Aberrant Maxillary Sinus Anatomy in a Cystic Fibrosis Patient

Timothy Baerg¹, Jennifer F Ha²,³,⁴, Sheena Bahrooloomi and David J Brown¹

¹University of Michigan Medical School, USA
²Department of Otolaryngology Head & Neck Surgery, University of Michigan Health System, USA
³Department of Surgery, University of Western Australia, Australia
⁴St John of God Hospital (Murdoch), Australia

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*Corresponding author: Jennifer Ha, Wexford Medical Center Suite, 17/18, Level 1, 3 Barry Marshall Parade, Murdoch 6150, Australia, Tel: +61863326868; Fax: +6186332 6800; E-mail: drjennha@yahoo.com.au

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Opinion

Cystic fibrosis (CF), a disease characterized by a heterozygous mutation in the Cystic Fibrosis Transmembrane Regulator (CFTR) gene, often involves the lungs, gastrointestinal tract, and upper respiratory tract. CF patients have an increased likelihood of developing chronic rhinosinusitis (CRS), especially those with a homozygous ΔF508 mutation [1]. CRS may result in decreased pneumatization of the frontal and sphenoid sinuses, manifesting in a decreased size or a complete absence of the sinus [1,2]. Independent of CRS, both homozygous and heterozygous mutations have been found to impair sinus development, leading to decreased anatomical volumes; indicating that paranasal size is correlated with CFTR function [2].

While CRS has been linked to abnormal sinus anatomy, CRS in the setting of CF has been linked to this phenomenon in a different way [3]. Lateral nasal wall medialisation, bilateral uncinated process demineralization, and middle meatus displacement have been described in CF with CRS, but these phenotypes are less commonly seen in patients with CRS only. These findings suggest that there is an elevated sinus pressure in CRS in the setting of CF. Moreover, erosion secondary to osteitis or nasal polyp and mucus mediated pressure can lead to the formation of pseudomucoceles [4]. CRS secondary to CF has been reported to occur in 63% of adult CF patients [1]. Due to prevalence and potential downstream complications that may occur, research characterizing the potential effects of aberrant sinus anatomy in the setting of CRS and CF has important clinical implications. We describe an unusual sinus anatomy secondary to CRS in a patient with CF.
A 16-year-old boy with a history of right nasal obstructions and purulent rhinorrhea on a background of left nasal septal deviation, and cystic fibrosis (CF) presented to our pediatric otorhinolaryngology clinic in August 2015. The flexible nasendoscopy confirmed large right-sided nasal polyposis (Figure 1), an apparent left inferior meatus window (Figure 2) and apparent maxillary antrostomy (Figure 3). However, he had no prior history of any sinonasal surgery. The computed tomography of his paranasal sinus confirmed the aberrant anatomy in the setting of his chronic rhinosinusitis (CRS) (Figure 4).

**Discussion**

Approximately 15 million Americans have at least one CFTR mutation, and one third of the mare expected to develop CRS [2]. While various studies have outlined that sinus size may decrease in the setting of CRS in CF, our patient’s findings have yet to be extensively explored. The aberrant anatomy in our patient may have developed as a result of the chronicity of the inflammatory process and bony erosion due to an expansile maxillary mucocele. The treatment of patients like ours is often conservative with systemic antibiotics, intranasal corticosteroids and nasal irrigations. Surgical treatment is reserved for large nasal polyposis when they significantly interfere with the patient’s quality of life.

The inferior meatus antrostomy procedure has been used to drain the maxillary sinus, but a maxillary antrostomy of the middle meatus is currently the standard of care. This procedure is effective for treating medically resistant CRS. However, in the setting of CRS, anterior bone displacement may occur, and ostial thickening may increase the challenge of surgery [5]. Since, however, the anterior portion of the sinus must be patent for effective mucociliary clearance, severe obstruction may be ablated through cutting instruments or cured powered microdebriders [5]. Complications of this are secondary to the stripping of the mucosa, leaving bone exposure on the posterior aspect of the lacrimal duct. An image guidance system should be used to aid in antrostomy.

**Conclusion**

To our knowledge, this is a first report of an aberrant inferior meatus ostium in the setting of CF with CRS. Since CRS is common in CF patients, it is important to be aware of anatomical aberrations for safe and successful surgical therapy. Through maxillary antrostomy, patients with this condition may avoid the downstream complications of CRS intervention.

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