacryoadenitis in a 50-year-old man who was treated successfully with surgery and antibiotics, along with a brief review of the literature.

**Keywords:** Abscess, *Burkholderia pseudomallei*, Dacryoadenitis, Melioidosis, Ocular, Unilateral.

*Brunei Int Med J. 2022;18:79-83*

# Brunei International Medical Journal (BIMJ) Official Publication of The Ministry of Health and Universiti Brunei Darussalam

## EDITORIAL BOARD

<b>Editor-in-Chief</b>	Ketan PANDE
<b>Sub-Editors</b>	Vui Heng CHONG William Chee Fui CHONG
<b>Editorial Board Members</b>	Muhd Syafiq ABDULLAH Alice Moi Ling YONG Ahmad Yazid ABDUL WAHAB Jackson Chee Seng TAN Pemasiri Upali TELISINGHE Pengiran Khairol Asmee PENGIRAN SABTU Dayangku Siti Nur Ashikin PENGIRAN TENGAH

## INTERNATIONAL EDITORIAL BOARD MEMBERS

Lawrence HO Khok Yu (Singapore)	Chuen Neng LEE (Singapore)
Wilfred PEH (Singapore)	Emily Felicia Jan Ee SHEN (Singapore)
Surinderpal S BIRRING (United Kingdom)	Leslie GOH (United Kingdom)
John YAP (United Kingdom)	Ian BICKLE (United Kingdom)
Nazar LUQMAN (Australia)	Christopher HAYWARD (Australia)
Jose F LAPENA (Philippines)	

### Advisor

Wilfred PEH (Singapore)

### Past Editors-in-Chief

Nagamuttu RAVINDRANATHAN  
Kenneth Yuh Yen KOK  
Chong Vui Heng  
William Chong Chee Fui

### Proof reader

John WOLSTENHOLME (CfBT Brunei Darussalam)

# Aim and Scope of Brunei International Medical Journal

The Brunei International Medical Journal (BIMJ) is a six monthly peer reviewed official publication of the Ministry of Health under the auspices of the Clinical Research Unit, Ministry of Health, Brunei Darussalam.

The BIMJ publishes articles ranging from original research papers, review articles, medical practice papers, special reports, audits, case reports, images of interest, education and technical/innovation papers, editorials, commentaries and letters to the Editor. Topics of interest include all subjects that relate to clinical practice and research in all branches of medicine, basic and clinical including topics related to allied health care fields. The BIMJ welcomes manuscripts from contributors, but usually solicits reviews articles and special reports. Proposals for review papers can be sent to the Managing Editor directly. Please refer to the contact information of the Editorial Office.

## Instruction to authors

### Manuscript submissions

All manuscripts should be sent to the Managing Editor, BIMJ, Ministry of Health, Brunei Darussalam; e-mail: editor-in-chief@bimjonline.com. Subsequent correspondence between the BIMJ and authors will, as far as possible via should be conducted via email quoting the reference number.

### Conditions

Submission of an article for consideration for publication implies the transfer of the copyright from the authors to the BIMJ upon acceptance. The final decision of acceptance rests with the Editor-in-Chief. All accepted papers become the permanent property of the BIMJ and may not be published elsewhere without written permission from the BIMJ.

### Ethics

Ethical considerations will be taken into account in the assessment of papers that have experimental investigations of human or animal subjects. Authors should state clearly in the Materials and Methods section of the manuscript that institutional review board has approved the project. Those investigators without such review boards should ensure that the principles outlined in the Declaration of Helsinki have been followed.

## Manuscript categories

### Original articles

These include controlled trials, interventional studies, studies of screening and diagnostic tests, outcome studies, cost-effectiveness analyses, and large-scale epidemiological studies. Manuscript should include the following; introduction, materials and methods, results and conclusion. The objective should be stated clearly in the introduction. The text should not exceed 2500 words and references not more than 30.

### Review articles

These are, in general, invited papers, but unsolicited reviews, if of good quality, may be considered. Reviews are systematic critical assessments of

literature and data sources pertaining to clinical topics, emphasising factors such as cause, diagnosis, prognosis, therapy, or prevention. Reviews should be made relevant to our local setting and preferably supported by local data. The text should not exceed 3000 words and references not more than 40.

### Special Reports

This section usually consist of invited reports that have significant impact on healthcare practice and usually cover disease outbreaks, management guidelines or policy statement paper.

### Audits

Audits of relevant topics generally follow the same format as original article and the text should not exceed 1,500 words and references not more than 20.

### Case reports

Case reports should highlight interesting rare cases or provide good learning points. The text should not exceed 1000 words; the number of tables, figures, or both should not be more than two, and references should not be more than 15.

### Education section

This section includes papers (i.e. how to interpret ECG or chest radiography) with particular aim of broadening knowledge or serve as revision materials. Papers will usually be invited but well written paper on relevant topics may be accepted. The text should not exceed 1500 words and should include not more than 15 figures illustration and references

three relevant references should be included. Only images of high quality (at least 300dpi) will be acceptable.

### **Technical innovations**

This section include papers looking at novel or new techniques that have been developed or introduced to the local setting. The text should not exceed 1000 words and should include not more than 10 figures illustration and references should not be more than 10.

### **Letters to the Editor**

Letters discussing a recent article published in the BIMJ are welcome and should be sent to the Editorial Office by e-mail. The text should not exceed 250 words; have no more than one figure or table, and five references.

### **Criteria for manuscripts**

Manuscripts submitted to the BIMJ should meet the following criteria: the content is original; the writing is clear; the study methods are appropriate; the data are valid; the conclusions are reasonable and supported by the data; the information is important; and the topic has general medical interest. Manuscripts will be accepted only if both their contents and style meet the standards required by the BIMJ.

### **Authorship information**

Designate one corresponding author and provide a complete address, telephone and fax numbers, and e-mail address. The number of authors of each paper should not be more than twelve; a greater number requires justification. Authors may add a publishable footnote explaining order of authorship.

### **Group authorship**

If authorship is attributed to a group (either solely or in addition to one or more individual authors), all members of the group must meet the full criteria and requirements for authorship described in the following paragraphs. One or more authors may take responsibility 'for' a group, in which case the other group members are not authors, but may be listed in an acknowledgement.

### **Authorship requirement**

### **DISCLAIMER**

All articles published, including editorials and letters, represent the opinion of the contributors and do not reflect the official view or policy of the Clinical Research Unit, the Ministry of Health or the institutions with which the contributors are affiliated to unless this is clearly stated. The appearance of advertisement does not necessarily constitute endorsement by the Clinical Research Unit or Ministry of Health, Brunei Darussalam. Furthermore, the publisher cannot accept responsibility for the correctness or accuracy of the advertisers' text and/or claim or any opinion expressed.

sign, and the analysis and interpretation of the data (where applicable); to have made substantial contributions to the writing or revision of the manuscript; and to have reviewed the final version of the submitted manuscript and approved it for publication. Authors will be asked to certify that their contribution represents valid work and that neither the manuscript nor one with substantially similar content under their authorship has been published or is being considered for publication elsewhere, except as described in an attachment. If requested, authors shall provide the data on which the manuscript is based for examination by the editors or their assignees.

### **Financial disclosure or conflict of interest**

Any affiliation with or involvement in any organisation or entity with a direct financial interest in the subject matter or materials discussed in the manuscript should be disclosed in an attachment. Any financial or material support should be identified in the manuscript.

### **Copyright transfer**

In consideration of the action of the BIMJ in reviewing and editing a submission, the author/s will transfer, assign, or otherwise convey all copyright ownership to the Clinical Research Unit, RIPAS Hospital, Ministry of Health in the event that such work is published by the BIMJ.

### **Acknowledgements**

Only persons who have made substantial contributions but who do not fulfill the authorship criteria should be acknowledged.

### **Accepted manuscripts**

Authors will be informed of acceptances and accepted manuscripts will be sent for copyediting. During copyediting, there may be some changes made to accommodate the style of journal format. Attempts will be made to ensure that the overall meaning of the texts are not altered. Authors will be informed by email of the estimated time of publication. Authors may be requested to provide raw data, especially those presented in graph such as bar charts or figures so that presentations can be constructed following the format and style of the journal. Proofs will be sent to authors to check for any mistakes made

# DACRYOADENITIS WITH MULTIPLE UNILATERAL OCULAR ABSCESSSES IN DISSEMINATED MELIOIDOSIS.

SYLVES P<sup>1</sup>, SEAH RB<sup>2</sup>, CAROLINE B<sup>2</sup>, YAP JY<sup>1</sup>, SHUAIBAH AG<sup>1</sup>, HANIDA H<sup>3</sup>.

<sup>1</sup>Department of Ophthalmology, Faculty of Medicine and Health Sciences Universiti Malaysia Sabah, Jalan UMS, 88400 Kota Kinabalu, Sabah, Malaysia.

<sup>2</sup>Department of Ophthalmology, Queen Elizabeth Hospital, 88586 Kota Kinabalu, Sabah, Ministry of Health Malaysia.

<sup>3</sup>Oculoplastic and Reconstructive Unit, Department of Ophthalmology, Queen Elizabeth Hospital, 88586 Kota Kinabalu, Sabah, Ministry of Health Malaysia.

## ABSTRACT

Melioidosis is caused by the gram-negative bacterium *Burkholderia pseudomallei* and has a wide range of manifestations in many organ systems. Ocular melioidosis is a rare manifestation of *Burkholderia pseudomallei* that commonly presents with orbital cellulitis rather than dacryoadenitis and multiple unilateral ocular abscesses, which are uncommon. The risk of blindness in ocular melioidosis is high because this bacterium is resistant to many antibiotics, requires a longer period of treatment with antibiotics and the diagnosis is not always straightforward. Here we describe a different presentation of ocular melioidosis manifested with left preseptal abscess, orbital cellulitis with abscess and dacryoadenitis in a 50-year-old man who was treated successfully with surgery and antibiotics, along with a brief review of the literature.

**Keywords:** Abscess, *Burkholderia pseudomallei*, Dacryoadenitis, Melioidosis, Ocular, Unilateral.

## INTRODUCTION

Melioidosis is caused by a gram-negative bacterium called *Burkholderia pseudomallei*, which is endemic in Southeast Asia and Northern Australia.<sup>1,2</sup> It has a wide range of manifestations involving many organs, such as the lungs, genitourinary system, skin, liver, pancreas, spleen, brain and joints, as well as

bone and ocular structures.<sup>2-4</sup> Ocular manifestation is rare, with prevalence estimated as 0.49–1.02%.<sup>5</sup> Even though ocular melioidosis is rare, the risk of blindness due to this infection is high and can lead to life-threatening conditions if not treated early because the bacterium is resistant to many antibiotics. Here we describe a different presentation of a case of disseminated melioidosis with ocular melioidosis manifested with left preseptal abscess, orbital cellulitis with abscess and dacryoadenitis in a patient who was treated successfully with surgery and antibiotics, along with a brief review of the literature.

**Corresponding author:** Dr. Sylvès Patrick, Senior Lecturer and ophthalmologist, Department of Ophthalmology, Faculty of Medicine and Health Sciences, Universiti Malaysia Sabah (UMS), 88400, Kota Kinabalu, Sabah, Malaysia  
Tel: +6010-5093168; E-mail: [sylves@ums.edu.my](mailto:sylves@ums.edu.my)

## CASE REPORT

A 50-year-old male farmer with underlying uncontrolled diabetes mellitus, hypertension, dyslipidaemia, end-stage renal failure, a history of old ischemic stroke and both eyes treated for proliferative diabetic retinopathy presented with three weeks of left upper eyelid swelling. The swelling was associated with left eye pain, redness and reduced vision. He also had lethargy and fever, with left knee pain and swelling. He had no double vision, upper respiratory symptoms, ear discharge, toothache or symptoms of increased intracranial pressure. He also had no history of trauma or insect bites and had not taken any immunosuppressive medication.

On examination, his Glasgow Coma Scale (GCS) was 15/15. On examination of the left eye, visual acuity (VA) was 6/24 (unaided) and intraocular pressure (IOP) was 55 mmHg. There was generalised erythematous upper eyelid swelling that was non-fluctuant and tender. The eyelid was fully ptotic (Figure 1a). There was hypoglobus with limited extraocular muscle movement in all gazes. The conjunctiva was inflamed and chemosed. However, there were no signs of optic nerve dysfunction. Anterior segment examination of the right eye was unremarkable. Both eyes' posterior segments showed multiple old laser marks of panretinal photocoagulation. Other systemic examinations were unremarkable except for the left scalp and knee joint redness and swelling.

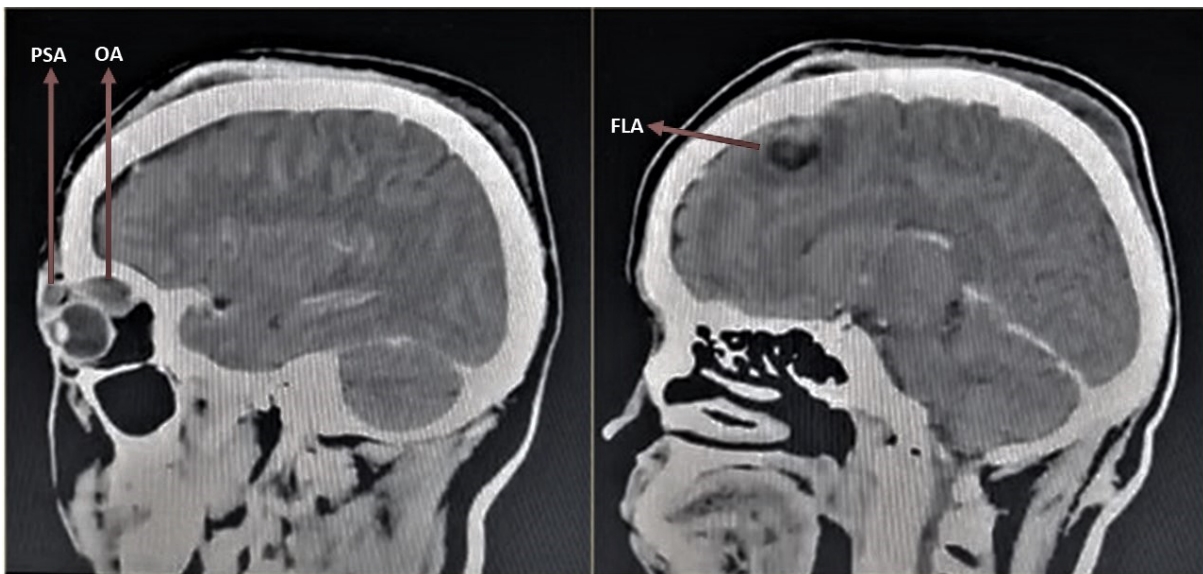
Blood investigations showed sign of infection with elevated liver enzymes

(transaminases). Viral hepatitis, retroviral and syphilis screenings were negative. Blood culture grew *Burkholderia pseudomallei*. Contrast-enhanced computed tomography (CECT) of the brain, orbit and paranasal sinuses showed left preseptal abscess, left orbital abscess, left frontal lobe abscess, left scalp abscess, left infratemporal abscess and left lacrimal gland enlargement with small well-defined hypodensities seen at bilateral internal capsules and left external capsule (Figure 2). Ultrasonography of the abdomen revealed no evidence of intra-abdominal abscess. Left knee ultrasonography showed heterogeneous hypoechoic synovial fluid.

Initially, the patient was treated with intravenous metronidazole and augmentin but the condition did not improve. Blood culture grew *Burkholderia pseudomallei* after 48 hrs and he was diagnosed with disseminated melioidosis with transaminitis. His antibiotics were immediately changed to intravenous ceftazidime (2 g od) and meropenem (2 g bd). He was co-managed by multi-disciplinary teams consisting of internal medicine, neurosurgical, otorhinolaryngology (ORL), orthopaedic and ophthalmology teams. Bedside aspiration of the scalp abscess was performed by the neurosurgical team but the left frontal lobe and left infratemporal abscesses were treated conservatively by the neurosurgical and ORL teams, respectively. The left knee underwent an arthrotomy washout performed by the orthopaedic team. He underwent incision and drainage of the preseptal and orbital abscesses via sub-brow anterior orbitotomy (Figure 1b) performed by the ophthalmology



**Figure 1: Dacryoadenitis and ocular swelling at a: Initial assessment; b: Postoperative Day 3; c: After 6 months of follow-up. (Click to enlarge)**



**Figure 2: Contrast-enhanced computed tomography showing left preseptal (PSA), orbital (OA) and frontal lobe (FLA) abscesses. The orbital abscess displaced the left eyeball downwards.**

gy team. Pus from all the sites above (scalp, knee and orbital) grew *Burkholderia pseudomallei*. He was also treated with moxifloxacin eyedrops every 4 h and with IOP-lowering eyedrops.

For diabetic controlled throughout the admission, he was treated with subcutaneous insulin (Mixtard, 14/10 units bd) and his other prescribed medications included amlodipine, 10 mg od; metoprolol, 25 mg bd; aspirin, 150 mg od; pantoprazole, 40 mg od; and hematinics. The intravenous ceftazidime was continued for 8 weeks and the meropenem for 3 weeks, along with trimethoprim/sulfamethoxazole (80 mg/400 mg, three tablets a day) for 6 months.

After completion of treatment, the patient's liver function test normalised. Repeated CECT of the brain and orbit/paranasal sinuses showed resolved left preseptal, orbital, frontal lobe, scalp and infratemporal abscesses, with normal left lacrimal gland. His left eye swelling had resolved completely with no more ophthalmoplegia and with a best-corrected VA of 6/6 (Figure 1c).

## DISCUSSION

The risk factors for ocular melioidosis are being male, a farmer, having diabetes mellitus and having renal disease.<sup>5</sup> All these risk factors were present in our case, in addition to the fact that the case occurred in an endemic region in Sabah, Malaysia. *Burkholderia pseudomallei* can be found in soil and surface water.<sup>6</sup> The patient may have acquired the infection via direct contact with contaminated soil or surface waters, especially through skin abrasions that occur during farming.<sup>3</sup>

Ocular melioidosis can be localised or part of disseminated melioidosis as was the case represented here. Disseminated melioidosis with ocular melioidosis has been reported in about 56% of cases.<sup>5</sup> Ocular melioidosis can be manifested as preseptal cellulitis, eyelid abscess, orbital cellulitis, orbital abscess, corneal ulcer, endophthalmitis, panophthalmitis, dacryocystitis, panuveitis, subretinal abscess and subconjunctival abscess.<sup>5,7</sup> However, the most common manifestation is orbital cellulitis, followed by preseptal cellulitis, endophthalmitis, panophthalmitis and panuveitis.<sup>5</sup>

Unilateral multiple ocular abscesses

are uncommon in ocular melioidosis. A case of bilateral multiple ocular abscesses was reported, manifested with bilateral orbital and eyelid abscesses that were successfully treated with surgical drainage and antibiotics. However, this was in a child with presumed ocular melioidosis.<sup>8</sup> Yaisawang et al., reported a case of culture-confirmed disseminated ocular melioidosis manifested with unilateral multiple orbital abscesses and necrotising fasciitis, with good outcome.<sup>5</sup> However, surgical drainage of the abscesses was not needed and the final visual outcome was not recorded.<sup>5</sup>

Unilateral multiple subconjunctival abscesses can occur in ocular melioidosis but such manifestation is also likely in other bacterial infections.<sup>7</sup> In our case, the abscesses was present in two different compartments (preseptal and orbital) of the eye were significant enough to need surgical drainage, which is unlikely to be the case in other bacterial infections. Knowledge of this manifestation will help ophthalmologists to prepare for melioidosis in those cases where there is no response to the standard antibiotic treatment.

Lacrimal gland involvement in melioidosis has been reported in the paediatric population in Sarawak, Malaysia.<sup>9</sup> However, this article needs to be interpreted cautiously because the authors described the lacrimal gland involvement as an inflamed tender swelling at the medial aspect of the lower eyelid, which refers more to dacryocystitis than dacryoadenitis. In our case, the left lacrimal gland was enlarged, as seen from the CECT, and this normalised after completion of treatment, suggesting lacrimal gland involvement in disseminated melioidosis.

A retrospective study conducted by Yaisawang et al. reported that about 73% of ocular melioidosis patients presented with significant visual impairment or blindness.<sup>5</sup> Even with adequate surgical intervention,

64% and 14% ended up legally blind and with enucleation, respectively.<sup>5</sup> This indicates that aggressive intervention is crucial in preserving the vision and saving the eye without undue delay. In our case, the plan for early surgical intervention was very important in relieving the high IOP, which can lead to blindness from irreversible glaucomatous optic neuropathy. After treatment with IOP-lowering agents, prolonged systemic antibiotics and surgical intervention, the visual and cosmetic outcomes were good.

## CONCLUSION

In conclusion, ocular melioidosis with multiple unilateral abscesses and dacryoadenitis is uncommon and can present as part of disseminated melioidosis, which carries a high risk of blindness and can even lead to mortality. The diagnosis is not always straightforward, which delays treatment and further worsens the prognosis. However, it is curable, with good outcomes following the appropriate surgical intervention and antibiotic treatment. Ophthalmologists need to be aware of the different ocular manifestations so that the appropriate treatment can be commenced without undue delay.

## CONFLICT OF INTEREST

The author reported no conflict of interest or financial liability.

## INFORMED CONSENT

Informed consent has been obtained from the patient in regards to the pictures and details included in this report.



## REFERENCES

- 1: Aldhous P. [Meloidosis? Never heard of it...](#) *Nature.* 2005;434:692–693. [Accessed on 2022 April 15].
  - 2: Pande K, Abd Kadir KA, Asli R, Chong VH. [Meloidosis in Brunei Darussalam.](#) *Trop Med Infect Dis.* 2018; 3(1):20. [Accessed on 2022 April 15].
  - 3: Cheng AC, Currie BJ. [Meloidosis: Epidemiology, Pathophysiology, and Management.](#) *Clinical Microbiology Reviews.* 2005:383-416. [Accessed on 2022 April 15].
  - 4: Chong VH, Lim KS, Sharif F. [Pancreatic Involvement in Meloidosis.](#) *JOP. J Pancreas (Online).* 2010; 11(4):365-368. [Accessed on 2022 April 15].
  - 5: Yaisawang S, Asawaphureekorn S, Chetchotisakd P, Wongratanacheewin S, Pakdee P. [Ocular involvement in melioidosis: a 23-year retrospective review.](#) *Journal of ophthalmic inflammation and infection.* 2018;8(1):1-9. [Accessed on 2022 April 15].
  - 6: Strauss J, Groves M, Mariappan M, Ellison D. *Melioidosis in Malaysia. The American journal of tropical medicine and hygiene.* 1969;18(5):698-702.
  - 7: Siripanthong S, Teerapantuwat S, Prugsanusak W, et al. *Corneal ulcer caused by Pseudomonas pseudomallei: report of three cases. Reviews of Infectious Diseases.* 1991;13(2):335-337.
  - 8: Kogilavaani J, Shatriah I, Regunath K. [Bilateral orbital abscesses with subdural empyema and cavernous sinus thrombosis due to melioidosis in a child.](#) *Asian Pacific Journal of Tropical Disease.* 2014;4:S851-S853. [Accessed on 2022 April 15].
  - 9: Mohan A, Podin Y, Tai N, et al. [Pediatric melioidosis in Sarawak, Malaysia: Epidemiological, clinical and microbiological characteristics.](#) *PLoS Neglected Tropical Diseases.* 2017;11(6):e0005650. [Accessed on 2022 April 15].
-