Introduction

Tonsillectomy is one of the most common otolaryngological operations. It is performed for several conditions, including sleep disorders (e.g., snoring or apnea), recurrent tonsillitis, complications such as peritonsillar abscess, and tumors suspected to be malignant. However, postoperative pain after tonsillectomy may be severe enough to impede a patient’s normal academic or working activities and compromise his or her nutritional status. Among the methods proposed to reduce postoperative pain are various forms of surgery using different types of equipment, local anesthesia before surgery, steroids, antibiotics, analgesics, and cold-water cooling. Nonetheless, a highly reliable and effective method for reducing postoperative pain has yet to be identified.

A non-steroidal anti-inflammatory drug (NSAID) mouthwash has been used in dentistry to reduce pain following periodontal surgery or pain caused by periodontitis, but it has been rarely used in otolaryngological procedures. Thus, we...
conducted a randomized, controlled study to evaluate the efficacy of NSAID mouthwash in reducing post-tonsillectomy pain.

**Subjects and Method**

The patients enrolled in this study were 15 years of age or older and underwent a tonsillectomy between January and November 2016 at Soonchunhyang University Bucheon Hospital. The indication for surgery was recurrent or chronic tonsillitis. Patients who underwent additional procedures such as uvulopharyngopalatoplasty, those with significant medical morbidities or an active tonsillar infection, patients who declined to participate, and those who did not complete the questionnaire were excluded from the study. Informed consent was obtained from all participants. This study was approved by the Institutional Review Board of Soonchunhyang University (IRB No. 2016-014).

The demographics of the 64 patients who were included in the study are listed in Table 1. This study was performed as single-blind, and the patients were randomly assigned to experimental (NSAID) and control (distilled water) groups (n=32 patients per each). Randomization was performed according to the randomized list, which was constructed using a computer-generated randomization sequence. All surgical procedures were performed by the same surgeon (J.Y.L.) with the same methods under general anesthesia. After adequate exposure of the oral cavity with a McIvor tongue retractor, each tonsil was dissected in the extracapsular plane using monopolar electrocautery with a coagulation mode of 20 W. Hemostasis was achieved by minimal spot electrocautery using bipolar forceps.

All patients were blinded as to whether they received NSAID (Diclomed®, 0.074% solution; Valeas S.P.A. Industria Chimica E Farmaceutici, Milano, Italy) or distilled water mouthwash postoperatively. Distilled water mouthwash was manufactured under the consultation with nutrition and pharmacy departments. It had similar color with NSAID mouthwash, but did not contain any harmful or pharmacological properties. The patients in both groups rinsed their mouth with 15 mL of the respective mouthwash twice a day for 30 sec each time, morning and evening, for 9 days after the operation. All of the patients were instructed to allow the solution to reach the tonsillar fossa. When they had difficulties in rinsing the tonsillar fossa, a spray bottle was provided.

After the operation, intravenous analgesics were administered to all patients (75 mg of diclofenac sodium mixed with 100 mL of 0.9% normal saline) at a dose interval of 8 h; oral analgesics were administered once every 8 h. Patients with unusually severe pain were administered additional diclofenac sodium via an intramuscular injection once or twice during their hospital stay. Patients in both groups were instructed to frequently rinse their mouth with ice water and 30X-diluted Povidone iodide® (7.5%; Firson Co., Ltd., Cheonan, Korea) mouthwash. Most of the patients were discharged on the first postoperative day (POD); the average hospital stay was 1.3 days. Upon discharge, all patients were prescribed a 1-week course of oral antibiotics, analgesics (acetaminophen and trimadole, 3 times a day), and either the NSAID mouthwash or distilled water mouthwash. They were also instructed to take oral acetaminophen if they experienced severe pain, even while taking the prescribed analgesics. Postoperatively, the patients received a questionnaire and were asked to record their subjective pain level on PODs 1–9 using a visual analog scale (VAS). A score of 0 indicated no pain or disturbances and a score of 10 unbearable pain. They were instructed to check the pain level in the morning 30 min after using

| Table 1. Demographics and variables of the patients |
|----------------------------------|--------|-----------------|-----------------|-----------------|--------|
| Variables                        | NSAID (n=32) | DW (n=32) | Total (n=64) | p value        |
| Average age (year)               | 32.5    | 29.6          | 31.1          |                 |
| Sex (%)                          |         |                |                |                 |
| Male                             | 15 (46.9) | 18 (56.3) | 33 (51.6) |                 |
| Female                           | 17 (53.1) | 14 (43.7) | 31 (48.4) |                 |
| Additional analgesics (%)        |         |                |                | <0.001*         |
| No                               | 26 (81.2) | 10 (31.3) | 36 (56.3) |                 |
| Yes                              | 6 (18.8) | 22 (68.7) | 28 (43.7) |                 |
| Post-operative bleeding (%)      |         |                |                | 1.000           |
| No                               | 30 (93.8) | 30 (93.8) | 60 (93.8) |                 |
| Yes                              | 2 (6.2)  | 2 (6.2)  | 4 (6.2)   |                 |

*statistically different between NSAID group and distilled water group. NSAID: non-steroidal anti-inflammatory drug, DW: distilled water
mouthwash. Completed questionnaires were collected at the patients’ first or second outpatient visit after discharge.

Statistical analyses were conducted using SPSS (version 14.0; SPSS Inc., Chicago, IL, USA) and R (version 3.1.3; The R Foundation for Statistical Computing, Vienna, Austria) software; all analyses were based on two-tailed tests. A p value < 0.05 was considered to indicate statistical significance.

Results

All patients who participated in the study completed the questionnaire. Except on POD 1, the pain level on PODs 2–9 was significantly lower in the NSAID mouthwash group than in the distilled water mouthwash group (Table 2). Five patients in the NSAID group reported scores of 9–10 on the VAS, indicating extreme pain, compared to 21 in the control group. These pain scores were usually recorded on PODs 3–4. In the NSAID group, pain increased until POD 3 but showed a decreasing tendency from POD 4 onwards. In the distilled water group, patients reported a decrease in pain beginning on POD 5 (Fig. 1). Two patients in each group experienced minor bleeding (Table 1).

As shown in Table 1, the two groups did not significantly differ in other variables except for the use of additional analgesics, which were taken by 6 patients in the experimental group and 22 in the control group. The significant difference between the groups with respect to their VAS scores remained even after statistical adjustment for the additional medications, which is shown in the last row of Table 2 (*). The linear regression analysis was used for the statistical adjustment.

Discussion

Although tonsillectomy is one of the most commonly performed procedures among otolaryngologists, postoperative pain can lead to morbidities such as dysphagia, dehydration, and local bleeding. In addition, severe pain may extend the hospital stay. While there is no standard method to effectively control postoperative pain following tonsillectomy, among the most common methods are the use of opiates or NSAID analgesics before surgery, local anesthetics administered to the surgical site preoperatively or postoperatively, and the intravenous injection of analgesics after surgery.4–8 The efficacy of several mouthwashes has been evaluated, but their effects on pain reduction have yet to be definitively determined.9,11,12 Diclomed® mouthwash is based on diclofenac sodium, a strong NSAID derived by choline diclofenac salification.

Table 2. Postoperative changes of VAS score

<table>
<thead>
<tr>
<th>VAS score</th>
<th>NSAID (n=32)</th>
<th>DW (n=32)</th>
<th>p value</th>
<th>p value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Postoperative 1-day</td>
<td>4.53 ± 2.23</td>
<td>5.81 ± 2.32</td>
<td>0.058</td>
<td>0.062</td>
</tr>
<tr>
<td>Postoperative 2-day</td>
<td>5.12 ± 2.47</td>
<td>6.62 ± 1.95</td>
<td>0.009*</td>
<td>0.015*</td>
</tr>
<tr>
<td>Postoperative 3-day</td>
<td>5.38 ± 2.21</td>
<td>7.47 ± 1.68</td>
<td>&lt; 0.001*</td>
<td>0.002*</td>
</tr>
<tr>
<td>Postoperative 4-day</td>
<td>4.69 ± 2.26</td>
<td>8.22 ± 1.54</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Postoperative 5-day</td>
<td>4.44 ± 2.15</td>
<td>8.25 ± 1.41</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Postoperative 6-day</td>
<td>4.09 ± 2.13</td>
<td>7.62 ± 1.39</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Postoperative 7-day</td>
<td>3.94 ± 2.21</td>
<td>7.38 ± 1.26</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Postoperative 8-day</td>
<td>3.44 ± 2.37</td>
<td>6.81 ± 1.55</td>
<td>&lt; 0.001*</td>
<td>&lt; 0.001*</td>
</tr>
<tr>
<td>Postoperative 9-day</td>
<td>3.12 ± 2.49</td>
<td>6.08 ± 1.74</td>
<td>&lt; 0.001*</td>
<td>0.019*</td>
</tr>
</tbody>
</table>

Data are given as mean ± standard deviations. *Statistically different between NSAID group and distilled water group, which demonstrates less post-tonsillectomy pain in NSAID group. †p values were obtained after adjustment for additional analgesics. The linear regression analysis was used for the statistical adjustment. VAS: visual analogue scale, NSAID: non-steroidal anti-inflammatory drug, DW: distilled water.
This modification allows rapid absorption of the diclofenac in the NSAID mouthwash by the oropharyngeal mucosa and thus a reduction in inflammation and pain by inhibiting cyclooxygenase enzyme and local prostaglandin production. Analgesic effect is related to its peripheral anti-inflammatory effect and probably by weakening the pain threshold under the thalamus cortex, then the pain is decreased or interrupted.\(^{10}\) This agent was also shown to inhibit the DNA synthase of Gram-positive and Gram-negative bacteria, including Pseudomonas aeruginosa and Escherichia coli.\(^{10,11,15}\) In the present study, we used this NSAID mouthwash to assess the effects of NSAID mouthwash versus distilled water mouthwash in the control of pain following tonsillectomy.

The accurate absorption percentage through the mucosa and half-life of NSAID mouthwash have not been clearly defined yet. In contrast with systemic NSAID, exact measurement of these parameters would be virtually difficult in topical agent. However, more rapid onset of analgesic effect can be anticipated because of direct contact and absorption through the oropharyngeal mucosa. According to the manufacturer’s pharmacokinetics, diclofenac was detectable within 10 min after contact with oral mucosa and peak concentration was reached within 45 min, with an average maximal concentration of 21.8±5.4 ng/mL. In addition, the systemic bioavailability of diclofenac solution after 7 days of mouth rinsing twice a day was 1/20–1/50 of that obtained with oral administration of 25 mg diclofenac tablet. However, these data are not formally published.

The lack of a significant difference in the VAS scores of the two groups on POD 1, immediately after surgery, may have been related to several factors, including throat discomfort caused by endotracheal intubation and swelling of the oral cavity mucosa due to the surgical placement of a tongue retractor. On PODs 2–9, the VAS scores of the two groups differed significantly and were better in the NSAID mouthwash group.

Two patients in each group experienced a minor episode of bleeding, but it was easily controlled with ice water gargling and did not necessitate a clinical visit or emergency evaluation.

The use of additional analgesics was significantly higher in the control group. However, the significance of their VAS scores remained even after statistical adjustment for the additional medications, which was better in the experimental group. These results demonstrate that the NSAID mouthwash significantly reduced postoperative pain compared to a distilled water mouthwash even in patients taking additional analgesics.

Potential limitations of this study are relatively small number of subjects in each group and the data which are drawn only by subjective symptom scores of the patients. Postoperative use of analgesics may be another limitation. The most ideal way to evaluate the effect of NSAID mouthwash is to use the agent in the experimental group only without administration of analgesics in both groups. However, this can induce intolerable pain in the control or even in the experimental group, and patients may demand pain control. Therefore, we provided basic analgesics in both groups after operation. We excluded pediatric patients because they cannot be relied on to clearly express their degree of pain or to use a mouthwash effectively to reach the tonsillar fossa. We also did not compare the NSAID mouthwash with other agents, but in Korea there are only a few commercially available mouthwashes (e.g., chlorhexidine gluconate and hydrogen peroxide) and their main purpose is to cleanse the surgical site and prevent bacterial or fungal infection. For these reasons, we restricted our comparison of the NSAID mouthwash to distilled water mouthwash.

In conclusion, the NSAID mouthwash significantly and quickly reduced postoperative pain compared to a distilled water mouthwash, thus providing a simple and effective method to relieve post-tonsillectomy pain when used in conjunction with routine medications. The postoperative use of an NSAID mouthwash may help patients achieve an earlier return to normal daily life.

Acknowledgments
This work was supported by the Soonchunhyang University Research Fund.

REFERENCES