Case Report

Outcome and Complications of Frontal Sinus Stenting: A Case Presentation and Literature Review

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1. Introduction

Because endoscopic surgery is now commonly used to manage nasal pathologies, many endoscopic techniques for the management of simple and complex frontal sinus diseases have been developed. Frontal sinus disease remains challenging to manage because of the complex anatomy and narrow outflow tract [1]. Failure of frontal sinus surgery has been frequently reported. [2] The most common causes of failure in the frontal recess are remnant frontal recess cells, a retained uncinated process, middle turbinate lateralization, and postoperative stenosis of the frontal sinus outflow due to formation of scar tissue, synechiae, or osteoneogenesis [2].

A limited number of studies suggest the success of frontal sinus stenting to improve outcome in endoscopic frontal sinus surgery [3–7]. However, failure and complications of frontal sinus stenting may occur. Infection, pain, edema, and stent obstruction were reported [8].

Still there are no guidelines or universally accepted indications for the use of frontal sinus stenting; the use of it may depend on the surgeon’s decision and the operative scenario. Hosemann demonstrated that frontal sinus outflow tract (FSOT) stenosis is more in patients with the diameter of the neo-ostium less than 5 mm, and this may be considered as an indication for stenting [1].

Double J stent, frontal sinus stent acting as a local drug-releasing system, Rains self-retaining silicon stent,
doxycycline-releasing stent, and H-shaped silicon stent are examples of many types used [5, 9].

2. Case Report

A 16-year-old male was referred to our Tertiary/Quaternary Care Hospital, King Abdullah Medical City, in 2015 with a history of allergic fungal rhinosinusitis (AFRS) and functional endoscopic sinus surgery (FESS) one year ago. The operative report included left frontal sinus stenting using a biliary T-tube stent. The patient came to the outpatient department complaining of bilateral nasal obstruction and decreased smell sensation along with left ocular symptoms. Ophthalmic symptoms included left eye pain, discomfort, and redness without eye movement restriction.

On examination, recurrent grade 3 nasal polyps were found bilaterally. The CT report showed a migrating frontal stent that pushes against the left lamina papyracea (Figure 1). The lamina papyracea was intact in all previous follow-up CT scans. In our institution, the patient underwent extended endoscopic sinus surgery with the Draf III procedure performed by the senior author (I.R.H.). Intraoperatively, granulation tissue was forming around the stent, which was removed, and the specimen was sent for histopathological analysis (Figures 2 and 3). The latter was initially interpreted as possible granulomatous invasive fungal sinusitis, but then proved to be foreign body granulation tissue in reaction to the stent (Figure 4).
3. Discussion

The effectiveness of frontal sinus stenting is still a subject of debate in the literature. A limited number of studies suggest the effectiveness of using the stent in frontal sinus surgery (Table 1) [3–5, 7]. Weber et al. showed high effectiveness in the stented group using a silicon stent versus the nonstented group: the neo-ostium of the frontal sinus was patent in 80% of the stented group evaluated by endoscopy and CT/MRT together while it was patent in only 33% of the nonstented group (Table 1) [3].

On the other hand, Banhiran et al. studied long-term effects of frontal sinus stenting by evaluating the postoperative patency of frontal ostium and the postoperative improvement in patients’ symptoms among stented and nonstented groups. The results showed no statistical difference between the two groups in ostium patency and symptoms improvement (Table 1) [10].

The optimal duration of stenting the frontal sinus is unknown. Orlandi et al. demonstrated that stent can be very well tolerated postoperatively and the mean length was 31.6 months but without identifying the optimal duration [8]. One case reported a success stenting kept in place for 21 months without any complications [11]. In our case, the stent was kept in the patient for 12 months and was complicated by ocular symptoms as well as granulation tissue formation (Table 1).

In our present case report, the use of the stent was accompanied by different complications. The first complication was migration: our case report is the first describing such a complication with using the stent. The CT scan showed the migration of the stent through lamina papyracea which caused the ocular symptoms: left eye pain, discomfort, and redness.

The second complication was granulomatous tissue formation: granulomatous tissue was formed around the stent resulted in misinterpretation of the specimen histopathologically as granulomatous invasive fungal sinusitis, due to the presence of foreign body-type multinucleated giant cells, which is the characteristic finding in both conditions (Figure 4).

Other complications of the frontal sinus stent have been reported. Examples include secondary infection and hypersensitivity [12], bacterial biofilm on stent, and toxic shock syndrome [13, 14].

On examination, the patient had recurrent grade 3 nasal polyp, and obviously the presence of frontal sinus stent was not helpful in providing the appropriate irrigation to the frontal recess. Instead of keeping the patency of the ostium, the stent resulted in unwanted outcome. As well, the granulation tissue formed around the stent was histopathologically misinterpreted as a granulomatous invasive fungal sinusitis. Further studies are strongly needed to judge the effectiveness of using the stent.

Data Availability

No data were used to support this study.

Conflicts of Interest

The authors have no conflicts of interest to declare.

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Table 1: Summary of studies showing the effectiveness, optimal duration, and complications of frontal sinus stenting.

| Title | Author | Type of study | Year | Result |
|-------|--------|---------------|------|--------|
| Endonasal Frontal Sinus Surgery with Permanent Implantation of a Place Holder | Weber et al. | Controlled prospective | 1997 | 80% patent FSOT in the stented group vs. 33% in the nonstented group |
| The Success of 6-Months Stenting in Endonasal Frontal Sinus Surgery | Weber et al. | Retrospective | 2000 | Long-term stenting of the FSOT prevented restenosis in revision cases but did not prevent polyp regrowth |
| Frontal Sinus Stenting | Rains | Prospective (no control) | 2001 | 94% patent FSOT |
| Toxic Shock Syndrome Associated with Frontal Sinus Stents | Chadwell et al. | Case report | 2001 | TSS can be complicated by the frontal sinus stent |
| Evidence of Bacterial Biofilms on Frontal Recess Stents in Patients with Chronic Rhinosinusitis | Perloff et al. | Prospective | 2004 | Evidence of bacterial biofilms on the stent of 6 patients |
| Long-Term Effect of Stenting after an Endoscopic Modified Lothrop Procedure | Banhiran et al. | Controlled prospective | 2006 | No difference between stented and nonstented groups |
| Prolonged Stenting of the Frontal Sinus | Orlandi et al. | Retrospective | 2009 | The mean length of stenting was 31.6 months |
| Failed Endoscopic Sinus Surgery | Huang et al. | Retrospective | 2009 | Explained the causes of failed endoscopic sinus surgery |
| Long-Term Stenting for Chronic Frontal Sinus Disease | Hunter et al. | Retrospective case series | 2010 | Three cases were successfully treated with long-term frontal sinus stenting |
| Twenty One Months of Frontal Sinus Stenting | Ivana et al. | Case report | 2012 | The stent was kept in place 21 months |
| Double J Stent of Frontal Sinus Outflow Tract in Revision Frontal Sinus Surgery | Mansour H. | Prospective (no control) | 2013 | Four of the 5 patients (6 out of 7 sinuses) had a patent frontal outflow tract |
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