

# 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  $48.0 \pm 38.9$  events/h and  $45.3 \pm 34.1$  events/h, respectively; the between-group difference was not significant ( $p = .831$ ). ESS improvement did not differ significantly between the TORS and coblation groups ( $3.4 \pm 3.0$  and  $2.8 \pm 4.3$ , 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.

1 **Trans-oral Robotic Surgery versus Coblation Tongue Base Reduction for**

2 **Obstructive Sleep Apnea Syndrome**

3

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19

20 **Abstract**

21 **Objectives.** To compare the efficacy of trans-oral robotic surgery (TORS) with that of  
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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27 tongue base reduction. Both groups received concomitant uvulopalatoplasty. Surgical  
28 outcomes were evaluated by comparing the initial polysomnography (PSG) parameters  
29 with the follow-up PSG data at least 3 months after the surgery. Epworth sleepiness scale  
30 (ESS) and complications were also compared between the 2 groups.

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  $48.0 \pm 38.9$  events/h and  $45.3 \pm 34.1$  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 ( $3.4 \pm 3.0$  and  $2.8 \pm 4.3$ , respectively,

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

43

#### 44 **Introduction**

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

55 and evaluative methods, multilevel surgery is becoming a more common method of  
56 successfully treating OSAS.<sup>7, 8</sup> Among these patients with multilevel collapse, the most  
57 frequently seen pattern was the concomitant collapse of palatal and tongue base  
58 (25.5%).<sup>6</sup> Uvulopalatopharyngoplasty (UPPP) is the most commonly reported surgery to  
59 address oropharyngeal obstruction. For dealing with tongue base obstruction, trans-oral  
60 robotic surgery (TORS) and coblation assisted tongue base reduction surgery were two  
61 of the most published tongue base tissue reduction procedures.

62 Several preoperative assessment strategies have been used. Friedman tonsil grading  
63 scale classifies the tonsil size into five grades (grade 0-IV) according to the location the  
64 tonsil relative to the surrounding structures.<sup>9</sup> Friedman tongue position (FTP) grading  
65 system is evaluated similarly to the modified Mallampati classification, but the tongue is  
66 evaluated in a neutral position without protrusion. Friedman staging system incorporates  
67 FTP, Friedman tonsil grading scale and BMI to classify OSAS patients into four stages:  
68 stage I includes patients with tonsils graded III-IV, FTP graded I-II and BMI<40 kg/M<sup>2</sup>;  
69 stage III includes patients with tonsils graded 0-II, FTP graded III-IV and BMI<40 kg/M<sup>2</sup>;  
70 stage IV includes patients with BMI>40 kg/M<sup>2</sup> or significant craniofacial or other anatomic  
71 abnormalities; stage II includes patients beyond stage I, III, IV.<sup>10</sup> Fiberoptic  
72 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.<sup>11</sup> 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.<sup>12</sup>

78 For most patients with oropharyngeal obstruction, uvulopalatopharyngoplasty (UPPP) is  
79 one of the most common and effective surgical procedures.<sup>13</sup> However, oropharyngeal  
80 obstruction combined with tongue base obstruction is recognized as the most important  
81 reason for failure after pharyngoplasty procedures.<sup>14</sup> 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.<sup>15</sup>

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.<sup>15</sup>

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.

91 <sup>16, 17</sup> However, there is a lack of fair comparison studies regarding the treatment efficacy  
92 and safety between TORS and coblation assisted tongue base reduction. Therefore, this  
93 study was conducted to compare the subjective and objective outcomes of TORS with  
94 endoscope-guided coblation tongue base reduction.

95

## 96 **Materials and methods**

97 Medical charts were retrospectively reviewed for OSAS patients admitted for TORS or  
98 coblation tongue base reduction surgery to a single tertiary hospital between 2012 and  
99 2017. Thirty three patients with age ranging from 18 to 62 years met the inclusion criteria  
100 (Table 1). Patients who were excluded were those without available postoperative  
101 polysomnography (PSG) data. PSG was performed at 3-12 months after the surgery.  
102 Patients who had previous upper airway surgery for OSAS were also excluded. This study  
103 was approved by the Institutional Review Board of the China Medical University Hospital  
104 (project approval number CMUH103-REC1-078).

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

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

111 The grades of airway collapse in Muller's maneuver were divided into four grades  
112 according to the percentage change in cross-sectional area: grade I  $\leq 25\%$  collapse,  
113 grade II  $>25\%$  and  $\leq 50\%$  collapse, grade III  $>50\%$  and  $\leq 75\%$  collapse, grade IV  
114  $>75\%$  collapse. VOTE classification was utilized for reporting DISE findings and the grade  
115 of collapse were classified as 0 ( $< 50\%$  obstruction); 1 (50–75% obstruction) and 2 ( $>$   
116 75% obstruction). Patients undergoing surgery must have at least partial tongue base  
117 collapse confirmed by Muller's maneuver and DISE. Details were also recorded from pre-  
118 operative and post-operative PSG data, and included AHI, apnea index (AI), lowest  
119 oxygen saturation (min-SpO<sub>2</sub>), cumulative time percentage with SpO<sub>2</sub>  $< 90\%$  (CT90)  
120 (Table 2). The success of the surgery was defined as achievement of  $\geq 50\%$  reduction of  
121 pre-operative AHI and a post-operative AHI  $< 20$ . Perioperative parameters, including the  
122 length of stay in hospital, the numeric rating scale (NRS) for pain intensity assessment  
123 on the first postoperative day and complications, were recorded.

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

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

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

129 was performed similar to the previous published literature.<sup>16, 18</sup> General anesthesia was

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

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

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

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

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

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

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

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

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

139 3D camera endoscope, tongue base tissue was grasped by robotic forceps and

140 cauterized with spatula monopolar electrode. The midline posterior glossectomy began

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

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

143 Endoscopic coblation assisted tongue base reduction surgery was performed similar to

144 previous reports.<sup>19, 20</sup> Under general anesthesia with nasotracheal intubation, the Molt

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

160

## 161 **Result**

162 Among the 33 patients in this analysis, 16 were in the TORS group (age of  $39.4 \pm 12.3$

163 years) and 17 patients were in the coblation group (age of  $38.7 \pm 11.5$  years). The male  
164 comprised 93.8% in the TORS group and 76.5% in the coblation group. The body mass  
165 index (BMI) at the time of admission was  $28.1 \pm 3.8$  kg/m<sup>2</sup> in the TORS group and  $27.3 \pm 5.5$   
166 kg/m<sup>2</sup> in the coblation group. There were no significant differences in tonsil grading scale,  
167 Friedman tongue position and Friedman staging system between the two groups before  
168 surgery. The grades of collapse in Muller's maneuver and DISE were similar in both  
169 groups. The Epworth sleepiness scale (ESS) was  $11.6 \pm 4.6$  in the TORS group and  
170  $10.8 \pm 5.1$  in the coblation group. All patients received polysomnography (PSG) for pre-  
171 operative evaluation. The baseline apnea-hyponea index (AHI) was  $50.4 \pm 19.6$  events/h  
172 and mean apnea index (AI) was  $34.3 \pm 20.3$  events/h in the TORS group; corresponding  
173 values were  $44.8 \pm 28.8$  events/h and  $28.2 \pm 26.7$  events/h, respectively, in the coblation  
174 group. The mean lowest oxygen saturation (min-SpO<sub>2</sub>) was  $73.7 \pm 7.0\%$  and mean  
175 cumulative time percentage with SpO<sub>2</sub> < 90% (CT90) was  $16.4 \pm 15.0\%$  in the TORS group;  
176 corresponding values were  $74.0 \pm 10.0\%$  and  $13.6 \pm 16.1\%$  in the coblation group.  
177 Demographics, baseline PSG data for both groups are summarized in Table 2. There  
178 were no significant between-group differences prior to treatment.  
179 The comparisons within-group (Table 3) and between-group (Table 4) were analyzed,  
180 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 ( $3.4 \pm 3.0$  and  $2.8 \pm 4.3$ , respectively,  $p = .646$ ; 95% CI =  $-3.30 \sim 2.08$ , Figure 2). The AHI  
183 reduced significantly from  $50.4 \pm 19.6$  events/h to  $25.5 \pm 19.4$  events/h in the TORS group  
184 ( $p = .002$ ). In the coblation group, the mean AHI reduced significantly from  $44.8 \pm 28.8$   
185 events/h to  $25.4 \pm 23.2$  events/h ( $p = .005$ ). The AHI reduction in the TORS and coblation  
186 groups were  $48.0 \pm 38.9$  events/h and  $45.3 \pm 34.1$  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-SpO<sub>2</sub> 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-SpO<sub>2</sub> 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-SpO<sub>2</sub> 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).

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

207

## 208 **Discussion**

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

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

217 pattern of the airway collapse.<sup>8, 17, 21</sup>

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  $45.7 \pm 21.7$  to  $12.8 \pm 8.2$  events/hour ( $p < .001$ ) postoperatively and the overall surgical  
222 success rate was 80%.<sup>20</sup> 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  $37.7 \pm 18.6$  to  $18.7 \pm 14.8$  events/hour  
225 ( $p < .001$ ) and the success rate was 55.6%.<sup>19</sup> 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%;  $p = .04$ ).<sup>17</sup>

230 O'Malley et al. developed a minimally invasive surgical procedure for management of  
231 tongue base neoplasms by using robotic surgical instruments.<sup>22</sup> 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

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

242 A study by Friedman et al. in 2012 was thought to be the first comparative study for the  
243 comparison of coblation and TORS in OSAS treatment.<sup>16</sup> It compared the effectiveness  
244 of TORS with that of coblation assisted submucosal minimally invasive lingual excision  
245 (SMILE). All the patients in the study received concomitant z-palatoplasty. The AHI  
246 reduction in the TORS and SMILE groups were  $60.5\% \pm 24.9\%$  and  $32.0\% \pm 43.3\%$  ( $p =$   
247  $.012$ ), respectively. The success rate in the TORS and SMILE groups were 66.7% and  
248 45.5%, respectively; the between-group difference was not significant ( $p = .135$ ).  
249 However, the techniques used by each group were different and not completely  
250 comparable.

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

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

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

266 As detailed in Table 4, the mean reduction of AHI, AI, ESS, CT90 and mean improvement  
267 of min-SpO<sub>2</sub> were similar for the TORS and coblation groups. The rate of surgical success  
268 in the TORS group were comparable to the coblation group (50.0% vs 58.8%, p= .611).  
269 Hwang et al. compared the tongue base coblation resection to TORS in OSAS patients  
270 and both groups were in combination with lateral pharyngoplasty.<sup>26</sup> 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/m<sup>2</sup> vs 27.3-28.1kg/m<sup>2</sup>). Moreover, preoperative mean ESS were  
275 lower (8.5-9.7 vs 10.8-11.6) and mean min-SpO<sub>2</sub> 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.<sup>15</sup>  
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 glossectomy<sup>27</sup>, lingual tonsillectomy<sup>28</sup>) and coblation (e.g. midline posterior

289 glossectomy<sup>20</sup>, SMILE<sup>29</sup>, channelling of the tongue<sup>30</sup>, Interstitial injections with  
290 needle coblation<sup>31</sup>). In the future, prospective, randomized, controlled trials that  
291 incorporate similar surgical technique will be needed to evaluate the efficacy of TORS  
292 compared with coblation tongue base reduction. Moreover, studies providing long-term  
293 results in the treatment of OSAS are also warranted.

294

## 295 **Conclusion**

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

300

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395

**Table 1** (on next page)

Inclusion criteria

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

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**Table 1:**  
Inclusion criteria

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≥ 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 retroglottal collapse in Muller's maneuver and DISE

Cannot tolerate CPAP

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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 groups†

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-SpO<sub>2</sub> = minimum oxygen saturation; CT90 = cumulative time percentage with SpO<sub>2</sub> < 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 <sup>2</sup>	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 <sub>2</sub> , %	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-SpO<sub>2</sub> = minimum oxygen saturation; CT90 = cumulative time percentage with SpO<sub>2</sub> < 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-SpO<sub>2</sub> = minimum oxygen saturation; CT90 = cumulative time percentage with SpO<sub>2</sub> < 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

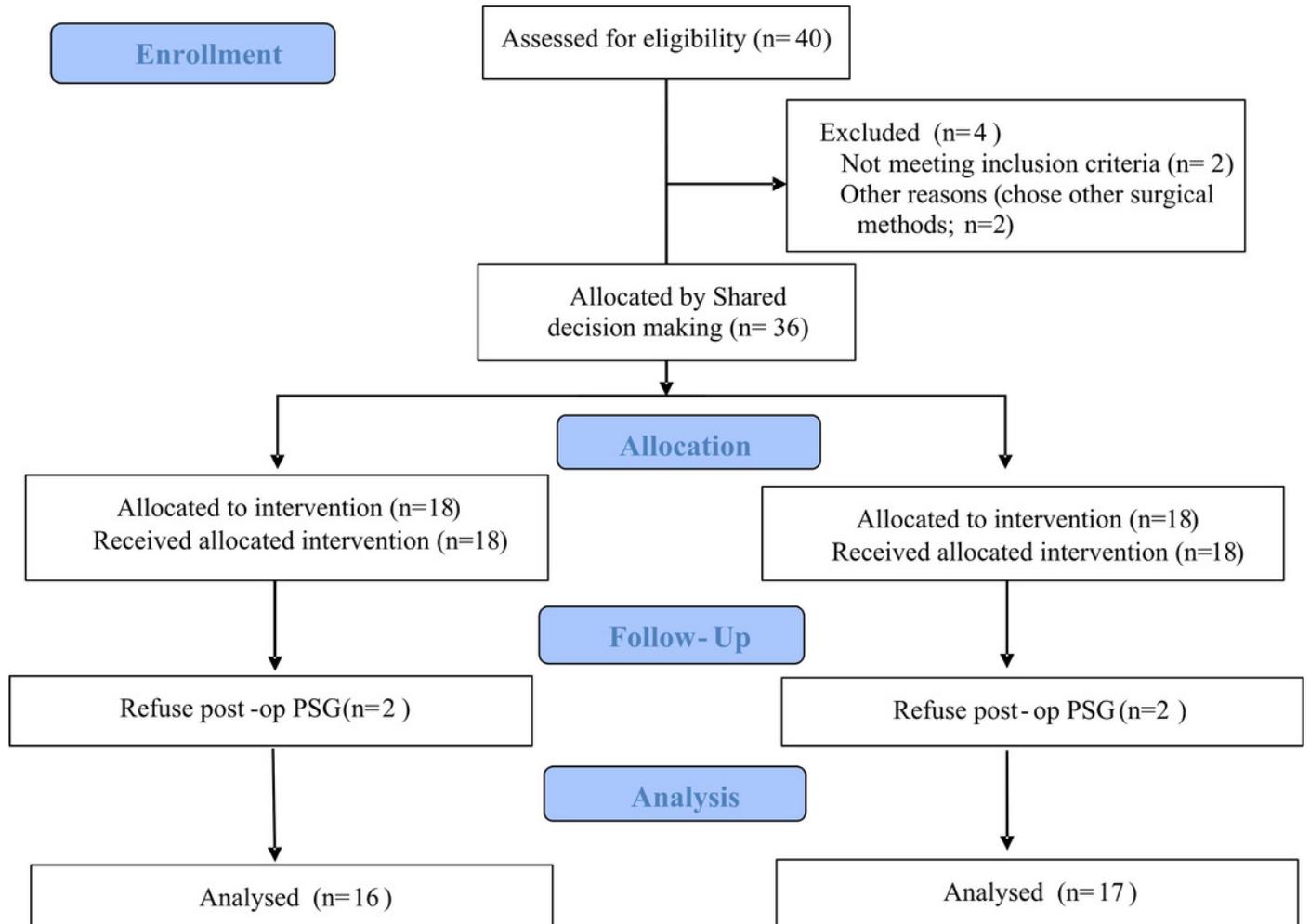
**TABLE 4.**  
Between-groups comparison of the treatment outcomes

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

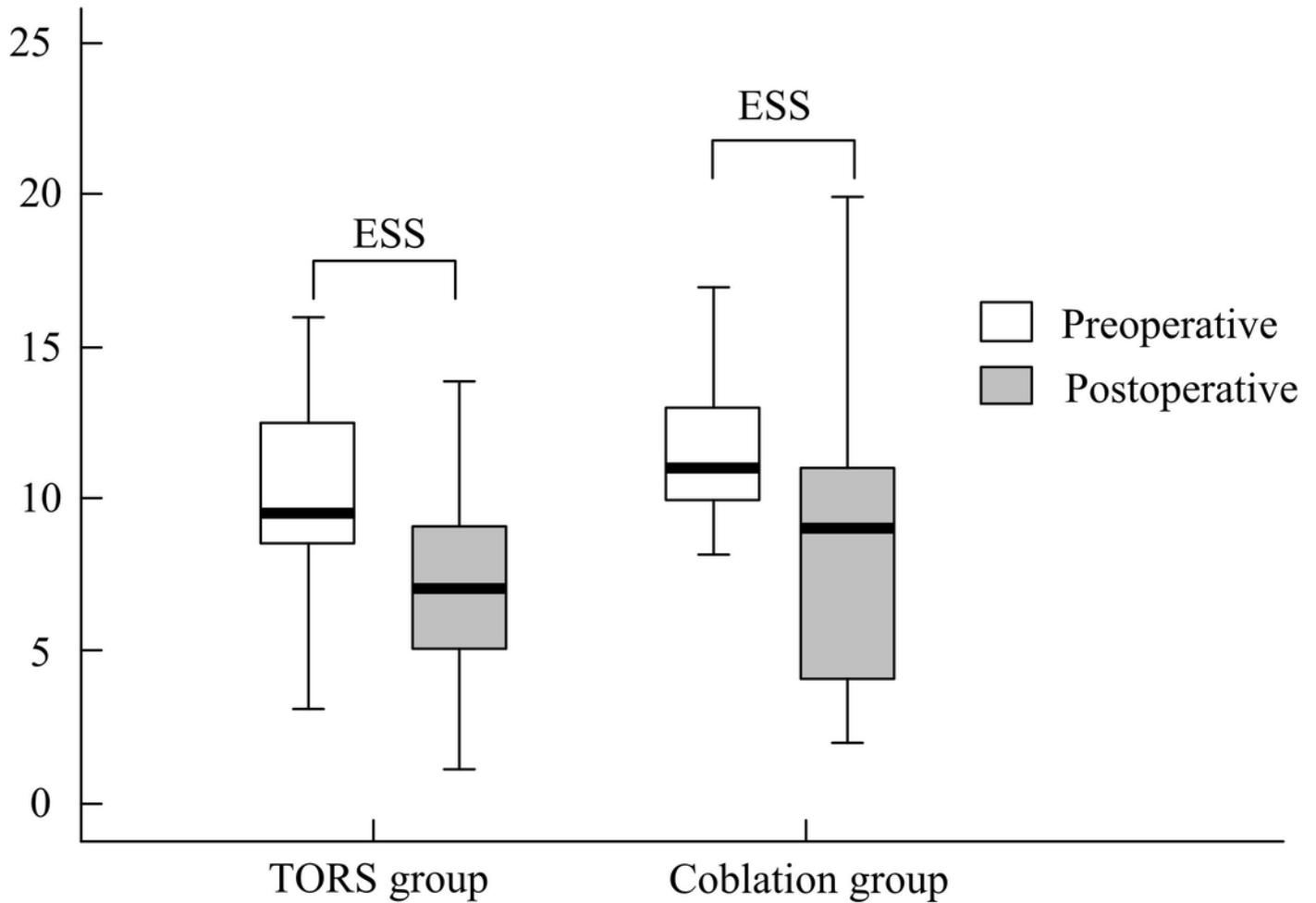
# Figure 1

The study flow diagram



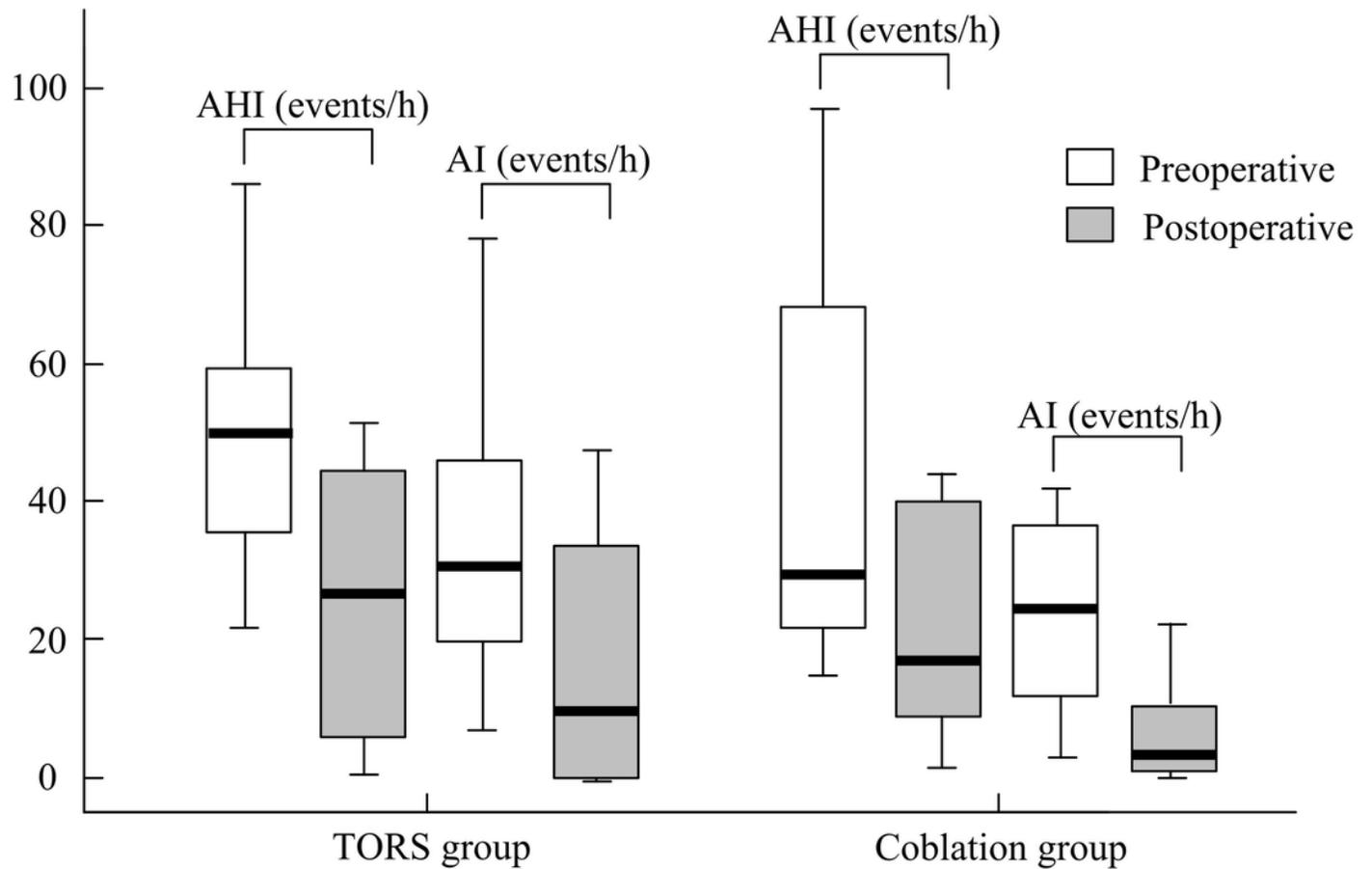
## Figure 2

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-SpO<sub>2</sub> and CT 90 between two groups

