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CASE REPORT

# Fungal abscess of the parotid gland – the value of microbiological assessment

AK Ogonowski Bizos, M White, D J Fagan, D V Pretorius

Otorhinolaryngology Division, Faculty of Health Sciences, University of Cape Town, South Africa

Corresponding author, email: ent.nsh@gmail.com

### **Summary**

Fungal parotitis is rare and the sequela parotid abscess exceedingly so. We report our experience with Candida glabrata and Candida albicans parotid gland abscesses in critically ill HIV-positive patients and highlight the value of microbiological assessment to tailor management.

Keywords: fungal abscess, parotid gland, microbiological assessment

#### Case 1

A 58-year-old male with a 6-month history of pain in the right temporomandibular joint presented in decompensated left ventricular failure. He had a fluctuant mass in the preauricular area associated with painful mastication. He had ischaemic heart disease, hypertension, cardiac failure and chronic kidney disease (CKD). He was HIV positive (CD4 count of 323 cell/Ul, viral load [VL] was lower than detectable [LDL]). He was admitted for medical management for the cardiac failure and was started on oral clindamycin, empirically awaiting microscopy, sensitivity and culture (MC&S) of the pre-auricular abscess aspirate.

The aspirate cultured a light growth of Candida glabrata sensitive to amphotericin B. Fine needle aspiration cytology (FNAC) of the mass documented the presence of acidfluorescent bacilli along with degenerate inflammatory cells in a necrotic background. Based on the FNAC findings, he was started on empiric tuberculosis treatment but with no confirmed diagnosis of mycobacterium tuberculosis on culture and continued on tuberculosis treatment throughout his admission and on discharge. On discussion with microbiology, antifungal treatment was not initiated as it was thought to be a contaminant. He remained apyrexial on clindamycin and was continued on this antibiotic prior to referral to the ear, nose and throat (ENT) department. At that time, a tender, well-localised cystic swelling over the right parotid gland was present. There was extensive candidiasis in the oral cavity. The facial nerve was intact and there was no trismus or odontogenic sepsis. Oral fluconazole and oral nystatin were commenced to treat the extensive oral candidiasis while awaiting culture results of a second aspirate. This aspirate again yielded a moderate growth of C. glabrata, which was sensitive to amphotericin B. The infectious diseases department recommended treatment with oral fluconazole 200 milligrams (mg) daily to treat his oral candidiasis. A clinical review and repeat aspiration on day 13 yielded 2 ml of pus; thereafter there was complete resolution of the abscess. On day 25 of admission, the patient

contracted COVID-19. He was transferred to a stepdown facility where he died.

#### Case 2

A 46-year-old female presented with multiple comorbidities including hypertension, poorly controlled type 2 diabetes mellitus (DM), HIV on antiretroviral therapy (CD4 count of 84 cell/Ul, VL of 303 copies/ml), and tuberculous meningitis on treatment.

She presented to the emergency unit acutely unwell, in a hyperosmolar hyperglycaemic state, precipitated by sepsis (pH 7,1, base excess -20, WCC 34). She had a week's history of a painful swelling at the left angle of the mandible.

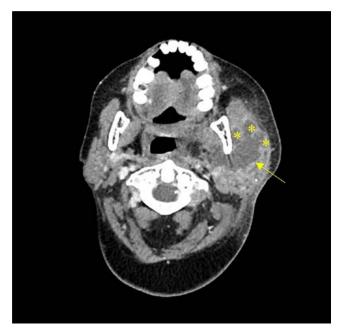


Figure 1: Contrasted CT scan depicting the left parotid gland abscess

Key: Arrow indicating intraparotid collection, asterisk indicating reactive lymphadenopathy

She had clinical features of acute parotitis with an associated abscess, with intact facial nerve and no trismus.

She was also noted to have oral candidiasis and odontogenic sepsis. Contrasted CT scan revealed a large multi-loculated, rim-enhancing, intraparenchymal collection of the parotid, with associated sub-centimetre, reactive cervical lymphadenopathy, and no evidence of sialolithiasis (Figure 1).

A bacterial suppurative parotid abscess was suspected, therefore amoxicillin/clavulanic acid therapy was initiated. She had an urgent incision and drainage (I&D) under general anaesthesia; 30 ml pus was drained from a collection involving both deep and superficial lobes of the parotid gland. Tissue specimens and pus were sent for MC&S, and both cultured Candida albicans sensitive to fluconazole. Histology from the abscess wall and parotid gland revealed moderate acute-on-chronic inflammation with no evidence of malignancy or granulomatous inflammation. Following consultation with the infectious diseases department on day 6 of admission, fluconazole was initiated and was continued for 1 month (400 milligrams daily po). She responded well to antifungal therapy and was discharged on day 14 with a small residual, granulating sinus at the site of previous I&D. At 2 months follow-up, she had complete resolution of the sinus and her surgical wound had healed with minimal scarring.

## **Discussion**

The most common suppurative infections of the parotid gland parenchyma in adults are of bacterial origin. <sup>1-6</sup> Fungal parotitis is extremely rare, with only 11 cases reported in the literature. <sup>3,5,8,9</sup> Of these, six cases of fungal parotitis were caused by *C. glabrata* and four cases were caused by *C. albicans*. <sup>3,5,7-9</sup>

All *C. glabrata* parotitis cases resulted in subsequent parotid gland abscesses.<sup>6</sup> Of the aforementioned cases, the ages ranged from 32 to 82 years. Underlying comorbidities included DM, HPT and CKD. Five of the six patients with *C. glabrata* parotitis were treated with fluconazole. All except one of the other patients with parotitis complicated with a parotid gland abscess due to *C. glabrata* fully recovered with a management approach of I&D, and fluconazole.<sup>3,5,8,9</sup>

Of the four previous cases presenting with suppurative parotitis due to *C. albicans*, three developed a parotid gland abscess. All were treated with fluconazole in addition to I&D with a resultant successful outcome; a negligible mortality rate has been reported. DM was the common underlying condition for all reported cases of *C. albicans* parotitis.<sup>4</sup> Of note, only two cases are documented to have HIV coinfection as a comorbidity.

There are currently no formal guidelines for medical management of fungal abscesses specifically relating to the head and neck region.

A review of the literature of case reports of candida parotitis show a variation in the dosage and duration of antifungal agents. There are four cases of previous parotid gland abscesses in which the dosage of antifungal agents is described and along with surgical management via incision and drainage, a successful outcome was reached.<sup>5,9</sup>

The principles of management of mycotic parotitis include timeous initiation of targeted antifungal therapy. In the majority of cases, high-dose fluconazole was considered

first line, and a 78% successful treatment rate has been reported.<sup>3-5,8,9</sup>

The surgical approach to a fungal parotid abscess should be graduated, where, in the absence of complications, less invasive, ultrasound-guided serial needle aspirations may suffice and will provide the necessary specimens for culture and sensitivity to direct antifungal therapy.<sup>6</sup> I&D may be required for source control in the presence of overwhelming sepsis, failed conservative treatment or with large collections with or without extension into the deep neck spaces.<sup>6</sup>

These two cases described depict the importance of primary management of a fungal parotid abscess via surgical source control and thereafter, culture directed antifungal therapy to avoid resistance. Mycotic infection should be considered as part of the differential in any patient presenting with a parotid abscess, especially in the context of immunosuppression and should be considered in the differential diagnosis.

Our approach was to manage parotid abscesses in two different ways as determined by the general clinical condition of the patient, with both having acceptable and clinically successful outcomes. Serial aspiration may be considered as initial first-line therapy. However, progression to incision and drainage may be considered in order to obtain source control in recalcitrant or complicated infections.<sup>6</sup>

# Conflict of interest

The authors declare no conflict of interest.

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#### Ethical approval

The author/s declare that this submission is in accordance with the principles laid down by the Responsible Research Publication Position Statements as developed at the 2<sup>nd</sup> World Conference on Research Integrity in Singapore, 2010. Informed verbal consent was obtained from one patient for being included in the case report, the second patient demised and informed verbal consent was obtained from the CEO of the hospital.

# **ORCID**

AK Ogonowski Bizos https://orcid.org/0000-0002-7887-8919

M White b <a href="https://orcid.org/0000-0003-3432-2086">https://orcid.org/0000-0001-3432-2086</a>
J Fagan b <a href="https://orcid.org/0000-0001-7924-7265">https://orcid.org/0000-0001-7924-7265</a>
V Pretorius b <a href="https://orcid.org/0000-0001-5907-5224">https://orcid.org/0000-0001-5907-5224</a>

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