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Clinical characteristics and outcomes of paediatric orbital cellulitis in Hospital Universiti Sains Malaysia: a five-year review

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ABSTRACT

Introduction: Limited data is available on paediatric orbital cellulitis in Asia. We aimed to describe demographic data, clinical presentation, predisposing factors, identified micro-organisms, choice of antibiotics and management in children with orbital cellulitis treated in a tertiary care centre in Malaysia.

Methods: A retrospective review was performed on children with orbital cellulitis aged below 18 years who were admitted to Hospital Universiti Sains Malaysia, Kelantan, Malaysia, between January 2013 and December 2017.

Results: A total of 14 paediatric patients fulfilling the diagnostic criteria for orbital cellulitis were included. Their mean age was 6.5 ± 1.2 years. Boys were more likely to have orbital cellulitis than girls (71.4% vs. 28.6%). Involvement of both eyes was observed in 14.3% of the patients. Sinusitis (28.6%) and upper respiratory tract infection (21.4%) were the most common predisposing causes. *Staphylococcus aureus* (21.4%) was the leading pathogen. Longer duration of hospitalisation was observed in those infected with Methicillin-resistant *Staphylococcus aureus* and *Burkholderia pseudomallei*. 10 (71.4%) patients were treated with a combination of two or three antibiotics. In this series, 42.9% had surgical interventions.

Conclusion: Young boys were found to be more commonly affected by orbital cellulitis than young girls. *Staphylococcus aureus* was the most common isolated micro-organism. Methicillin-resistant *Staphylococcus aureus* and *Burkholderia pseudomallei* caused severe infection. Sinusitis and upper respiratory tract infection were the most common predisposing factors. A majority of the children improved with medical treatment alone. Our findings are in slight disagreement with other published reports on paediatric orbital cellulitis, especially from the Asian region.

Keywords: Asian children, orbital abscess, orbital cellulitis

INTRODUCTION

Orbital cellulitis in children is a serious infection that can result in significant complications including blindness, cavernous sinus thrombosis, meningitis, subdural empyema and brain abscess.⁽¹⁻³⁾ Prompt diagnosis and early treatment are important to prevent potential vision or life-threatening conditions.

Numerous reports on paediatric orbital cellulitis have been published in the literature.⁽⁴⁻¹²⁾ Despite the abundance of reports on the topic, among Asian countries, data was only available from Singapore, India and Taiwan.⁽¹²⁻¹⁵⁾ These series included data on both adults and children.⁽¹³⁻¹⁵⁾ In this study, we described demographics, clinical characteristics, causative organisms, treatment and visual outcomes among paediatric orbital cellulitis cases treated in Malaysia, and reviewed the relevant literature.

METHODS

A retrospective case series was conducted on all children aged under 18 years who were diagnosed with orbital cellulitis and admitted to Hospital Universiti Sains Malaysia, Kelantan, Malaysia. Data was collected from 1 January 2013 till 31 December 2017. The study was conducted in accordance with the Declaration of Helsinki.

Cases were identified through a database search of medical records using the International Classification of Diseases Ninth and Tenth Revision codes (376.01, H05.0) for orbital cellulitis. A total of 44 paediatric patients with the diagnosis of orbital cellulitis were identified on the admission computers, and their medical records were retrieved. Children who were diagnosed with orbital cellulitis and admitted to the ward were included in our study. We excluded those who had pre-existing ocular diseases such as corneal opacity, congenital cataracts, strabismus, refractive errors, optic disc or macular problems. A final list was reviewed, and 14 patients were carefully selected. Data documented included age, gender, laterality,

possible identified cause of infection, presenting clinical manifestations, imaging results, microbiology results, treatment options (medical treatment alone or in combination with surgical drainage), length of hospital stay and final visual outcome. The institution required patients' paediatric orbital cellulitis registry forms to be completed by a consultant paediatric ophthalmologist. Data was analysed using SPSS Statistics version 22.0 (IBM Corp, Armonk, NY, USA).

RESULTS

A total of 14 paediatric patients with orbital cellulitis in 16 eyes were recruited according to the study protocol. Table I summarises the clinical profile of all patients involved in this study. The mean age of the patients was 6.5 ± 1.2 years. The majority were male (10 eyes, 71.4%). 7 (50.0%) patients had left eye involvement and 2 (14.3%) patients had bilateral involvement.

Lid swelling and conjunctival chemosis were the most common presenting features (100.0% and 78.6%, respectively). 5 (35.7%) patients had presenting visual acuity of 6/12 (20/40) and worse. 11 (78.6%) patients had final visual acuity of 6/9 (20/32) or better. The final visual acuity of three young patients aged 39 days, one year and two years was not documented, as they were uncooperative during the examination. The mean duration of hospitalisation was 13.2 ± 11.9 days (Table II).

6 (42.9%) patients had been treated with oral antibiotics prior to referral to the centre. The most commonly prescribed antibiotics were Augmentin and ceftriaxone. Other antibiotics included metronidazole, cloxacillin, ceftazidime and/or vancomycin. The majority of the patients (71.4%, $n = 10$) received intravenous antibiotics for seven days or less, with 2 (14.3%) patients receiving antibiotics for six weeks due to intracranial complications.

The most commonly isolated organisms were *Staphylococcus* species in 4 (28.6%) patients and *Streptococcus* species in 3 (21.5%) patients. Blood cultures were negative for all

patients, except for one patient aged 39 days with methicillin-resistant *Staphylococcus aureus*. All patients were promptly treated with intravenous antibiotics upon admission. 8 (57.1%) patients were successfully treated with medical treatment alone, while 6 (42.9%) patients were treated with a combination of medical treatment and surgical interventions.

DISCUSSION

There have been limited reviews of paediatric orbital cellulitis in Asian countries.⁽¹²⁻¹⁵⁾ Yang et al described a report involving 20 children with orbital cellulitis treated in Singapore,⁽¹²⁾ while Liu et al, Babu et al and Pandian et al reported data from Taiwan^(13,14) and India,⁽¹⁵⁾ respectively, but their inclusion criteria were slightly different from those in this study. Table II summarises published reports on paediatric orbital cellulitis from Asian countries, Australia, the Middle East, United Kingdom, Scotland, Canada and the United States, including data from the present study.^(4-7,9,12,13,15)

The mean age of presentation of our patients was 6.5 ± 1.2 years, which was consistent with other published reports.^(5-7,12) 71.4% of our patients were boys, a similar observation to the reports of Ferguson et al and Nageswaran et al.^(4,5) In contrast, a Scotland study of 243 children found that slightly more girls than boys had orbital cellulitis.⁽⁶⁾

Left eye involvement was observed in 50% of our patients. A higher percentage of left eye involvement was also documented by Nageswaran et al (56.1%),⁽⁵⁾ Fanella et al (65.8%),⁽⁷⁾ and Atfeh and Khalil (61.0%).⁽⁹⁾ In our study, both eyes were involved in 2 (14.3%) patients. One of the two patients had an abscess in the medial canthal area, while the second developed severe septicaemia. Bilateral eye involvement was also reported by Nageswaran et al (34.1%)⁽⁵⁾ as well as Atfeh and Khalil (4.0%).⁽⁹⁾

Sinusitis and upper respiratory tract infection were the most common predisposing factors, occurring in 28.6% and 21.4% of our patients, respectively. These findings were

consistent with Yang et al's report of a 30.0% incidence of sinusitis and 25.0% incidence of upper respiratory infection in their review involving 20 Singaporean children diagnosed with orbital cellulitis.⁽¹²⁾ In contrast, a very high incidence of sinusitis was reported among Australian and Canadian children (91.2% and 94.7%, respectively).^(4,7) Trauma was documented in 14.3% of the cases in this study. On the contrary, Ferguson et al and Yang et al reported no trauma in their studies.^(4,12)

Final visual acuity of 6/9 (20/32) or better was documented in 78.6% of our patients. The remaining patients did not cooperate for the assessment of visual acuity during our assessment. A good final visual outcome of 90.0% was reported by Yang et al.⁽¹²⁾ Our patients stayed longer in the hospital (mean 13.2 ± 11.9 days) compared to other published data.^(4,5-7,9) This was mainly due to difficult access to our hospital, poor clinical response to the initial antibiotic treatment and the necessity for a longer course of antibiotics. Duration of antibiotic course in our patients was fairly similar to that in the report by Nageswaran et al.⁽⁵⁾

Staphylococcus aureus (21.4%) was the most common pathogen in our series. This observation was in keeping with studies conducted in Australia (14.7%) and Singapore (25.0%).^(4,12) Both Fanella et al and Atfeh and Khalil discovered that *Staphylococcus aureus* and non-haemolytic streptococci were the leading pathogen in their series.^(7,9) We also encountered cases infected with methicillin-resistant *Staphylococcus aureus* and *Burkholderia pseudomallei*.^(16,17) Both patients exhibited severe manifestations of infection, leading to a longer duration of antibiotics and length of hospitalisation.^(16,17) Infection due to *Haemophilus influenzae* was documented by Nageswaran et al (7.3%) and Crosbie et al (6.7%).^(5,6) Other infections were caused by Group C β -haemolytic streptococci, *Eikenella corrodens*, *Arcanobacterium haemolyticum* and *Moraxella catarrhalis*.⁽⁵⁾

Melioidosis is an uncommon cause for orbital cellulitis in children. Our patient in this study presented with bilateral orbital abscesses, subdural empyema and cavernous sinus

thrombosis.⁽¹⁷⁾ A clinical assumption of melioidosis was made in this patient based on poor clinical response to conventional antibiotic therapy, borderline indirect haemagglutination assay titre and a favourable response after commencing ceftazidime treatment. It is pertinent to have a high index of suspicion for melioidosis when dealing with such patients who show poor response to conventional treatment in children, especially those living in endemic areas.

Broad choices for antibiotics were commenced based on clinical and laboratory results. Combinations of three antibiotics including Augmentin, metronidazole, ceftazidime, cefuroxime and vancomycin were prescribed for 28.6% of our patients. Dual combinations including Augmentin-cloxacillin, ceftriaxone-Augmentin, ceftriaxone-cloxacillin, ceftriaxone-metronidazole and ceftriaxone-ceftazidime were also started in our patients. An ampicillin-cloxacillin combination was used to treat 15.0% of patients in the study described by Yang et al.⁽¹²⁾ Patients treated with clindamycin-cephalosporin and cloxacillin-cefotaxime combinations represented 42.2% of the population in the study by Fanella et al.⁽⁷⁾ Nageswaran et al described that 46.3% of their patients were prescribed with an ampicillin-sulbactam combination, while another 26.8% was treated with nafcillin-ceftizoxime.⁽⁵⁾ The broad selection of antibiotics in the above studies was consistent with observations by Lee and Yen, who suggested that different communities have distinctive flora and varying resistance profiles.⁽¹⁸⁾

Surgical intervention was considered in 42.9% of our patients, while the others improved with medical treatment alone. Fanella et al reported that 21.1% of their patients underwent surgical interventions,⁽⁷⁾ which was inconsistent with the higher percentages of surgical interventions observed in Singapore (65%), Australia (76.4%), Scotland (83.3%) and the United States (70.7%). Indications for surgical intervention included larger abscesses with mass effect, poor clinical response to initial medical treatment and concurrent intracranial involvement.⁽¹²⁾ This phenomenon also suggests that ophthalmologists have different threshold levels for surgical interventions in managing paediatric orbital cellulitis. However, the information gathered from

this study is limited to that from a single institution. Multicentre involvement would be beneficial to reflect a broader spectrum of the illness.

In conclusion, paediatric orbital cellulitis was more likely to affect young boys in our series. Methicillin-resistant *Staphylococcus aureus* and *Burkholderia pseudomallei* caused severe infection, while sinusitis and upper respiratory tract infection were the most common predisposing factors. A majority of the population improved with medical treatment alone. The findings from this study were slightly different from other published data on paediatric orbital cellulitis.

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Table I. Summary of clinical cases (n = 14).

No.	Age; gender	Clinical presentation; onset; laterality	CT result	Cause	Culture; investigation	Antibiotic (duration)	Surgery	Hospital LOS	Presenting VA; final VA
1	6 yr; M	Lid swelling, ptosis, high-grade fever; 3 days; right eye	Extraconal abscess (subperiosteal abscess)	Upper respiratory tract infection and sinusitis	No growth from eye swab, blood and urine samples; TWC 10.6, PLT 383, ESR 96, CRP NA	IV ceftriaxone (1 wk), IV Augmentin (5 days)	Nil	12 days	6/6; 6/6
2	5 yr; M	Lid swelling, erythema, ptosis, high-grade fever, intracranial abscess; 7 days; left eye	Lid and cerebral abscess	Trauma – hit by toys, underlying sinusitis	No growth from pus, blood and urine samples; TWC 14.7, PLT 444, ESR 106, CRP positive	IV ceftriaxone (7 wk), IV metronidazole (6 wk)	Open lid incision and drainage	7 wk	Not cooperative; 6/6
3	2 yr; M	Lid swelling, ptosis, chemosis, high-grade fever; 2 days; both eyes	Medial canthus abscess	Upper respiratory tract infection and exudative tonsillopharyngitis	<i>Streptococcus Pneumoniae</i> (pus); TWC 19.6, PLT 271, ESR NA, CRP positive	IV Augmentin (5 days), IV cloxacillin (5 days)	Nil	1 wk	NA; NA
4	6 yr; M	Lid swelling, ptosis, chemosis; 1 day; right eye	Lid abscess	No identified cause	No growth from eye swab, blood and urine samples; TWC NA, ESR NA, CRP NA	IV Augmentin (5 days)	Nil	5 days	6/9; 6/6
5	14 yr; M	Lid swelling, ptosis, chemosis, low-grade fever; 1 day; right eye	Periorbital tissue thickening extending to right temporal region	Trauma and laceration wound at the temporal eyelid	<i>Streptococcus beta-haemolytic group A</i> ; <i>Staphylococcus aureus</i> (pus); TWC 32.6, PLT 150, ESR NA, CRP NA	IV ceftriaxone (2 wk), IV cloxacillin (2 wk)	Open lid incision and drainage	2 wk	6/6; 6/6
6	13 yr; M	Lid swelling, chemosis, low-grade fever; 5 days; right eye	Not done	Upper respiratory tract infection	No growth from eye swab, blood and urine samples; TWC 7.6, PLT 206, ESR NA, CRP NA	IV Augmentin (5 days)	Nil	6 days	6/12; 6/6
7	39 days; M	Facial swelling, proptosis, chemosis, high-grade fever; 1 day;	Periorbital and facial abscess	Gingivostomatitis	Methicillin-resistant <i>Staphylococcus aureus</i> (from eye swab and blood sample); TWC 13.4, PLT	IV ceftazidime (14 days), IV metronidazole (14 days), IV	Image-guided endoscopic drainage	23 days	NA; NA

		right eye			511, ESR NA, CRP positive	vancomycin (10 days)			
8	10 yr; F	Lid swelling, chemosis, high-grade fever; 3 days; left eye	Soft tissue thickening and lacrimal sac abscess	Dacryocystitis	No growth from eye swab, blood and urine samples; TWC 12.01, PLT 270, ESR NA, CRP NA	IV Augmentin (5 days), IV ceftazidime (1 wk), IV metronidazole (1 wk)	Nil	11 days	6/6; 6/6
9	6 yr; M	Lid swelling, chemosis, ophthalmoplegia, low-grade fever; 1 day; left eye	Lid abscess and orbital cellulitis	Pansinusitis	No growth from eye swab, blood and urine samples; TWC 29.7, PLT 896, ESR NA, CRP NA	IV Augmentin (5 days), IV ceftazidime (5 days), IV metronidazole (5 days)	Nasal decongestion and left antral washout	8 days	Not cooperative; 6/9
10	11 yr; F	Lid swelling, chemosis, no fever; 7 days; left eye	Not done	Infected wound from a ruptured upper lid abscess	<i>Staphylococcus aureus</i> (pus); TWC 9.9, PLT 263, ESR NA, CRP NA	IV Augmentin (5 days)	Nil	7 days	6/21; 6/6
11	4 yr; M	Lid swelling, chemosis, low-grade fever; 4 days; left eye	Heterogeneous enhancement of anterior orbital tissue with fluid collection	Dacryoadenitis	No growth from eye swab, blood and urine samples; TWC 10.4, PLT 225, ESR NA, CRP NA	IV Augmentin (1 wk), IV ceftazidime (1 wk), IV metronidazole (1 wk)	Nil	16 days	6/12; 6/6
12	1 yr; M	Lid swelling, chemosis, ptosis; 3 days; left eye	Soft tissue thickening	No identified cause	No growth from eye swab, blood and urine samples; TWC 8.4, PLT 276, ESR NA, CRP negative	IV Augmentin (1 wk)	Open eyelid incision and drainage	7 days	NA; NA
13	7 yr; F	Lid swelling, chemosis, proptosis, ophthalmoplegia, diplopia; 3 days; left eye	Enhancing soft tissue thickening with intraorbital abscess	Sinusitis and dacryoadenitis	<i>Staphylococcus aureus</i> (pus); TWC 12.1, PLT 242, ESR 18, CRP NA	IV ceftriaxone (1 wk)	Nil	9 days	6/12; 6/6
14	11 yr; F	Periorbital swelling, blurred vision, proptosis,	Bilateral orbital abscess, intracranial	No identified cause	<i>Burkholderia pseudomallei</i> ; TWC NA, PLT NA, ESR NA, CRP	IV ceftazidime (6 wk), IV metronidazole	Bifrontal craniotomy and evacuation of	49 days	6/60; 6/6

		ophthalmoplegia, high-grade fever, intracranial abscess; 3 days; both eyes	abscess, subdural empyema and cavernous sinus thrombosis		NA	(6 wk)	pus within subdural space, open eyelid incision and drainage		
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CRP: C-reactive protein; ESR: erythrocyte sedimentation rate; F: female; IV: intravenous; LOS: length of stay; M: male; NA: not applicable; PLT: platelet; TWC: total white cell count; VA: visual acuity

Table II. Comparison of published studies on paediatric orbital cellulitis.

Variable	No. (%) / mean \pm standard deviation								
	Present study	Yang et al ⁽¹²⁾	Liu et al ⁽¹⁵⁾	Babu et al ⁽¹³⁾	Ferguson et al ⁽⁴⁾	Atfeh & Khalil ⁽⁹⁾	Crosbie et al ⁽⁶⁾	Nageswaran et al ⁽⁵⁾	Fanella et al ⁽⁷⁾
Country; yr	Malaysia; 2019	Singapore; 2009	Taiwan; 2006	India; 2016	Australia; 1999	United Kingdom; 2015	Scotland; 2016	United States; 2006	Canada; 2011
Population	14 children	20 children	Total 27 (8 children + adults)	Total 14 (5 children + adults)	Total 52 (34 children) + adults)	Total 54 (40 children + adults)	Total 234 (including 30 children who fulfilled Chandler classification)	41 children	38 children
Age (yr)	6.5 \pm 1.2	5.5	0–18	NA	NA	5.8 \pm 4.5	4.7 \pm 4.4	7.5	7.5
Gender									
Male	10 (71.4)	NA	Male > female	2 (40.0)	25 (73.5)	NA	(51.9)	30 (73.2)	NA
Female	4 (28.6)	NA		3 (60.0)	9 (26.5)	NA	(48.1)	11 (26.8)	NA
Laterality									
Right	5 (35.7)	NA	NA	3 (60.0)	NA	(35.0)	NA	4 (9.8)	13 (34.2)
Left	7 (50.0)	NA	NA	2 (40.0)	NA	(61.0)	NA	23 (56.1)	25 (65.8)
Both	2 (14.3)	NA	NA	0 (0)	NA	(4.0)	NA	14 (34.1)	0 (0)
Antibiotics before admission	6 (42.9)	NA	NA	NA	11 (32.4)	NA	NA	33 (80.5)	22 (57.9)
Fever on/after presentation (> 38°C)	6 (42.9)	20 (100.0)	NA	NA	9 (26.5)	NA	NA	27 (65.9)	NA
TWC > 15,000	3 (21.4)	13 (65.0)	NA	NA	NA	NA	NA	19 (46.3)	13 (34.2)
Predisposing factors									
Sinusitis	4 (28.6)	6 (30.0)	NA	4 (80.0)	31 (91.2)	NA	NA	NA	36 (94.7)

Diabetes mellitus	0 (0)	0 (0)	NA	1 (20.0)	0 (0)	NA	NA	NA	NA
Other									
<i>Dental abscess</i>	1 (7.1)	0 (0)	NA	NA	1 (2.9)	NA	NA	NA	NA
<i>Trauma</i>	2 (14.3)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
<i>Foreign body</i>	0 (0)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
<i>Dacryocystitis</i>	1 (7.1)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
<i>Dacryoadenitis</i>	2 (14.3)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
<i>Upper respiratory tract infection</i>	3 (21.4)	5 (25.0)	NA	NA	3 (8.8)	NA	NA	NA	NA
<i>Uveitis</i>	0 (0)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
<i>Tumour</i>	0 (0)	0 (0)	NA	NA	0 (0)	NA	NA	NA	NA
Ophthalmologic signs									
Lid swelling	14 (100.0)	20 (100.0)	NA	NA	NA	NA	NA	NA	NA
Reduce visual acuity	5 (35.7)	NA	NA	NA	11 (32.4)	NA	NA	NA	NA
Proptosis	4 (28.6)	8 (40.0)	NA	NA	26 (76.5)	NA	NA	25 (61.0)	NA
Chemosis	11 (78.6)	10 (50.0)	NA	NA	5 (14.7)	NA	NA	NA	NA
Ptosis	7 (50.0)	NA	NA	NA	NA	NA	NA	30 (73.2)	NA
Ophthalmoplegia	3 (21.4)	15 (75.0)	NA	NA	24 (70.6)	NA	NA	19 (46.3)	NA
None	0 (0)	NA	NA	NA	0 (0)	NA	NA	11 (26.8)	NA
Visual acuity on presentation									
6/60	1 (7.1)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/21	1 (7.1)	1 (5.0)	NA	NA	NA	NA	NA	NA	NA
6/18	0 (0)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/12	3 (21.4)	9 (45.0)	NA	NA	NA	NA	NA	NA	NA
6/9	1 (7.1)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/6	3 (21.4)	7 (35.0)	NA	NA	NA	NA	NA	NA	NA
Not cooperative	5 (35.7)	2 (10.0)	NA	NA	NA	NA	NA	NA	NA
Final visual acuity									
6/21	0 (0)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/18	0 (0)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/12	0 (0)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/9	1 (7.1)	0 (0)	NA	NA	NA	NA	NA	NA	NA
6/6	10 (71.4)	18 (90.0)	NA	3(60.0)	NA	NA	NA	NA	NA
Not done	3 (21.4)	2 (10.0)	NA	NA	NA	NA	NA	NA	NA
Length of hospitalisation (day)									
All patients	13.2 ± 11.9	NA	NA	NA	6.7	3.9 ± 3.5	NA	5.8 ± 2.9	7.0 ± 2.7

8–14	1 (7.1)	NA	NA	NA	NA	NA	NA	NA	NA
15–30	1 (7.1)	NA	NA	NA	NA	NA	NA	NA	NA
> 30	2 (14.3)	NA	NA	NA	NA	NA	NA	NA	NA
Micro-organism									
Aerobes									
α -non-haemolytic streptococci	0 (0)	NA	0 (0)	NA	0 (0)	8 (30.8)*	0 (0)	7 (17.1)	1 (2.6)
Group A β -haemolytic streptococci	2 (14.3)	NA	0 (0)	NA	3 (8.8)	NA	0 (0)	3 (7.3)	0 (0)
<i>Streptococcus pyogenes</i>	0 (0)	NA	0 (0)	NA	0 (0)	NA	9 (30.0)	0 (0)	1 (2.6)
<i>Streptococcus anginosus</i>	0 (0)	NA	0 (0)	NA	0 (0)	NA	3 (10.0)	0 (0)	0 (0)
<i>Streptococcus pneumoniae</i>	0 (0)	NA	0 (0)	NA	1 (2.9)	NA	1 (3.3)	0 (0)	0 (0)
<i>Staphylococcus aureus</i>	3 (21.4)	5 (25.0)	3 (37.5)	NA	5 (14.7)	8 (30.8)*	0 (0)	3 (7.3)	1 (2.6)
Coagulase-negative <i>Staphylococcus</i>	0 (0)	0 (0)	3 (37.5)	NA	0 (0)	NA	0 (0)	0 (0)	0 (0)
Methicillin-resistant <i>Staphylococcus aureus</i>	1 (7.1)	NA	0 (0)	NA	0 (0)	NA	0 (0)	0 (0)	0 (0)
<i>Burkholderia pseudomallei</i>	1 (7.1)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	0 (0)	0 (0)
<i>Haemophilus influenzae</i>	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	2 (6.7)	3 (7.3)	0 (0)
Group C β -haemolytic streptococci	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	2 (4.9)	0 (0)
<i>Eikenella corrodens</i>	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	2 (4.9)	0 (0)
<i>Arcanobacterium hemolyticum</i>	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	1 (2.4)	0 (0)
<i>Moraxella catarrhalis</i>	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	1 (2.4)	0 (0)
<i>Pseudomonas aeruginosa</i>	0 (0)	1 (5.0)	0 (0)	NA	0 (0)	NA	0 (0)	0 (0)	0 (0)
<i>Corinebacterium</i>	0 (0)	0 (0)	0 (0)	NA	0 (0)	NA	0 (0)	0 (0)	0 (0)
Anaerobes	0 (0)	0 (0)	0 (0)	NA	4 (11.8)	2 (7.7)*	0 (0)	8 (19.5)	0 (0)
No growth	7 (50.0)	0 (0)	NA	NA	NA	14 (53.8)*	8 (26.7)	NA	3 (7.9)
No specimen	0 (0)	NA	NA	NA	NA	NA	3 (10.0)	NA	NA

Values of n have been omitted when they were not available. *n = 26 (based on positive culture). †n = 8. NA: not available; TWC: total white cell count