



# Enfermedades Infecciosas y Microbiología Clínica

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Diagnosis at first sight

## Recurrent frontal tumour in a patient with repeated craniotomies\*

Tumoración frontal recidivante en paciente con craneotomías de repetición

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### Case report

32-Year-old male, intranasal drug user, who was involved in a road traffic accident when he was 18 years of age, with comminuted right frontal and supraorbital bone fractures and subsequent recurrent CSF leak (fistula), associated with recurrent meningitis caused by *H. influenzae* and *S. pneumoniae*, which required up to 3 prior surgeries. He was admitted in April 2011 due to pneumococcal (*S. pneumoniae*) meningitis associated with fluctuating swelling of the forehead and right frontal epidural abscess (Figs. 1 and 2).

A right frontal craniotomy and sinus drainage was performed, obtaining cultures positive for *S. pneumoniae*; after 6 weeks of treatment with intravenous ceftriaxone and an asymptomatic period with no recurrences, a right frontal cranioplasty (CustomBone®) was implanted in September 2012.

After one year symptom-free, the patient was readmitted in November 2013 with recurrent pneumococcal meningitis and with an underlying previous epidural abscess to the cranioplasty that had to be removed, debridement of the right frontal sinus and intravenous treatment with ceftriaxone followed by oral levofloxacin. Due to intolerance, the patient switched to clindamycin, which he received until he completed 6 months of treatment (June 2014).

### Clinical course

The patient was readmitted in October 2014 due to recurrence of fluctuating swelling with spontaneous supranasal suppuration and cultures positive for coagulase-negative *Staphylococci*. After administering vancomycin + ceftriaxone, and despite the brain MRI not showing obvious intracranial collections (Fig. 3), another frontal craniotomy, drainage and debridement of the left frontal sinus, supranasal sinus and residual right supraorbital sinus sealed with



Fig. 1. Right frontal region with fluctuating swelling.

wax were performed, and vancomycin and gentamycin were administered. The intraoperative cultures were negative and the universal 16S PCR undetectable, so intravenous treatment was completed for 6 weeks with IV teicoplanin 600 mg/24 h and ceftriaxone 2 g/12 h.

### Final comment

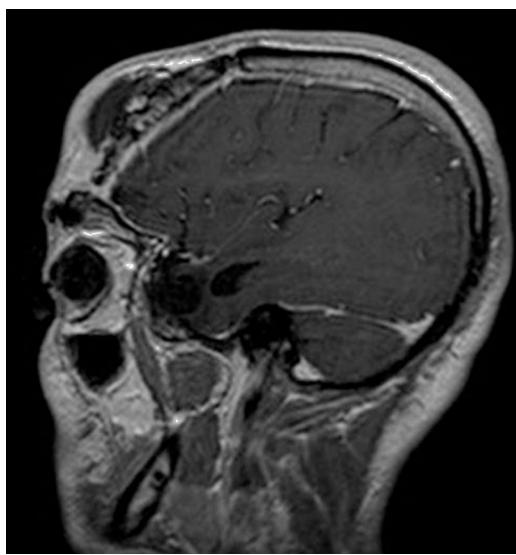
Suppurative osteomyelitis of the frontal bone, also known as Pott's Puffy Tumour, is a rare and little-known condition that is seldom reported in the medical literature. It is characterised by fluctuating swelling of the forehead secondary to a subperiosteal abscess caused by frontal bone osteomyelitis. It is most commonly seen in children and adolescents in whom the most widespread predisposing factor is acute or chronic frontal sinusitis.<sup>1</sup> It is much less common in adults, however, and the main risk factors include frontal bone trauma, prior craniotomy or cranioplasty, mastoiditis, nasal drug abuse, tooth infection and ethmoid sinusitis.<sup>2,3</sup>

The pathogenesis of this condition is closely related to the unique anatomy of the sinuses and the ease of propagation through the mucosal venous drainage of the frontal sinus towards the

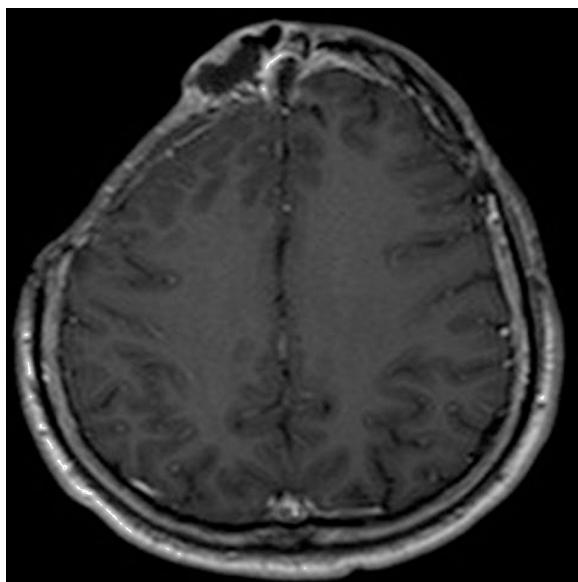
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**Fig. 2.** Brain MRI with gadolinium, sagittal plane.



**Fig. 3.** Brain MRI with gadolinium, coronal plane (October 2014).

diploë. The solutions of continuity in the sinus wall secondary to the risk factors described above facilitate progression of the infection. The clinical picture varies depending on the spread of the infection, and it may manifest as swelling of the frontal region without intracranial spread if it progresses towards the external table of the frontal bone, as an epidural abscess if it progresses towards the internal table or as orbital cellulitis if it spreads towards the supraorbital region.<sup>4,5</sup>

It is diagnosed on the basis of clinical and radiological findings. Typical symptoms in adults include fever, purulent rhinorrhoea, frontal or orbital cellulitis, fluctuating swelling or frontal fistula, as well as neurological symptoms such as meningitis or frontal focal symptoms.<sup>3</sup>

The CT scan is the gold standard for frontal bone osteomyelitis diagnosis, while MRI is effective at identifying intracranial complications. Labelled leucocyte scintigraphy may be useful in monitoring treatment response.<sup>6</sup>

The bacteria involved in the pathogenesis of the condition are mostly found in the saprophytic flora of the sinuses and lead to infection by anaerobic *S. aureus* and *Streptococcus* sp., Gram-negative bacteria and fungi.<sup>2</sup>

Treatment involves a combined medical-surgical approach: early broad-spectrum antibiotic treatment and the surgical drainage of intracranial collections (external or endoscopic) depending on the location and extent of the complications.<sup>7</sup>

Prognosis is variable and is determined by potential neurological sequelae and the high probability of recurrence, associated with significant morbidity owing to the need for multiple surgical procedures as shown in the case presented above.<sup>3,8</sup>

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### Conflicts of interest

The authors declare that they have no conflicts of interest.

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### References

1. Aínsa D, Pons S, Muñoz A, Vega MI, Otero MCL. Tumor inflamatorio de Pott: una complicación infrecuente de la sinusitis frontal. An Pediatr (Barc). 2014;80:317–20.
2. Akiyama K, Karaki M, Mori N. Evaluation of adult Pott's puffy tumor: our five cases and 27 literature cases. Laryngoscope. 2012;122:2382–8.
3. Salomao J, Cervante TP, Bellas AR, Boechat MC, Pone SM, de Pone MV, et al. Neurosurgical implications of Pott's puffy tumor in children and adolescents. Childs Nerv Syst. 2014;30:1527–34.
4. Nicoli T, Makitie A. Frontal sinusitis causing epidural abscess and puffy tumor. N Engl J Med. 2014;370:e18.
5. Ibarra S, Aguirrebengoa K, Pomposo I, Bereciartua E, Montejo M, González de Zarate P. Osteomyelitis of the frontal bone (Pott's puffy tumor). A report of 5 patients. Enferm Infect Microbiol Clin. 1999;17:489–92.
6. Ketenci I, Unlu Y, Tecer B, Vural A. The Pott's puffy tumor: a dangerous sign for intracranial complications. Eur Arch Otorhinolaryngol. 2011;268:1755–63.
7. Jung J, Lee HC, Park IH, Lee HM. Endoscopic endonasal treatment of a Pott's puffy tumor. Clin Exp Otorhinolaryngol. 2012;5:112–5.
8. Kombogiorgas D, Solanki GA. The Pott puffy tumor revisited: neurosurgical implications of this unforgotten entity. Case report and review of the literature. J Neurosurg. 2006;105 Suppl. 2:S143–9.