Modified Z-palatoplasty for correction of acquired nasopharyngeal stenosis following palatal surgery: our experience in nine patients

Mohamed Eesa¹, Ehsan Hendawy¹, and Mohammad Waheed El-Anwar¹

¹Zagazig University Faculty of Human Medicine

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Abstract

Abstract Purpose: to evaluate the safety and the efficacy of a modified Z-palatoplasty (ZPP) technique in management of post-surgical nasopharyngeal stenosis (NPS), describing its steps and results. Methods: This prospective study was conducted on patients with snoring± OSA due to acquired post-surgical NPS of grade I and II. Surgical repair using a modified ZPP was employed on the patients and the pre and postoperative results were statistically compared and adverse events were recorded. Results: The grade of NPS improved significantly postoperatively (p= 0.00136) throughout a follow up of one year. Postoperatively, there was statistically significant improvement of AHI (p= 0.0005), VAS of nasal obstruction (p<0.0001) and VAS of snoring (p<0.0001). While transient VPI and dysphagia disappeared within 3 months postoperatively. Conclusion: The utilized procedure appears fast, low cost, and easily applicable, and it does not require implants, special tools or suture materials. Furthermore, it gives promising results, with tolerable pain, and rapid recovery without significant or persistent complications.

Title

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Succinct key points

- Nasopharyngeal stenosis (NPS) is one of the challenging problems that can complicate palatal surgeries. Numerous diverse surgical interventions have been used to correct NPS; but, most are complicated and result in inconsistent outcomes
- The aim of the current study wasto evaluate the safety and the efficacy of a modified Z-palatoplasty (ZPP) technique in the management of post-surgical nasopharyngeal stenosis (NPS), describing its steps and results.
- Surgical repair using a modified ZPP was employed on patients with snoring \pm OSA due to acquired post-surgical NPS of grade I and II then the pre and postoperative results were statistically compared and adverse events were recorded.
- The grade of NPS improved significantly postoperatively (p=0.00136) throughout a follow up of one year-with significant improvement of AHI (p=0.0005), VAS of nasal obstruction (p<0.0001) and VAS of snoring (p<0.0001). While transient VPI and dysphagia disappeared within 3 months postoperatively.
- The utilized procedure appears fast, low cost, and easily applicable, and it does not require implants, special tools or suture materials. Furthermore, it gives promising results, with tolerable pain, and rapid recovery without significant or persistent complications.

Key words: Nasopharyngeal stenosis, palatal surgery, obstructive sleep apnea, snoring, apnea hypoapnea index, UPPP, velopharynx, ZPP.

Introduction

Nasopharyngeal stenosis (NPS) is a rare condition of obstruction of the communication between the nasopharynx and oropharynx due to concentric scar contracture of the tonsillar pillars, soft palate, and posterior pharyngeal wall. Mostof NPS nowadays are secondary to adenotonsillectomy, uvulopalatopharyngoplasty (UPPP), or radiotherapy for nasopharyngeal carcinoma¹. Symptoms of NPS vary from nasal obstruction, difficulty in nose blowing to snoring, hyponasal speech, obstructive sleep apnea (OSA), daytime fatigue, anosmia, rhinorrhea, and dysphagia². With increasing use of electrocautery and different palatal surgeries techniques, NPS incidence also increases³. Definitive treatment is often very difficult and may induce more scarring and restenosis. Nonetheless, not much has been written about this challenging complication and unique and standardized management has not yet been presented. Some articles discussed and presented diagrams of flaps that are difficult to understand and would appear to be difficult to reproduce and create, while others presented the idea of stenting⁴ that would appear very difficult to fit for a long time into this irregular, very mobile area. Z-palatoplasty (ZPP)⁵ was first described in 2001 for snoring and OSA but their Z-plasty requires intact anterior and posterior pillars and is meant for patients with intact tonsils. In 2004, Friedman et al described a modified ZPP especially in tonsillectomized patients⁶. Both techniques were dedicated for primary treatment of snoring and OSA. Now, the ZPP role has been diminished due to emergence of many lateral wall addressing techniques. We thought to use a modified form of ZPP as a salvage treatment for post-surgical NPS mainly after UPPP. Our aim was to evaluate the efficacy and safety of a new modified ZPP, describing its steps and results.

Materials and methods

This prospective study was conducted from April 2016 until December 2019 on patientswith snoring ± OSA due to post-surgical NPS of grade I and II.

NPS was classified into; **grade** I: minimal scar tissue, no soft palate lengthening, just adherence between lateral aspects of the palate and posterior pharyngeal wall, **grade** II: moderate scar tissue, soft palate lengthening, with a small opening in the soft palate (1-2 cm in diameter) and **grade** III: excessive scarring with complete fusion of soft palate and tonsillar pillars with the posterior pharyngeal wall (atresia) or stenosis leaving a pinpoint hole less than 1cm in diameter³.

Patients with grade III NPS, severe craniofacial anomalies affecting airway, limited mouth opening (interincisive distance <1.5cm), and/or unfit for general anaesthesiawere excluded.

This study was conducted according to the declaration of Helsinki on Biomedical Research Involving Human Subjects. An informative consent was signed by all patients. The [Blinded for review] University institutional review board approval was obtained.

Outcome measures:

- 1. A subjective measure of NPS by upper airways endoscopy and oral cavity exploration.
- 2. Baseline, 4weeks, 3months, and 1-year Visual Analog Scale (VAS 0–10) for nasal breathing, snoring by bed partner, and dysphagia (0 "no complaints" to 10 "severe complaint").
- 3. Baseline and 1-year follow-up polysomnography.
- 4. Short term morbidity and adverse events especially VPI.

Surgical technique (figure 1,2) Under general anaesthesia and patient's supine position, mouth gag was inserted. The operative site was infiltrated with 1% lidocaine with 1:100,000 epinephrine solution. A butterfly mucosal flaps were designed, marked, elevated and removed from the oral surface of the soft palate with caution to preserve the soft palate nasal mucosa. Then, the soft palate was divided in the midline paying attention not to injure the posterior pharyngeal wall mucosa. Excessive soft palate scar tissue was removed as required to facilitate eversion and suturing of the nasal mucosal surface to the oral side. Vicryl suture was placed on the edge of each side of the divided palate to facilitate its eversion. The closure was done in a centrifugal direction by vicryl in 2 layers; 1st one is submucosal inverted sutures and 2nd one is horizontal mattress mucosal sutures without tension and keeping the stitches on the oral side, not on theedge.By

this way, we aim to change the scar contracture tension lines to anterolateral vectors and to widen the anteroposterior and lateral oropharyngeal air spaces at palatal level. All patients received postoperative paracetamol, antibiotics and steroids, and asked to maintain on soft diet and fluids for 2 weeks. The SPSS program version 20 (Chicago, Illinois, USA) was used for statistical analysis. P-value <0.05 was considered significant.

Results

Eleven patients were included; nine completed the follow-up; 7 males and 2 females. Their mean age was 47.3±5.944 years (range=39-56). Eight patients were caused by UPPP while one caused by adenotonsillectomy. The duration between the causative operation and the NPS repair ranged between 6 and 21 years (mean=1.56±4.6). Preoperatively, the NPS was grade 1 in two patients (22.2%) and grade 2 in 7 patients (77.8%).

The operative bleeding was non-significant and recovery was eventless in all cases. The mean postoperative hospitalization was 1.8 ± 1.03 (range=1-4) days. Noinfection, primary or secondary hemorrhages were reported. Postoperative pain was tolerated and controlled by paracetamol orally and was completely relieved within 2 weeks. All patients resumed normal diet after 2 weeks. Five patients developed transient VPI that was noted occasionally with drinking fluids, but did not interfere with normal diet and disappeared completely within 3 months with no permanent VPI. No patients developed palatal fistula or restenosis >50%.

The grade of NPS improved significantly postoperatively ($X^2=13.2$, p=0.00136) throughout the follow up of one year. Preoperatively, all patients had OSA that was objectively documented by polysomnography with AHI ranged between 5.6 and 38.8 (mean= 23.2 ± 9.37). Postoperatively, the AHI showed significant improvement to a mean of 12.35 ± 3.75 (t=5.0089, p=0.0005) (Table 1).

The encountered complications were temporary and related to postoperative dry throat, and inability to clear the throat. Dysphagia showed early worsening but it improved completely at three months postoperatively and remained throughout the follow-up period (Table 2).

Discussion

NPS is a challenging problem that can complicate palatal surgeries. Diversesurgical interventions have been used to correct NPS; but, most are complicated and result in inconsistent outcomes⁴.

Theend-goal of repair remains the expansion of the nasopharyngeal-oropharyngeal communication through removal/release of the scar and mucosal coverage of denuded surfaces to limit recurrence. MacKenty described the first use of superiorly based pharyngeal mucosal flaps folded on themselves in 1927². This was subsequently modified by Kazanjian et al⁷ to include local oropharyngeal mucosal flaps. In more severe NPS cases, aggressive strategies included use of radial forearm and jejunal free flaps to treat recurrent stenosis after conventional flaps failure. Due to technical challenges of free tissue transfer, other local approaches developed; bilateral Z-plasty⁸ and bivalved palatal transposition flap⁹.

The NPS repair is difficult because it is rare clinical entity. Clinical series are usually limited to just a few patients and comparative studies are non-existent. Few reports in the literature described the usage of anyone specific technique for its surgical correction with lacking follow-up beyond 6 months in most series. In addition, there are no standardized outcome measures yet to detect the effectiveness of each surgical method. The success has been based mostly on subjective relief of symptoms⁹. But, we used the subjective VAS of symptoms, grading of stenosis and objective evaluation by AHI.

Most acquired NPS arise after UPPP. In the current study, UPPP was the cause in 88.9% of cases and males were more affected than females. Similarly, the two studied cases by Toh et al⁹ and the three repaired by Magdy et al⁴ were males.

Successful NPS correction relies on adequate scar tissue removal and/or lysis, and coverage of the raw mucosal surfaces⁹. So, we ensured complete scar tissue removal with mucosa preservation to use it to cover the raw

area and avoid the presence of two raw areas on the nasopharynx and palate.

In our study, there was no necessity for postoperative obturation that appears very difficult to fit for a long time into this irregular, very mobile and sensitive mucosal area and may be difficult to be tolerated by the patients beside its foreign body effect and reaction.

The current procedure was performed through the transoral rout without extra tool or suture material and so no financial implications.

Avoidance of excess palatal dissection reduced postoperative edema and soavoided ICU admission. Our follow-up period (12 months at least) exceeds these of past studies and thus the results appear promising.

Recently, Cammaroto et al¹⁰ published a technique for grade III NPS correction that represents a modification of bivalved palatal transposition flaps originally described by Toh et al⁹. However, our technique which represents a modified form of ZPP originally described by Friedman et al dedicated mainly for grade I and II NPS.

Thus, our technique for NPS repair showed satisfactory outcome and easy applicability. However, further studies on larger number of patients and in comparison to other techniques are recommended.

Conclusion

The described procedure appears to be effective, fast, low cost, and easily applicable, and it does not require implants, special tools, or sutures materials without significant complications.

Conflict of interest

The authors declare no conflict of interest.

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Figure legends

Figure 1: A; Mucosal flaps designed and marked. B; Elevating and removing mucosa from the oral surface of soft palate. C; Dividing soft palate in midline. D; Eversion and suturing the nasal mucosa to the oral one.

Figure 2: Pre and post-operative view of the palate.

Tables

Table 1 .Preopeative and 1-year postoperative follow up.

Outcome measures	Outcome measures	Preop	1 year postop	Test	P value
Grade of stenosis	Grade 0	0	6	$X^2 = 13.2$	0.00136 S
	Grade 1	2	3		
	Grade 2	7	0		
\mathbf{AHI}	\mathbf{AHI}	27.3 ± 6.276	12.35 ± 3.75	t = 5.0089	$0.0005 \; \mathrm{S}$
Nasal	Nasal	$6.78 {\pm} 0.916$	0.3 ± 0.67	t=17.1295	< 0.0001 S
breathing	breathing				
Snoring	Snoring	$6.25{\pm}0.968$	0.67 ± 0.94	t=12.4064	$< 0.0001 \; S$
Dysphagia	Dysphagia	3.5 ± 0.866	0	t=12.1247	< 0.0001 S

Legends: AHI, apnea hypoapnea index; S, significant

 ${f Table~2}$. Preoperative and postoperative VAS of symptoms

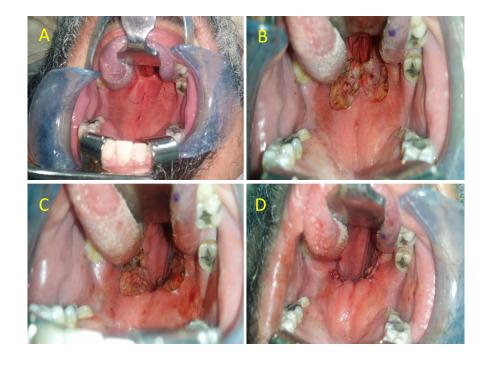
VAS		Preop VAS	One month postop	3 months post	1 year post	Statistical test
Nasal breathing	Range	5-8	1- 4	0-2	0-1	
	Mean	$6.78 \pm\ 0.916$	2.44 ± 0.83	0.3 ± 0.67	0.2 ± 0.4	F=159.669
Snoring	Range	5-8	1-5	0-2	0-2	
	Mean	$6.25 \pm\ 0.968$	3 ± 1.3	0.67 ± 0.94	$0.67 {\pm} 0.94$	F = 58.112
Dysphagia	Range	2-5	3-6	0	0	
	Mean	$3.5 \pm\ 0.866$	$4.2 \pm\ 0.916$	0	0	F=113.788

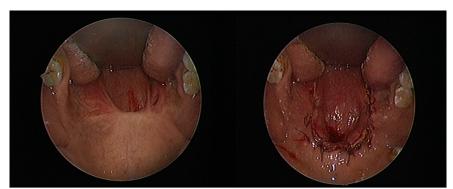
 $Legend: \ F = \ ANOVA \ test; \ S = \ significant$

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Table 1.docx available at https://authorea.com/users/341855/articles/468760-modified-z-palatoplasty-for-correction-of-acquired-nasopharyngeal-stenosis-following-palatal-surgery-our-experience-in-nine-patients

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