Chronic Invasive Fungal Sinusitis Mimicking a Sinonasal Malignancy: A Case Report

Lacy S. Brame, DO¹, Aniruddha C. Parikh, MD²; Kibwei A. McKinney, MD²; Rusha J. Patel, MD² ¹Oklahoma State University Center for Health Sciences, Tulsa, OK, 74107; ²University of Oklahoma Health Sciences Center, Department of Otolaryngology – Head & Neck Surgery, Oklahoma City, OK, 73104

Introduction

- Chronic invasive fungal sinusitis (CIFS) is a rare form of invasive fungal sinusitis.
- It is an insidious and invasive fungal infection which can extend intracranially and cause significant morbidity and mortality when untreated.
- Classically, CIFS has been associated with diabetes.
- The prevalence of CIFS may potentially be increasing given the growing use of long-term steroids and immunocompromised populations.

Methods



- The authors present a case of an immunocompetent patient with CIFS which mimicked a sinonasal malignancy.
- A comprehensive literature review was performed.

Case Report

- A 78-year-old male presented with a nasopharyngeal and sphenoid sinus mass. Nasopharyngoscopy showed submucosal fullness within the nasopharynx.
- He underwent a nasopharyngeal biopsy which was benign.
- He then underwent a sphenoidotomy with biopsy which was also benign but showed inflammation and fungal elements. There was no evidence of fungal invasion on frozen pathology.
- Final pathology of the second biopsy showed fungal angioinvasion concerning for invasive fungal sinusitis.
- MRI demonstrated involvement of the right temporal lobe and cavernous sinus.
- He was admitted and underwent endoscopic sinus surgery which included an extended dissection of the pterygopalatine and infratemporal fossae, the nasopharynx to the base of the clivus, and a sphenoid sinus drill out.
- He was treated with long-term anti-fungal therapy and has

Figure 3: Postoperative MRI demonstrating right temporal lobe involvement and the extent of surgical dissection within the nasopharynx and sphenoid sinus.



Figure 4: Postoperative MRI demonstrating involvement of the right cavernous sinus with disease surrounding the right internal carotid artery.

Results of Literature Review

been doing well postoperatively.



Figure 1: Preoperative CT demonstrating a nasopharyngeal and sphenoid sinus mass with erosion of the floor of the sphenoid sinus.



- A total of 16 articles were reviewed which discussed immunocompetency status in the presence of CIFS.
- A total of 95 cases were reported. 63 were immunocompetent and 32 were immunocompromised.
- Many articles did not discuss their inclusion criteria for immunocompetency. Moreover, many studies did not specify if patients were treated with corticosteroids for sinusitis, which may be a potential cause of immunocompromise.

Conclusions

- CIFS is a rare disease which can affect both immunocompetent and immunocompromised patients.
- Radiologically, it may mimic a sinonasal mass which may lead to a delay in diagnosis.
- CIFS should be considered a part of the differential diagnosis when evaluating atypical sinonasal lesions, especially when bony erosion is noted.
- Endoscopic biopsy and pathologic evaluation with fungal staining will assist in making the diagnosis.

Figure 2: Preoperative MRI demonstrating enhancement of the right temporal lobe and cavernous sinus.

 MRI may assist in preoperative diagnosis by identifying specific changes in the sinonasal cavity and identifying intracranial involvement.

References

- deShazo et al. A new classification and diagnostic criteria for invasive fungal sinusitis. Arch Otolaryngol Head Neck Surg. 1997 Nov;123(11):1181-8. doi: 10.1001/archotol.1997.01900110031005. PMID: 9366697.
- Li et al. Chronic invasive fungal rhinosinusitis vs sinonasal squamous cell carcinoma: the differentiating value of MRI. Eur Radiol. 2020 Aug;30(8):4466-4474. doi: 10.1007/s00330-020-06838-1. Epub 2020 Apr 11. PMID: 32279114.
- Alotaibi et al. Chronic Invasive Fungal Rhinosinusitis in Immunocompetent Patients: A Retrospective Chart Review. Front Surg. 2020 Dec 16;7:608342. doi: 10.3389/fsurg.2020.608342. PMID: 33392248; PMCID: PMC7772145.