

Trans-oral robotic surgery versus coblation tongue base reduction for obstructive sleep apnea syndrome

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Objectives. To compare the efficacy of trans-oral robotic surgery (TORS) with that of coblation assisted tongue base reduction surgery in patients with obstructive sleep apnea syndrome (OSAS). **Subjects and Methods.** The medical charts were retrospectively reviewed for all OSAS patients admitted to one institution for surgical intervention between 2012 and 2017. We analyzed 33 cases; 16 patients received TORS and 17 received coblation surgery for tongue base reduction. Both groups received concomitant uvulopalatoplasty. Surgical outcomes were evaluated by comparing the initial polysomnography (PSG) parameters with the follow-up PSG data at least 3 months after the surgery. Epworth sleepiness scale (ESS) and complications were also compared between the 2 groups.

Results. The success rate ($\geq 50\%$ reduction of pre-operative AHI and post-operative AHI < 20) in the TORS group and coblation group were 50% and 58%, respectively, and there was no significant difference ($p = .611$). The AHI (mean \pm SD) reduction in the TORS and coblation groups were 48.0 ± 38.9 events/h and 45.3 ± 34.1 events/h, respectively; the between-group difference was not significant ($p = .831$). ESS improvement did not differ significantly between the TORS and coblation groups (3.4 ± 3.0 and 2.8 ± 4.3 , respectively, $p = .646$). The rates of minor complication were higher in the TORS group (50%) than that of the coblation group (35.3%) without statistical significance ($p = .393$).

Conclusion. TORS achieved comparable surgical outcomes compared to coblation assisted tongue base reduction surgery in OSAS patients. Multilevel surgery using either TORS or coblation tongue base reduction combined with uvulopalatoplasty is an effective approach for the management of OSAS.

Trans-oral Robotic Surgery versus Coblation Tongue Base Reduction for Obstructive Sleep Apnea Syndrome

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19

20 **Abstract**

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22 coblation assisted tongue base reduction surgery in patients with obstructive sleep apnea
23 syndrome (OSAS).

24 **Subjects and Methods.** The medical charts were retrospectively reviewed for all OSAS
25 patients admitted to one institution for surgical intervention between 2012 and 2017. We
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31 **Results.** The success rate ($\geq 50\%$ reduction of pre-operative AHI and post-operative AHI
32 <20) in the TORS group and coblation group were 50% and 58%, respectively, and there
33 was no significant difference ($p = .611$). The AHI (mean \pm SD) reduction in the TORS and
34 coblation groups were 48.0 ± 38.9 events/h and 45.3 ± 34.1 events/h, respectively; the
35 between-group difference was not significant ($p = .831$). ESS improvement did not differ
36 significantly between the TORS and coblation groups (3.4 ± 3.0 and 2.8 ± 4.3 , respectively,

p= .646). The rates of minor complication were higher in the TORS group (50%) than that of the coblation group (35.3%) without statistical significance (p= .393). **Conclusion.** TORS achieved comparable surgical outcomes compared to coblation assisted tongue base reduction surgery in OSAS patients. Multilevel surgery using either TORS or coblation tongue base reduction combined with uvulopalatoplasty is an effective approach for the management of OSAS.

Introduction

Obstructive sleep apnea syndrome (OSAS) is a common disorder which affects 3-7% of adult men and 2-5% of adult women.¹ OSAS results from upper airway collapse during sleep. Clinical symptoms include fragmented sleep and excessive daytime sleepiness.² Continuous positive airway pressure (CPAP) is thought to be the gold standard treatment for OSAS,³ but some patients cannot tolerate it and may seek surgical treatment instead.⁴ Different levels and degrees of obstruction in OSAS patients lead to variable response to surgical intervention.⁵ In one study, Vroego et al. analyzed the upper airway collapse patterns in patients with sleep-disordered breathing by using drug-induced sleep endoscopy (DISE) and multilevel collapse was disclosed in 68.2% of all patients.⁶ As the intricacies of airway collapse are better understood, due to improvements in diagnostic

and evaluative methods, multilevel surgery is becoming a more common method of successfully treating OSAS.^{7, 8} Among these patients with multilevel collapse, the most frequently seen pattern was the concomitant collapse of palatal and tongue base (25.5%).⁶ Uvulopalatopharyngoplasty (UPPP) is the most commonly reported surgery to address oropharyngeal obstruction. For dealing with tongue base obstruction, trans-oral robotic surgery (TORS) and coblation assisted tongue base reduction surgery were two of the most published tongue base tissue reduction procedures.

Several preoperative assessment strategies have been used. Friedman tonsil grading scale classifies the tonsil size into five grades (grade 0-IV) according to the location the tonsil relative to the surrounding structures.⁹ Friedman tongue position (FTP) grading system is evaluated similarly to the modified Mallampati classification, but the tongue is evaluated in a neutral position without protrusion. Friedman staging system incorporates FTP, Friedman tonsil grading scale and BMI to classify OSAS patients into four stages: stage I includes patients with tonsils graded III-IV, FTP graded I-II and BMI<40 kg/M²; stage III includes patients with tonsils graded 0-II, FTP graded III-IV and BMI<40 kg/M²; stage IV includes patients with BMI>40 kg/M² or significant craniofacial or other anatomic abnormalities; stage II includes patients beyond stage I, III, IV.¹⁰ Fiberoptic nasopharyngoscopy with Muller's maneuver, which mimics the pathophysiological status

73 of OSAS during wakefulness by asking the patient to block bilateral nostrils and inhale
 74 with mouth closed, can identify the level and degree of upper airway collapse.¹¹ Drug-
 75 induced sleep endoscopy (DISE), which is recognized as a breakthrough in evaluation of
 76 OSAS patients, can provide direct identification of airway collapse during intravenous
 77 anesthesia. The VOTE classification is utilized for the findings of DISE.¹²

78 For most patients with oropharyngeal obstruction, uvulopalatopharyngoplasty (UPPP) is
 79 one of the most common and effective surgical procedures.¹³ However, oropharyngeal
 80 obstruction combined with tongue base obstruction is recognized as the most important
 81 reason for failure after pharyngoplasty procedures.¹⁴ For tongue base obstruction,
 82 multiple procedures have been proposed and could be simply categorized into tongue
 83 base volume reduction and tongue suspension. Among these procedures, trans-oral
 84 robotic surgery (TORS) and coblation assisted tongue base reduction surgery proved to
 85 be the most published therapeutic methods in the field of the tongue base reduction.¹⁵

86 TORS can provide a 3D visual field and the operator can easily access the tongue base
 87 area and perform surgery using delicately controlled robotic instruments. Nevertheless,
 88 the high cost of TORS makes operators and patients hesitant to make use of it.¹⁵

89 Endoscopic coblation assisted tongue base reduction surgery has been reported to be a
 90 useful procedure for tongue base obstruction and it has a lower cost compared to TORS.

^{16, 17} However, there is a lack of fair comparison studies regarding the treatment efficacy and safety between TORS and coblation assisted tongue base reduction. Therefore, this study was conducted to compare the subjective and objective outcomes of TORS with endoscope-guided coblation tongue base reduction.

Materials and methods

Medical charts were retrospectively reviewed for OSAS patients admitted for TORS or coblation tongue base reduction surgery to a single tertiary hospital between 2012 and 2017. Thirty three patients with age ranging from 18 to 62 years met the inclusion criteria (Table 1). Patients who were excluded were those without available postoperative polysomnography (PSG) data. PSG was performed at 3-12 months after the surgery. Patients who had previous upper airway surgery for OSAS were also excluded. This study was approved by the Institutional Review Board of the China Medical University Hospital (project approval number CMUH103-REC1-078).

Detailed profiles were constructed for each patient and included the following variables: age, sex, body mass index (BMI), tonsil grade, Friedman tongue position, Friedman stage, pre-operative and post-operative Epworth sleepiness scale (ESS). Post-operative ESS was recorded at the date for post-operative PSG. Nasopharyngoscopy with Muller's

maneuver and drug-induced sleep endoscopy (DISE) were performed in all patients to evaluate the site of obstruction and the pattern of the airway collapse.

The grades of airway collapse in Muller's maneuver were divided into four grades according to the percentage change in cross-sectional area: grade I $\leq 25\%$ collapse, grade II $>25\%$ and $\leq 50\%$ collapse, grade III $>50\%$ and $\leq 75\%$ collapse, grade IV $>75\%$ collapse. VOTE classification was utilized for reporting DISE findings and the grade of collapse were classified as 0 ($< 50\%$ obstruction); 1 (50–75% obstruction) and 2 ($> 75\%$ obstruction). Patients undergoing surgery must have at least partial tongue base collapse confirmed by Muller's maneuver and DISE. Details were also recorded from pre-operative and post-operative PSG data, and included AHI, apnea index (AI), lowest oxygen saturation (min-SpO₂), cumulative time percentage with SpO₂ $< 90\%$ (CT90) (Table 2). The success of the surgery was defined as achievement of $\geq 50\%$ reduction of pre-operative AHI and a post-operative AHI < 20 . Perioperative parameters, including the length of stay in hospital, the numeric rating scale (NRS) for pain intensity assessment on the first postoperative day and complications, were recorded.

In this study, 16 patients received TORS and 17 patients received coblation surgery for tongue base reduction (Figure 1). All patients received conventional uvulopalatopharyngoplasty combined with tongue base reduction for multilevel

obstruction in these patients. All of the surgeries were performed by a single surgeon.

The surgical procedure of trans-oral robotic surgery for tongue base volume reduction

was performed similar to the previous published literature.^{16, 18} General anesthesia was

introduced via nasotracheal intubation. The anesthesia machine was positioned at the

left side foot of the bed. The surgical cart of the da Vinci surgical system (Intuitive Surgical,

Sunnyvale, California, USA) approached the patient from the right-hand side with an

angle of 45 degrees to the bed. The scrub nurse stood next to the patient's left hand and

the first assistant sat at the head of the bed. The operative surgeon was at the operative

console and used open-surgery hand movements which were precisely replicated in the

operative field by the robotic instruments. The laryngeal advanced retractor system

(Fentex, Tuttlingen, Germany) was used to expose the tongue base area. The size of the

tongue blade was chosen accordingly to well expose the tongue base. Under 30 degree

3D camera endoscope, tongue base tissue was grasped by robotic forceps and

cauterized with spatula monopolar electrode. The midline posterior glossectomy began

from the foramen cecum and advanced posteriorly to vallecula without injury to epiglottis

mucosa, laterally to 1cm from the midline and 1.5 cm inferior to the tongue base surface.

Endoscopic coblation assisted tongue base reduction surgery was performed similar to

previous reports.^{19, 20} Under general anesthesia with nasotracheal intubation, the Molt

mouth gag (Sklar, West Chester, Pennsylvania, USA) was applied to the left side of labial commissure. We placed a silk suture through the anterior tongue and the silk was held by a Kelly forceps. The first assistant could easily retract the tongue forward by holding the Kelly forceps. A 70 degree rigid endoscope (Karl Storz, Tuttlingen, Germany) was applied to expose the tongue base area and kept in position by an endoscope holder (Karl Storz, Tuttlingen, Germany). With the aid of the endoscope holder, the surgeon could perform the procedure bimanually and thus decrease the operation time and the morbidity related to blood loss. The Coblator II ENT Surgery System and PROCISE MAX coblation wand (Arthrocare ENT, Sunnyvale, California, USA) were used for the midline posterior glossectomy. The targeted resection area was the same as the TORS mentioned above. We used the Statistical Packages for the Social Sciences version 24.0 (IBM Corp.; Armonk, NY, USA) for statistical analysis of the data. The descriptive statistic was used to present the outcome values. The Mann-Whitney and Wilcoxon test were used for comparing numerical variables, and Fisher's test was used for categorical variables. A p value of less than .05 was considered to be statistically significant.

Result

Among the 33 patients in this analysis, 16 were in the TORS group (age of 39.4 ± 12.3

years) and 17 patients were in the coblation group (age of 38.7 ± 11.5 years). The male comprised 93.8% in the TORS group and 76.5% in the coblation group. The body mass index (BMI) at the time of admission was 28.1 ± 3.8 kg/m² in the TORS group and 27.3 ± 5.5 kg/m² in the coblation group. There were no significant differences in tonsil grading scale, Friedman tongue position and Friedman staging system between the two groups before surgery. The grades of collapse in Muller's maneuver and DISE were similar in both groups. The Epworth sleepiness scale (ESS) was 11.6 ± 4.6 in the TORS group and 10.8 ± 5.1 in the coblation group. All patients received polysomnography (PSG) for pre-operative evaluation. The baseline apnea-hyponea index (AHI) was 50.4 ± 19.6 events/h and mean apnea index (AI) was 34.3 ± 20.3 events/h in the TORS group; corresponding values were 44.8 ± 28.8 events/h and 28.2 ± 26.7 events/h, respectively, in the coblation group. The mean lowest oxygen saturation (min-SpO₂) was $73.7 \pm 7.0\%$ and mean cumulative time percentage with SpO₂ < 90% (CT90) was $16.4 \pm 15.0\%$ in the TORS group; corresponding values were $74.0 \pm 10.0\%$ and $13.6 \pm 16.1\%$ in the coblation group. Demographics, baseline PSG data for both groups are summarized in Table 2. There were no significant between-group differences prior to treatment. The comparisons within-group (Table 3) and between-group (Table 4) were analyzed, respectively. Statistically significant improvement of ESS was observed in both groups.

181 ESS improvement did not differ significantly between the TORS and coblation groups
 182 (3.4 ± 3.0 and 2.8 ± 4.3 , respectively, $p = .646$; 95% CI= $-3.30 \sim 2.08$, Figure 2). The AHI
 183 reduced significantly from 50.4 ± 19.6 events/h to 25.5 ± 19.4 events/h in the TORS group
 184 ($p = .002$). In the coblation group, the mean AHI reduced significantly from 44.8 ± 28.8
 185 events/h to 25.4 ± 23.2 events/h ($p = .005$). The AHI reduction in the TORS and coblation
 186 groups were 48.0 ± 38.9 events/h and 45.3 ± 34.1 events/h, respectively; the between-
 187 group difference was not significant ($p = .831$; 95% CI= $-12.67 \sim 23.73$, Figure 3). The
 188 mean AI reduced significantly in both TORS and coblation group ($p = .005$ and $p = .018$,
 189 respectively), but the mean AI reduction did not differ significantly between the groups as
 190 well ($p = .481$; 95% CI= $-18.53 \sim 14.60$, Figure 3). The min-SpO₂ improved from
 191 $73.7 \pm 7.0\%$ to $83.9 \pm 5.7\%$ ($p = .001$) in the TORS group and from $74.0 \pm 10.0\%$ to
 192 $80.6 \pm 12.5\%$ in the coblation group ($p = .045$). The improvement of min-SpO₂ was
 193 $10.2 \pm 7.9\%$ in the TORS group and $6.6 \pm 12.5\%$ in the coblation group. There were no
 194 significant statistic differences in the improvement of min-SpO₂ between the two groups
 195 ($p = .355$; 95% CI= $-4.11 \sim 10.82$, Figure 4). The TORS group patients had more reduction
 196 of CT90 percentage, but the difference was not significant. The success rate in the TORS
 197 group and coblation group were 50% and 58%, respectively, and there was no statistically
 198 significant difference ($p = .611$; 95% CI= $-4.93 \sim 13.79$, Figure 4).

The average numeric rating scales (NRS) for pain evaluation on the first postoperative day were similar in both groups ($p = .428$). In the TORS group, the length of stay in hospital was longer compared with the coblation group ($p = .002$). There were no major complications (e.g., intra-operative or post-operative bleeding, airway compromise, prolonged intubation, pneumonia and pharyngeal laceration, tongue limitation) in either group. No tracheotomies were performed for airway management perioperatively. The rates of minor complication, including transient dysphagia, pharyngeal edema and dysgeusia, in the TORS and coblation groups were 50% and 35.3%, respectively.

Discussion

Our results demonstrate that the surgical outcomes of trans-oral robotic surgery (TORS) were comparable to coblation assisted tongue base reduction surgery in obstructive sleep apnea syndrome (OSAS) patients. The PSG outcomes and success rate were similar for the TORS and coblation groups.

Multilevel surgery is thought to be a successful management for patients suffering from obstructive sleep apnea syndrome (OSAS). Because of the better understanding of the complexity of the upper airway collapse during sleep in OSAS patients, surgeons can determine correct surgical management according to the site of obstruction and the

217 pattern of the airway collapse.^{8, 17, 21}

218 In a retrospective study, 25 moderate-to-severe OSAS patients with retropalatal and
 219 tongue base obstruction received coblation endoscopic lingual lightening and modified
 220 uvulopalatopharyngoplasty (relocation pharyngoplasty). AHI decreased significantly from
 221 45.7 ± 21.7 to 12.8 ± 8.2 events/hour ($p < .001$) postoperatively and the overall surgical
 222 success rate was 80%.²⁰ Coblation lingual tonsil removal technique proved to be an
 223 effective procedure in a cohort of Korean OSAS patients with retroglossal obstruction.
 224 The average AHI decreased significantly from 37.7 ± 18.6 to 18.7 ± 14.8 events/hour
 225 ($p < .001$) and the success rate was 55.6%.¹⁹ Another study compared combined coblation
 226 endoscopic lingual lightening and relocation pharyngoplasty to relocation pharyngoplasty
 227 alone in OSAS patients (AHI > 20, Friedman stage III), and reported that combined surgery
 228 had better improvement in AHI (-65.5 vs -53.2 ; $p = .047$) and higher surgical success rate
 229 than relocation pharyngoplasty alone (73% vs 50%; $p = .04$).¹⁷

230 O'Malley et al. developed a minimally invasive surgical procedure for management of
 231 tongue base neoplasms by using robotic surgical instruments.²² Trans-oral robotic
 232 surgery (TORS) can offer clear 3D visualization and gain adequate access to tongue
 233 base, larynx and hypopharynx and provide meticulous tissue resection. A preliminary
 234 study in 2010 conducted by Vicini et al. reported that TORS for tongue base resection in

OSAS patients is practical and well tolerated. Ten patients were included and the AHI decreased from 38.3 ± 23.5 to 20.6 ± 17.3 events/hour.¹⁸ Further study for demonstration of the feasibility of TORS performed in forty four patients with OSAS reported significant improvement of mean AHI (24.6 ± 22.2 events/hour) and mean ESS (5.9 ± 4.4).²³ The latest systematic review and meta-analysis by Meccariello et al. concluded that TORS seems to be a promising and safe technology for the management of OSAS and the mean failure rate was 34.4% (29.5–46.2%).²⁴

A study by Friedman et al. in 2012 was thought to be the first comparative study for the comparison of coblation and TORS in OSAS treatment.¹⁶ It compared the effectiveness of TORS with that of coblation assisted submucosal minimally invasive lingual excision (SMILE). All the patients in the study received concomitant z-palatoplasty. The AHI reduction in the TORS and SMILE groups were $60.5\% \pm 24.9\%$ and $32.0\% \pm 43.3\%$ ($p = .012$), respectively. The success rate in the TORS and SMILE groups were 66.7% and 45.5%, respectively; the between-group difference was not significant ($p = .135$). However, the techniques used by each group were different and not completely comparable.

To the best of our knowledge, there is a lack of fine and matched studies regarding the treatment efficacy and safety between TORS and coblation adopting similar technique in

tongue base resection. Our retrospective comparison of TORS with coblation in the treatment of OSAS patients with multilevel obstruction found that both groups had similar surgical results.

The demographics and preoperative polysomnographic data did not differ significantly between the two groups (Table 2) at baseline. According to the within-group outcomes showed in Table 3, statistically significant improvement of apnea-hyponea index (AHI), apnea index (AI), Epworth Sleepiness scale (ESS) and minimum oxygen saturation (min-SpO₂) were noted in both the TORS and coblation groups. It confirmed that either TORS tongue base resection or coblation assisted tongue base resection combined with concomitant uvulopalatoplasty can offer reliable surgical results. The cumulative time percentage with SpO₂ < 90% (CT90) were decreased in both groups but only significantly reduced in the TORS group which could be related to small sample size or poor correlation of CT90 to AHI.²⁵

As detailed in Table 4, the mean reduction of AHI, AI, ESS, CT90 and mean improvement of min-SpO₂ were similar for the TORS and coblation groups. The rate of surgical success in the TORS group were comparable to the coblation group (50.0% vs 58.8%, p= .611). Hwang et al. compared the tongue base coblation resection to TORS in OSAS patients and both groups were in combination with lateral pharyngoplasty.²⁶ They reported that

271 the surgical success rates did not differ significantly between the two groups (56.3% in
272 TORS vs 62.1% in coblation, $p = .711$). Our success rates are lower than those in that
273 study. However, the preoperative BMI of patients in that study was lower than in our study
274 group (25.8-26.8kg/m² vs 27.3-28.1kg/m²). Moreover, preoperative mean ESS were
275 lower (8.5-9.7 vs 10.8-11.6) and mean min-SpO₂ were higher (78.5-79.8 vs 73.7-74.0) in
276 their study than those in our study group, which might suggest the severity of OSAS
277 is greater in our patients. In our study, the average pain scores (numeric rating scales)
278 on the first postoperative day were comparable in both groups ($p = .428$). In the TORS
279 group, the length of stay in hospital was longer compared with the coblation group ($p =$
280 $.002$). There was no major complication in either group. The rates of minor complication
281 were higher in the TORS group (50%) than that of the coblation group (35.3%) without
282 statistical significance. According to a review article, slightly better outcomes were
283 observed in TORS compared to coblation, but the higher rate of minor complications and
284 the significant costs of TORS are two aspects which surgeons will need to consider.¹⁵
285 This study has some limitations. First, the retrospective analysis used in this study is a
286 possible source for selection bias. Second, it is difficult to make comparisons among
287 studies because of different surgical techniques utilized by TORS (e.g. midline posterior
288 glossectomy²⁷, lingual tonsillectomy²⁸) and coblation (e.g. midline posterior

glossectomy²⁰, SMILE²⁹, channelling of the tongue³⁰, Interstitial injections with
 needle coblation³¹). In the future, prospective, randomized, controlled trials that
 incorporate similar surgical technique will be needed to evaluate the efficacy of TORS
 compared with coblation tongue base reduction. Moreover, studies providing long-term
 results in the treatment of OSAS are also warranted.

Conclusion

TORS resulted in comparable objective and subjective outcomes compared to coblation
 assisted tongue base reduction surgery in OSAS patients. Multilevel surgery with either
 TORS or coblation tongue base reduction combined with uvulopalatoplasty is effective in
 reducing disease severity in moderate-to-severe OSAS cases.

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Table 1 (on next page)

Inclusion criteria

AHI = Apnea-Hypopnea Index; DISE = Drug-Induced Sleep Endoscopy; CPAP = Continuous positive airway pressure

Table 1:

Inclusion criteria

≥ 18 years old
Symptoms of obstructive sleep apnea syndrome (snoring, disrupted sleep, daytime sleepiness)
Preoperative AHI > 20
Friedman tongue position grade 3 or 4
Partial or complete retropalatal and retroglossal collapse in Muller's maneuver and DISE
Cannot tolerate CPAP

AHI = Apnea-Hypopnea Index; DISE = Drug-Induced Sleep Endoscopy; CPAP = Continuous positive airway pressure

Table 2 (on next page)

Demographics, Baseline data of the 2 groupst

BMI=body mass index (weight in kilograms divided by height in meters squared); FTP = Friedman tongue position; DISE = Drug-Induced Sleep Endoscopy; ESS = Epworth Sleepiness scale; AHI = Apnea-Hypopnea index; AI = Apnea index; Min-SpO2 = minimum oxygen saturation; CT90 = cumulative time percentage with SpO2 < 90% † All values are presented as mean ± standard deviation

TABLE 2.
Demographics, Baseline data of the 2 groups†

	TORS group (n=16)	Coblation group (n=17)	p value
age, years	39.4±12.3	38.7±11.5	.861
Male: Female ratio, n (%)	15:1 (93.8:6.3)	13:4 (76.5:23.5)	.335
BMI, kg/m ²	28.1±3.8	27.3±5.5	.645
Tonsil grade	2.0±1.3	1.9±0.8	.764
FTP	3.4±0.6	3.3±0.6	.831
Friedman stage	2.4±0.6	2.7±0.6	.207
Grade of collapse in Muller maneuver			
Retropalatal area	3.1±0.9	3.6±0.6	.122
Retroglossal area	2.6±0.8	2.3±0.8	.349
Grade of collapse in DISE			
Velum	2.0±0.0	1.6±0.6	.423
Oropharynx	1.4±0.5	1.3±0.6	.807
Tongue base	1.4±0.5	1.3±0.6	.807
Epiglottis	0.8±0.8	0.3±0.6	.351
ESS	11.6±4.6	10.8±5.1	.917
AHI, events/hour	50.4±19.6	44.8±28.8	.517
AI, events/hour	34.3±20.3	28.2±26.7	.498
Min-SpO ₂ , %	73.7±7.0	74.0±10.0	.932
CT90, %	16.4±15.0	13.6±16.1	.641

BMI=body mass index (weight in kilograms divided by height in meters squared); FTP = Friedman tongue position; DISE = Drug-Induced Sleep Endoscopy; ESS = Epworth Sleepiness scale; AHI = Apnea-Hypopnea index; AI = Apnea index; Min-SpO₂ = minimum oxygen saturation; CT90 = cumulative time percentage with SpO₂ < 90%

† All values are presented as mean ± standard deviation

Table 3 (on next page)

Within-group comparison of the treatment outcomes†

AHI = Apnea-Hypopnea index; AI = Apnea index; ESS = Epworth Sleepiness scale; Min-SpO2 = minimum oxygen saturation; CT90 = cumulative time percentage with SpO2 < 90%

† All values are presented as mean ± standard deviation * p < .05 is considered statistically significant

TABLE 3.

Within-group comparison of the treatment outcomes†

	TORS group (n = 16)			Coblation group (n = 17)		
	Preoperative	Postoperative	p value	Preoperative	Postoperative	p value
AHI	50.4±19.6	25.5±19.4	.002*	44.8±28.8	25.4±23.2	.005*
AI	34.3±20.3	13.6±16.7	.005*	28.2±26.7	14.7±23.8	.018*
ESS	11.6±4.6	7.6±3.5	< .001*	10.8±5.1	8.06±5.5	.017*
Min-SpO2	73.7±7.0	83.9±5.7	< .001*	74.0±10.0	80.6±12.5	.045*
CT90	16.4±15.0	5.7±7.8	.004*	13.6±16.1	8.8±19.3	.248

AHI = Apnea-Hypopnea index; AI = Apnea index; ESS = Epworth Sleepiness scale; Min-SpO2 = minimum oxygen saturation; CT90 = cumulative time percentage with SpO2 < 90%

† All values are presented as mean ± standard deviation

* p < .05 is considered statistically significant

Table 4(on next page)

Between-groups comparison of the treatment outcomes

SD = standard deviation; AHI = Apnea-Hypopnea index; AI = Apnea index; ESS = Epworth Sleepiness scale; Min-SpO2 = minimum oxygen saturation; CT90 = cumulative time percentage with SpO2 < 90%; NRS = numerical rating scale

TABLE 4.
Between-groups comparison of the treatment outcomes

	TORS group (n=16)	Coblation group (n=17)	p value
AHI reduction, mean \pm SD	48.0 \pm 38.9	45.3 \pm 34.1	.831
AI reduction, mean \pm SD	64.2 \pm 40.6	53.2 \pm 42.1	.481
ESS reduction, mean \pm SD	3.4 \pm 3.0	2.8 \pm 4.3	.646
Min-SpO2 improvement, %, mean \pm SD	10.2 \pm 7.9	6.6 \pm 12.5	.355
CT90 reduction, %, mean \pm SD	10.6 \pm 11.4	4.8 \pm 15.6	.268
Success rate, n (%)	8(50.0)	10(58.8)	.611
Day 1 pain score(NRS), mean \pm SD	2.7 \pm 0.8	2.5 \pm 0.7	.428
Hospital stay, days, mean \pm SD	5.5 \pm 1.1	4.3 \pm 0.7	.002
Major complication, n (%)	0(0)	0(0)	
Minor complication, n (%)	8(50.0)	6(35.3)	.393

SD = standard deviation; AHI = Apnea-Hypopnea index; AI = Apnea index; ESS = Epworth Sleepiness scale; Min-SpO2 = minimum oxygen saturation; CT90 = cumulative time percentage with SpO2 < 90%; NRS = numerical rating scale

Figure 1

The study flow diagram

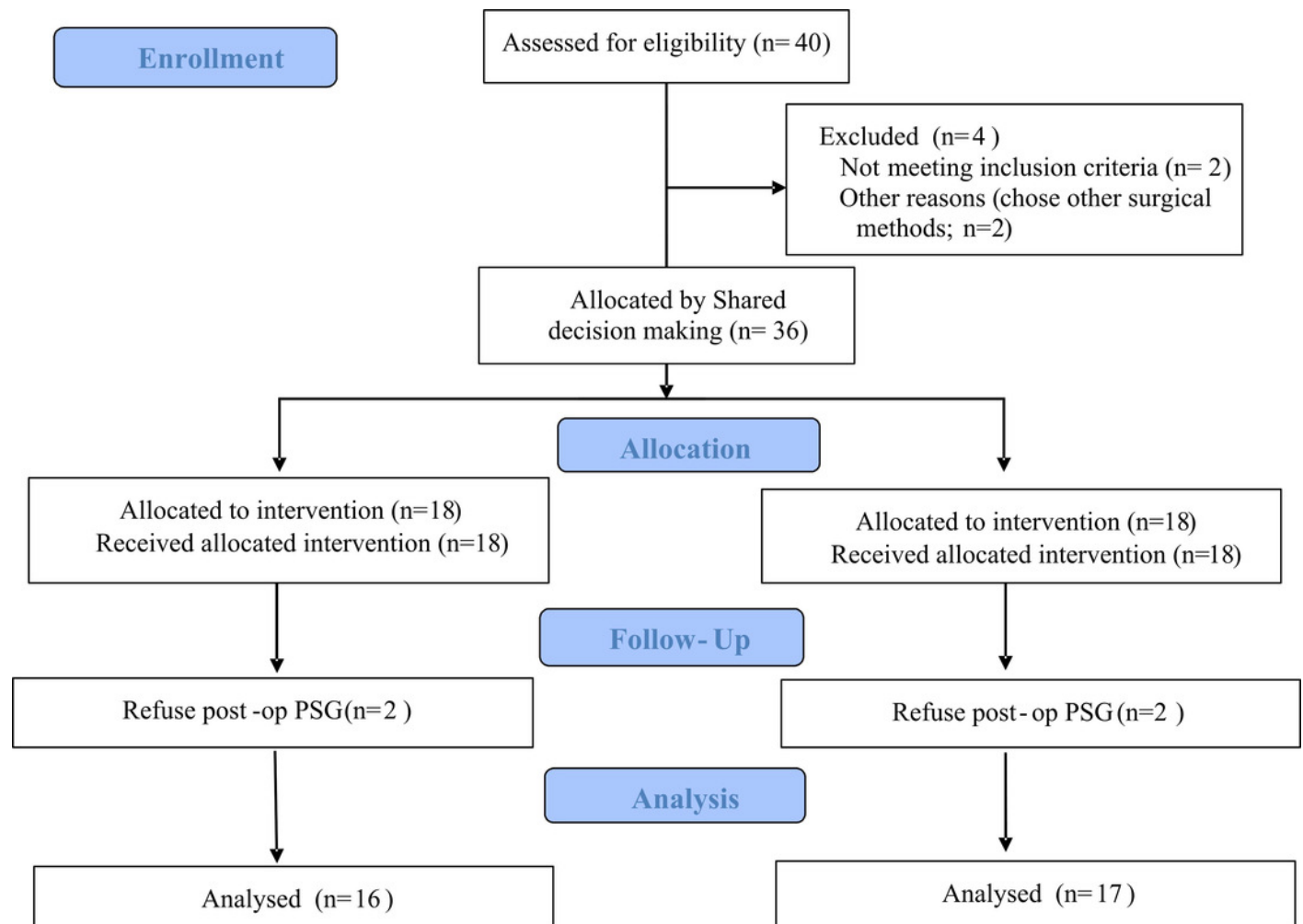


Figure 2

The treatment outcome of ESS between two groups

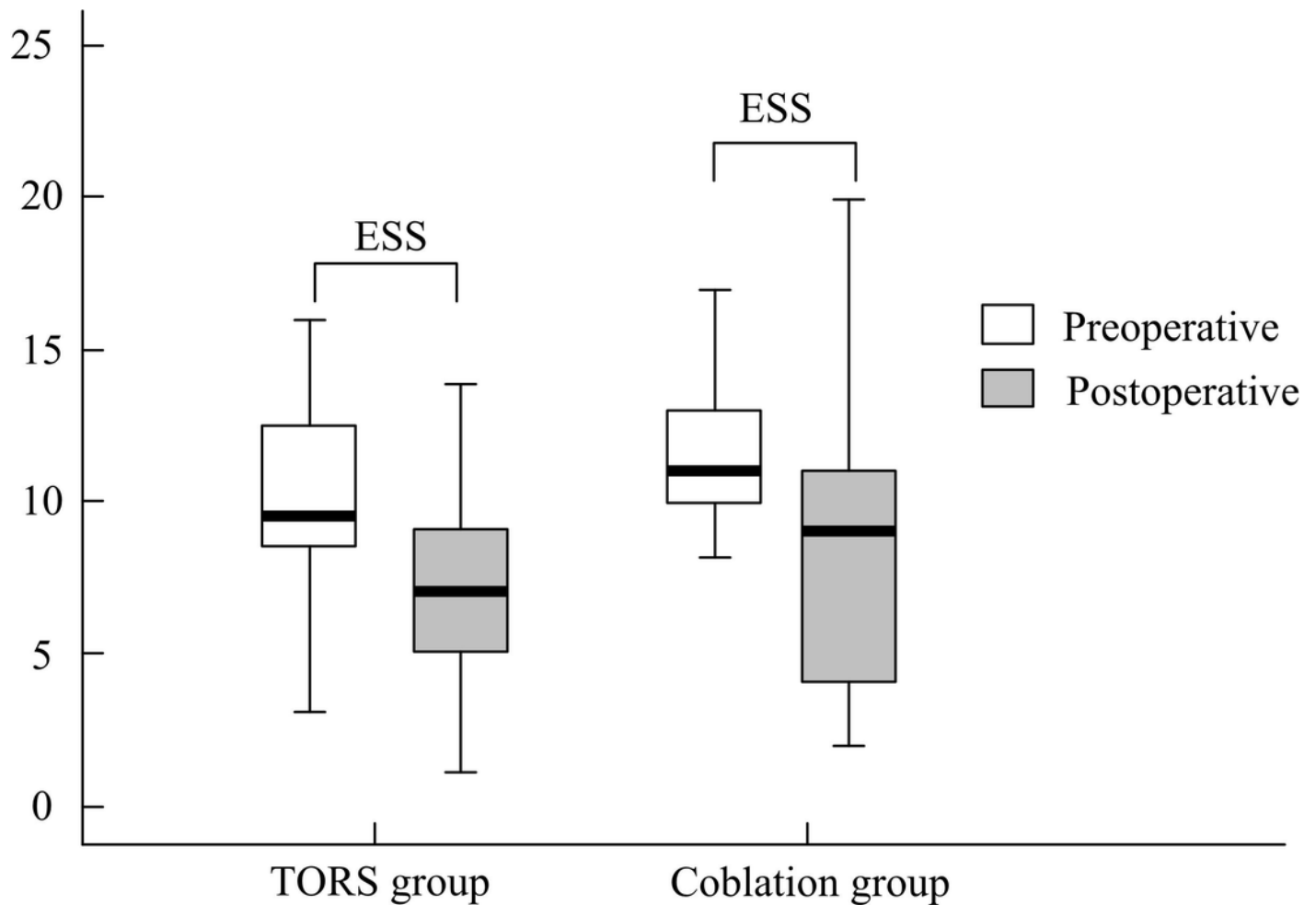


Figure 3

The treatment outcome of AHI and AI between two groups

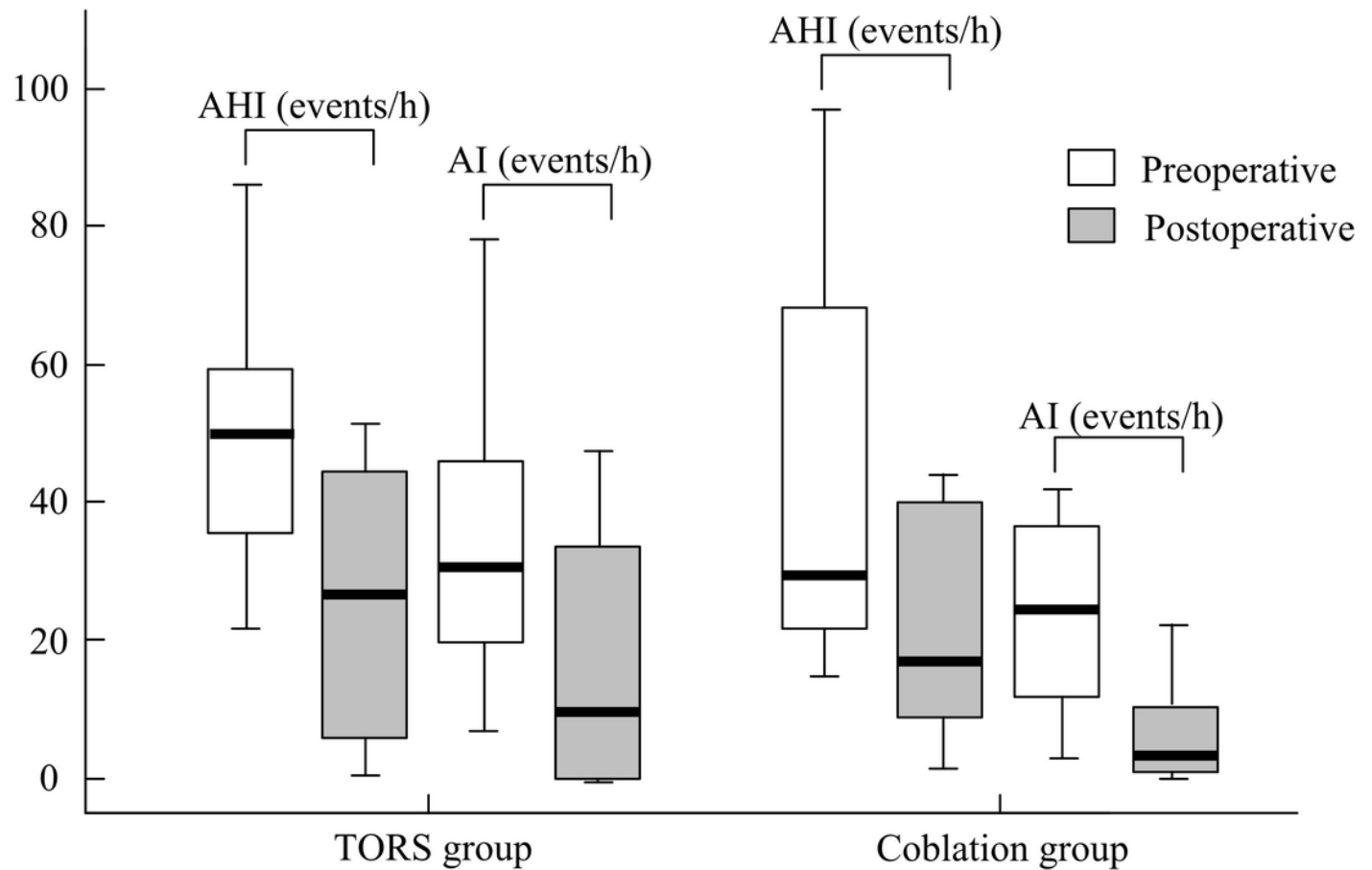


Figure 4

The treatment outcome of min-SpO2 and CT 90 between two groups

