**Mycobacterium avium-intracellulare** (MAI) sinusitis is rarely diagnosed. There are only nine previous cases documented in the literature, most of which are in patients with acquired immune deficiency syndrome (AIDS). Awareness of MAI sinusitis may prove valuable to the practicing otolaryngologist, and an elevated index of suspicion is necessary in all immunocompromised patients with sinusitis, especially those refractory to medical management. Unless specific investigations for MAI are ordered, the diagnosis may be missed and complications may result. We present the first case report of MAI sinusitis in a diabetic or organ transplant patient and the first in a non–HIV-infected immunocompromised adult. We also present a discussion of MAI and a literature review of MAI sinusitis.

**Case Report**

A 40-year-old male with type 1 diabetes, on immunosuppressive treatment 10 years post–renal transplantation, presented in August 2005 with a 6-week history of left-sided nasal obstruction, left-sided facial pain, hyposmia, and a mild left-sided headache.

He had no previous history of sinus disease or nasal trauma. His past medical history included the diagnosis of type 1 diabetes at the age of 7 years and a cadaveric renal transplant in 1994 at the age of 29 years while on hemodialysis for chronic renal failure. Other complications of his diabetes included retinopathy, a toe amputation owing to gangrene, and hypertension from his nephropathy. The patient was on multiple medications, including immunosuppressives (cyclosporine, prednisone, and mycophenolate mofetil) and insulin. The patient was human immunodeficiency virus (HIV) negative, and no other infectious diseases were identified.

His physical examination was remarkable for what was described as a papillomatous-like growth in his left nasal cavity, presenting on the septum and lateral nasal wall. The growth was biopsied, and the tissue sample was directly stained with auramine-O (fluorochrome technique) and showed + acid-fast bacilli (AFB). The sample was subsequently smeared, after cultures grew, using the Kinyoun acid-fast stain and showed non-tuberculous AFB; it was then probed for MAI. This isolate grew on both the Liquid–bioMérieux BacT/ALERT (bioMérieux Industry, France) (bioMérieux Incorporated, Durham, NC) and solid culture Lowenstein-Jensen media specific for mycobacteria. DNA probes (Gen-Probe AccuProbe system; Gen-Probe Incorporated, San Diego, CA) confirmed the diagnosis of MAI. The patient was subsequently referred to Infectious Diseases in September 2005, where he was followed monthly and started on a triple-antibiotic regimen consisting of rifampin 300 mg once daily, azithromycin 625 mg once daily, and ethambutol 800 mg once daily.

His symptoms persisted with no improvement despite treatment with the standard triple-antibiotic regimen for 5 months. As a result, the patient was referred to the St. Paul’s Hospital Sinus Centre. His physical examination was unremarkable except for a narrow left nasal cavity that was almost obstructed with the papillomatous-like growth. No pus, polyps, or other abnormalities were seen. A computed tomographic (CT) scan revealed complete opacification of his left frontal sinus, almost complete opacification of his left sphenoid sinus, mild-moderate right and left maxillary sinus mucosal thickening, and a left-sided nasal spur (Figure 1).

Owing to his persistent symptoms of sinusitis refractory to medical management, image-guided functional endoscopic sinus surgery (FESS) was carried out together...
with a nasal septal reconstruction. The left agger nasi cap was removed, and the frontal sinus was widely opened. The mucous membrane within the frontal sinus was thickened and fragile. Copious saline and antibiotic steroid solution lavage was carried out intraoperatively for the accessed sinuses. His triple-antibiotic therapy was continued postoperatively.

The patient showed an immediate improvement of his symptoms postoperatively. At 6 weeks postoperatively, he had no signs or symptoms of sinusitis. His sense of smell improved, and his nasal obstruction, headache, and facial pain were alleviated. On endoscopic examination, his nasal cavity and ethmoid, maxillary, and frontal sinuses appeared clear bilaterally, with no signs of infection. The patient’s sinuses appeared well healed from his surgery. His triple-antibiotic therapy was discontinued at that point. Four months after surgery, a CT scan showed resolution of the left sphenoid opacification (Figure 2). At the 1-year follow-up, he continued to have no signs or symptoms of sinusitis, with healthy and clean-appearing sinuses on endoscopic nasal examination.

**Discussion**

MAI is an atypical or nontuberculous mycobacterium. There is some controversy in the literature regarding the nomenclature, but in the clinical and health care realm, it is generally accepted that MAI is synonymous with *Mycobacterium avium* complex (MAC), and both terms represent an infection by either *Mycobacterium intracellulare* or *Mycobacterium avium*.5,10–12

The present case report is the first to document MAI sinusitis in a diabetic or organ transplant patient and the first in a non–HIV-infected immunocompromised adult. A previous literature review exists on nontuberculous mycobacterium sinusitis.7 Table 1 is an updated review of all previous cases of MAI sinusitis published in the English-language literature.1–7 There are two other cases

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**Figure 1.** Preoperative computed tomographic scans showing complete opacification of the left frontal sinus (A, B), mucosal thickening of the maxillary sinuses (C, D), and significant opacification of the left sphenoid sinus (E, F).
published in the international non–English-language literature, both in AIDS patients. Other cases of non-
MAI nontuberculous mycobacterial sinusitis have been
reported involving *Mycobacterium chelonae*,
*Mycobacterium kansasii*, and *Mycobacterium chelonae-
abscessus* complex.

MAI is omnipresent in the environment (soil, water,
animals, dust, and dairy products). Unlike *Mycobacterium tuberculosis*, human-to-human transmis-
sion is rare. Inhalation, ingestion, and direct
inoculation are thought to be the primary methods for
MAI colonization of the respiratory and gastrointestinal
tracts. It is possible that the proximal location of the
nasal cavity and paranasal sinuses in the aerodigestive tract in relation to the environment enables MAI to colonize
these areas more easily.

Standard cultures often do not grow mycobacteria. Mycobacteria-specific cultures should be obtained from the
sinuses for definitive diagnosis—AFB stains and histopathology alone are much less sensitive and specific than mycobacteria-specific cultures. With sufficient
cultures, molecular testing techniques such as DNA
probing can diagnose MAI within hours.

MAI infections in immunocompromised individuals are not uncommon, and its treatment in this population can be challenging. MAI is an opportunistic organism that is relatively more pathogenic than other environmental mycobacteria and very resistant to first-line antimycobac-
terial medications. Treatment with multiple antibiotic agents allows rapid clearance of MAI from the blood-
stream and decreases the danger of drug resistance. Long-term antibiotic therapy with a macrolide (azithro-
mycin or clarithromycin), in combination with companion
drugs (ethambutol and rifampin), is an accepted
protocol in the literature. Nonetheless, surgical
treatment of nontuberculous infections is often considered essential, as was the case in the present report.

MAI has a predilection to disseminate in the immu-


![Figure 2. Preoperative computed tomographic (CT) scans showing significant opacification of the left sphenoid sinus (A, B) versus postoperative CT scans showing complete resolution of the opacification (C, D).](image-url)
Table 1. Previous Cases of *Mycobacterium avium-intracellulare* Sinusitis

<table>
<thead>
<tr>
<th>Year</th>
<th>Study</th>
<th>Sinus</th>
<th>Age (yr)</th>
<th>Sex</th>
<th>Presentation</th>
<th>Risk Factors</th>
<th>Definitive Management</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>1992</td>
<td>Zurlo et al1</td>
<td>NR</td>
<td>NR</td>
<td>NR</td>
<td>Disseminated MAI</td>
<td>AIDS</td>
<td>Incision and drainage of subgaleal abscess and antimycobacterial agents</td>
<td>NR</td>
</tr>
<tr>
<td>1994</td>
<td>Naguib et al2</td>
<td>Frontal</td>
<td>33</td>
<td>M</td>
<td>Persistent acute frontal headaches and unilateral facial edema secondary to frontal sinus osteomyelitis and subgaleal abscess</td>
<td>AIDS</td>
<td>Drainage of maxillary and ethmoid sinuses; failed antibiotics and antimycobacteria-specific agents</td>
<td>Initial remission then relapse, death secondary to disseminated MAI 9 mo later</td>
</tr>
<tr>
<td>1995</td>
<td>Sussman3</td>
<td>Bilateral maxillary Child, age NR</td>
<td>F</td>
<td></td>
<td>Fever of unknown origin and history of chronic rhinosinusitis</td>
<td>AIDS</td>
<td>Drainage of maxillary and ethmoid sinuses; failed antibiotics and antimycobacterial agents</td>
<td>Death secondary to disseminated MAI 6 mo after presentation</td>
</tr>
<tr>
<td>1997</td>
<td>Weiss et al4</td>
<td>Frontal, maxillary, and ethmoid</td>
<td>13</td>
<td>F</td>
<td>Periorbital cellulitis and subperiosteal abscess (without osteomyelitis) secondary to frontoethmoid sinusitis</td>
<td>None</td>
<td>Endoscopic sinus surgery and external frontal sinus trephination with postoperative antimycobacterial agents</td>
<td>Resolved at 5 mo postoperatively</td>
</tr>
<tr>
<td>1997</td>
<td>Ferguson et al5</td>
<td>Pansinus disease</td>
<td>38</td>
<td>M</td>
<td>Acute pansinusitis and history of disseminated MAI</td>
<td>AIDS</td>
<td>Revision endoscopic sinus surgery with postoperative triple antimycobacterial agents</td>
<td>Death secondary to AIDS and disseminated MAI infection 2 mo postoperatively</td>
</tr>
<tr>
<td>2002</td>
<td>Mra et al6</td>
<td>Bilateral sphenoid</td>
<td>56</td>
<td>F</td>
<td>Orbital cellulitis with cranial nerve involvement and cavernous sinus thrombosis</td>
<td>None</td>
<td>Operative sphenoid sinus drainage; postoperative antibacterials, antimycobacteria-specific agents, and warfarin</td>
<td>Gradual partial recovery of eye at 1 yr postoperatively</td>
</tr>
<tr>
<td>2007</td>
<td>Brown et al7</td>
<td>Bilateral maxillary, ethmoid, and sphenoid</td>
<td>8</td>
<td>M</td>
<td>Recurrent rhinosinusitis with polyposis and headaches</td>
<td>CF</td>
<td>Endoscopic sinus surgery with postoperative antibiotics</td>
<td>Clinically well at 1 mo postoperatively</td>
</tr>
</tbody>
</table>

AIDS = acquired immune deficiency syndrome; CF = cystic fibrosis; MAI = *Mycobacterium avium-intracellulare*; NR = not reported.

*MAI and *Mycobacterium kansasii* coinfection.
many known diseases that MAI can cause in immunocompromised hosts, standard medical textbooks seldom mention MAI sinusitis.\textsuperscript{3,12} MAI infection in the sinuses can serve as a nidus for disseminated disease.\textsuperscript{2–5,11} MAI disseminates via the heterogeneous route and has a predilection to spread to many sites, including the bone marrow, spleen, and liver.\textsuperscript{2–5,11}

**Conclusion**

The present case report is the first to document MAI sinusitis in a non–HIV-infected immunocompromised adult. It is also the first case report of MAI sinusitis documented in a diabetic or organ transplant patient. Nontuberculous mycobacterial sinusitis should be considered in all immunocompromised patients presenting with signs and symptoms of sinusitis, especially those refractory to standard medical treatment.\textsuperscript{2,5} Routine cultures do not grow mycobacteria; thus, mycobacteria-specific cultures should be obtained from the sinuses in combination with molecular techniques such as DNA probing.\textsuperscript{5,17,20} AFB stains and histopathology alone are much less sensitive and specific.\textsuperscript{5,17,20} MAI infection of the sinuses can serve as a nidus for disseminated disease. A delay in diagnosis can result in high mortality owing to an increased likelihood of dissemination to vital organs in untreated immunocompromised patients. Meticulous FESS in conjunction with antimycobacterial triple medical therapy appears to be an effective and safe treatment option.

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**References**
