Palatal Procedures for Obstructive Sleep Apnea

Kathleen Yaremchuk, MD, MSA

KEYWORDS

- Uvulopalatopharyngoplasty Laser-assisted uvulopalatoplasty
- Radiofrequency volumetric tissue reduction Palatal implants
- Lateral pharyngoplasty Cautery-assisted palatal stiffening operation
- Z palatoplasty

KEY POINTS

- Uvulopalatoplasty is used to treat patients with obstructive sleep apnea (OSA) who have narrowing of the retro-palatal area of the pharynx.
- There have been many descriptions of procedures for the palate to decrease its length and move it anteriorly to increase the anterior posterior dimensions of the inferior margin of the palate.
- New techniques using radio frequency, laser, and implants have been used for snoring and OSA; but results have been similar to more surgically oriented techniques.

The procedure uvulopalatopharyngoplasty (UPPP) was first described for the treatment of snoring by Ikematsu in 1964.¹ Much later, in 1981, UPPP was described by Fujita and colleagues as "a new surgical approach" to treat obstructive sleep apnea.² Until then, permanent tracheostomy had been the only consistently effective surgical treatment in adult sleep apnea³ but resulted in psychosocial issues that were unacceptable to many patients. Fujita and colleagues described 12 predominantly male (11 of 12) patients with a history of excessive daytime sleepiness and loud habitual snoring. The velopharyngeal space was identified as the area of functional collapse of the pharynx during apneas. Clinically, the patients had a shallow oropharyngeal space with a relatively large uvula and redundant mucosa of the surrounding tissue.²

Fujita and colleagues⁴ subsequently described a series of 66 patients (63 men) treated for obstructive sleep apnea with UPPP. The mean apnea index (AI) preoperatively was 59. Significant improvements occurred after UPPP, although great variability was noted in individual patient response. Two subgroups were identified: responders (33 of 66) showed a significant decrease in AI of 84% (58.3–9.5), whereas

Disclosure Statement: The author has nothing to disclose. Department of Otolaryngology/Head and Neck Surgery, Henry Ford Hospital, 2799 West Grand Boulevard, Detroit, MI 48202, USA *E-mail address:* kyaremc1@hfbs.org nonresponders had little improvement (60.3–55.4). Despite recognizing the clear variability in response, they were unable to identify a sleep or respiratory parameter that differentiated the response to UPPP.

Some explored whether patients with mild sleep apnea might be more likely to benefit from surgical treatment. In an evaluation of 37 unselected patients with mild obstructive sleep apnea who underwent UPPP, Senior and colleagues⁵ found that only 40% had at least a 50% postoperative reduction in the respiratory event index (REI). Other patients had an increase in average REI from 161.6 \pm 5.0 to 26.7 \pm 18.4. Subjective assessment of sleepiness similarly was not improved. Again, the issue of responders and nonresponders arose.

In a review of the literature, Sher and colleagues⁶ noted that reports of case series and few controlled trials tend to limit the ability to advocate for change in surgical practice across the specialty. Their review found that UPPP was effective, at best, in less than 50% of patients with obstructive sleep apnea.

The variability in results of UPPP and the inability to predict which patients would respond to surgery became frustrating to otolaryngologists as many sleep medicine physicians refrained from referring patients for surgical interventions. Because of a success rate that was quoted as a 50:50 chance of improvement, many otolaryngologists attempted to improve surgical success results and to decrease the postoperative period in patients with obstructive sleep apnea with modifications of the traditional UPPP.

A breakthrough began when Friedman and colleagues^{7,8} classified patients with obstructive sleep apnea with a staging system based on body mass index (BMI), tonsil size, and palate position. Stage I was defined as a palate position 1 or 2 combined with tonsil size 3 or 4. Stage II was defined as palate position 3 or 4 and tonsil size 3 or 4. Stage III patients had palate position 3 or 4 and tonsil size 0, 1, or 2. Any patient with a BMI greater than 40 was stage III. In a retrospective analysis, UPPP alone had an 80% success rate in stage I patients, 37.9% success in stage II, and 8.1% success in stage III (Fig. 1, Table 1).

Multiple palatal procedures were developed to improve surgical success. An attempt to review all published procedures would be difficult because most are single series by individual surgeons and did not include large enough numbers or the technique was not adopted by others to reach significance within the specialty. Some of the procedures included new technology that was developed or applied in a novel way.

UVULOPALATOPHARYNGOPLASTY

Fujita described the UPPP as a procedure to remove redundant mucosa and preserve the muscular layer to enlarge the oropharyngeal space.² The procedure required general anesthesia, and patients were admitted to the hospital. The UPPP was performed by making an incision through the mucosa of the soft palate lateral to the glossopalatal arch from the inferior pole of the tonsillar fossa toward the uvula ending at its tip. The incision was extended on the pharyngeal side of the uvula and the pharyngopalatal arch toward the inferior pole of the tonsil. The mucosa of the soft palate, tonsillar fossa, and the lateral aspect of the uvula were undermined with sharp dissection and excised. The mucosal edges between the anterior and postural palatal arches were reapproximated with interrupted sutures. This maneuver brought the palatal arch forward or anteriorly with an increase in the anterior posterior dimension of the oropharyngeal space. If the uvula was elongated with the maneuver, then it was shortened or removed. If redundant tissue was in the posterior pharyngeal wall, an additional

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