Community acquired methicillin sensitive *Staphylococcus aureus* bacteremia, meningitis and brain abscess: A unique presentation

Carlos Gonzalez, Juan Roa, Nehad Shabarek

**ABSTRACT**

Introduction: *Staphylococcus aureus* causing isolate central nervous system (CNS) infection is rare. It is mostly related to either neurosurgical intervention or previous local or systemic infection rather than spontaneous isolated infection in adults. Case Report: We report a case of a previously healthy 19-year-old male with no risk factors of *Staphylococcus aureus* infection, who was found to have methicillin sensitive *Staphylococcus aureus* (MSSA) bacteremia, meningitis and cerebral abscess, without an apparent source of infection. Conclusion: Our case is a unique presentation of a young male presenting with isolated CNS infection by MSSA. This case highlights the challenge of early diagnosis of brain abscess, an entity that presents very often with non-specific signs, the diagnosis of which can be easily missed, with repercussions on long-term disability and mortality.

**Keywords:** *Staphylococcus aureus*, Bacteremia, Meningitis, Brain abscess

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**INTRODUCTION**

*Staphylococcus aureus* is the most virulent of the staphylococcal species with unique versatility to cause various types of infections, from community acquired mild skin infections to highly lethal infections such as necrotizing pneumonia, endocarditis or central nervous system (CNS) infections. CNS infections caused by *Staphylococcus aureus* are in general uncommon. Various series report *Staphylococcus aureus* as the fourth or fifth most common etiology for neuro-infection; more commonly as cause of brain abscess and less commonly as an isolated cause of meningitis (less than 5%) [1–5, 6]. In either case, CNS infection caused by *Staphylococcus aureus* is related to recent neurosurgical intervention or associated with systemic or local infection classically described as a complication of endocarditis or soft tissue infection [1–2, 6]. Although there are a few case reports of CNS infection in patients without an apparent source, most of them are related to community acquire methicillin resistant *Staphylococcus aureus* (CA-MRSA) infections [7]. To date, we have not found any reported case of confirmed meningitis with brain abscess, caused by MSSA infection in a previous healthy patient with no
predisposing infection or risk factors. We report a case of MSSA bacteremia with meningitis and brain abscess in a previous healthy young male without any risk factors and no obvious source of infection, the detection of which was delayed due to a normal brain non-contrast computed tomography (CT) scan findings.

CASE REPORT

A 19-year-old Hispanic male student, presented to the emergency department (ED) complaining of dull frontal and occipital headache associated with fever, few episodes of vomiting, poor oral intake and myalgia for two days. His past medical history was only remarkable for a recent dental procedure. He denied any sick contacts and drug abuse. Initial physical examination was remarkable for tachycardia of 123 bpm and temperature of 38°C. No meningeal signs or focal deficits were found on the initial presentation. The remaining physical examination was unremarkable. Initial laboratory work up was positive for mild hyperglycemia with blood sugar 143 mg/dL, Na 132 mEq/dL and neutrophilia 92% with WBC count 11x10^3/mm³. Brain CT scan and lumbar puncture (LP) were performed to exclude meningitis or any intracranial pathology. Both the investigations were negative. The patient received symptomatic management with IV fluids and non-steroidal anti-inflammatory drugs (NSAIDs) for fever, with notable improvement. He was discharged home from the ED with symptomatic management. One day after the initial presentation, blood cultures were reported positive for MSSA. Multiple attempts to contact the patient and his family were unfortunately unsuccessfully.

Four days after the initial presentation, the patient was brought to the ED again, after being found unresponsive. Patient’s mother stated he was doing well until early morning when he complained of malaise, fever and headache. Physical examination was positive for temperature of 40.5°C. Other vital signs were normal. The patient was lethargic, found to have nuchal rigidity and was only responsive to painful stimuli. Glasgow coma scale (GCS) was 9/15. Mild patchy folliculitis was noted in the suprapubic area with crusted lesions. He was intubated and transferred to the medical intensive care unit (MICU) with a diagnosis of possible meningitis and MSSA bacteremia based on his previous blood culture. Repeat LP and blood culture were performed. Lumbar puncture showed xanthochromia with RBC 11/mm³, WBC count 817/mm³ (neutrophils 90%, lymphocytes 2%, monocytes 8%), glucose 28 mg/dL and protein 126 mg/dL. Blood cultures grew MSSA with the same pattern of sensitivity as the first culture. Non-contrast brain CT scan was again performed, with normal findings (Figure 1). The patient was initiated on bacterial meningitis treatment with ceftriaxone 2 g q12hr, nafcillin 2 g q4hr, and dexamethasone 4 mg q6hr. Ceftriaxone was switched to gentamicin 60 mg q8hr on day-2.

During the hospital course in MICU, transthoracic echocardiogram (TTE) was negative for endocarditis, urine toxicology was negative and HIV status was negative. The patient was extubated four days after admission, at which time a left palpebral ptosis was noted. His pupils were symmetric, 3 mm in diameter, and reactive. He remained lethargic and a contrast enhanced magnetic resonance imaging (MRI) was ordered due to his lack of neurological improvement. The MRI scan of brain showed a focal lesion centered at the left anterior thalamus with contiguous extension into the subthalamic region and left cerebral peduncle of the left midbrain, with a small focus of contralateral extension into the right thalamus (left lesion measure: 3.5 cm height, 2 cm antero-posterior and 1.8 transverse, right lesion 0.4 cm diameter) (Figure 2). These findings were compatible with brain abscess.

After eight days of medical treatment and observation with the patient was sent for stereotactic biopsy and aspiration of the abscess due to no neurological improvement. The procedure was done without complication and patient started showing signs of improvement. He became more alert and followed commands. Pathology reported gliotic brain tissue with organizing necrosis and hemorrhage without purulent inflammation. Fluid culture was negative.

Ten days after treatment with nafcillin, the patient presented with diffuse erythematous rash involving the palm of the hands. Biopsy of the lesions was remarkable for interface dermatitis, compatible with a drug eruption. Nafcillin was then switched to vancomycin 1 g q12hr. Patient remained stable and was transferred to medical floor and subsequently to rehabilitation.
facilit to continue antibiotic treatment for 6–8 weeks or until resolution of the brain abscess. Patient was seen four weeks after with complete neurological recuperation and resolution of the brain abscess on repeat MRI scan.

**DISCUSSION**

*Staphylococcus aureus* infections remains a major cause of morbidity and mortality, with presentations ranging from simple cutaneous infection to serious life-threatening infections. Isolated meningitis caused by *Staphylococcus aureus* is rare accounting for less than 5% of all meningitis presentations [2]. On the other hand, *Staphylococcus aureus* is a major cause of cerebral abscess being the second most commonly found bacteria in patients with cerebral abscess, with *Streptococcus milleri* being the most common, and anaerobes the third most common [1, 3–5]. CNS infection caused by *Staphylococcus aureus* is usually associated with prior neurosurgical intervention or with typical risk factors such as endocarditis, soft tissue infection, skin abscesses, AIDS or intravenous drug use [8, 6]. In this case, the only infection that could be associated with the MSSA bacteremia was the mild suprapubic foliculitis, which we determined to be a very unlikely source as it was a mild cutaneous infection in an immunocompetent host. The patient was evaluated by oral maxillofacial surgery (OMS) department who determined that the prior dental work was an unlikely cause of the brain abscess and neurological infection that ensued.

The initial presentation of a patient with cerebral abscess could be very non-specific with only generalized symptoms. This makes the initial recognition very challenging. Different series have reported headache as the most common symptom in a patient with a brain abscess (41–63% cases). Fever is an inconsistent finding, only reported in 25% cases [3, 5]. Altered mental status (AMS) and focal neurologic deficit are inconsistent findings reported in different series with a range for AMS being 18–48% and from 25–65% for focal neurologic deficit [1, 3–5]. Other symptoms, especially gastrointestinal symptoms are present in about 27% cases [5]. In our patient, non-contrast CT scan of the head was performed and yielded a negative result on both presentations to the ED. A contrast MRI scan done several days after admission showed the thalamic abscess. The initial negative finding on CT scan and the absence of focal neurological deficit contributed to the delay in the diagnosis of brain abscess. Despite this, non-contrast CT scan is shown to be a good initial test for a patient with suspicion of cerebral abscess, even though in the early phase, non-contrast CT scan could be negative [7, 9, 10]. When it is used in combination with contrast CT scan, the sensitivity is very high, as was demonstrated in a series of 50 cases where positive CT scan finding was reported in 100% of the cases [1]. When CT scan is not diagnostic, MRI scan with contrast is in general the next step. Contrast MRI scan is highly accurate for detection of brain abscesses [11].

Treatment of brain abscess is based on pertinent antibiotic therapy and neurosurgical mechanical drainage in most cases. Recommended initial empiric antibiotic treatment is a third generation cephalosporin in combination with metronidazole. If there is suspicion or risk factors for *Staphylococcus aureus* infection, vancomycin should be added until culture identification and sensitivities are available. Vancomycin remains the drug of choice when there is a suspicion for *Staphylococcus aureus* infection despite poor penetration in the CNS [1–5, 10]. As a result, some authors have proposed linezolid as a good alternative for vancomycin, as it has better penetration in the CNS [12, 7, 13]. In general, if the abscess is more than 2–2.5 cm, neurosurgical intervention is indicated. Stereotactic aspiration is the treatment of choice for abscesses located in the brain stem [1, 5, 14].

**CONCLUSION**

Our case highlights the challenge of early diagnosis of brain abscess, an entity that presents very often with non-specific signs, the diagnosis of which could be easily missed, with repercussions on long-term disability and mortality. The mortality rate in patients with brain abscess remains high despite early detection and improvement in treatment. Long-term disability is also very high. Hence an early diagnosis and neurosurgical
intervention could prove significant in the final outcome for these patients.

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Author Contributions
Carlos Gonzalez – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Juan Roa – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published
Nehad Shabarek – Substantial contributions to conception and design, Acquisition of data, Analysis and interpretation of data, Drafting the article, Revising it critically for important intellectual content, Final approval of the version to be published

Guarantor
The corresponding author is the guarantor of submission.

Conflict of Interest
Authors declare no conflict of interest.

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