Tonsillotomy versus Tonsillectomy for Chronic Recurrent Tonsillitis in Children

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Abstract

Objective: This study was conducted to compare two different surgical methods; partial tonsil resection using the bipolar technique, tonsillotomy (TT), and total tonsillectomy (TE) (blunt dissection) for recurrent tonsillitis in children. The frequency of recurrent throat infections was determined during postoperative follow-up.

Methods: A total of 393 patients were included in this study. TT was performed on 174 patients (100 males, 74 females) and TE on 219 patients (112 males, 107 females). Following surgery, an analysis was made of treatment outcomes of patients with upper respiratory tract infections. The patients were followed up for 12-48 months. Their parents were also contacted by telephone to determine the frequency of reinfection and their satisfaction with the surgery. Also, parents of the patients completed a questionnaire pertaining to postoperative satisfaction (scale of 1-10).

Results: In the TT group, 14 (8.1%) patients had recurrent tonsillitis postoperatively. In the TE group, 12 (5.4%) patients required antibiotherapy due to recurrent pharyngitis after the surgery. There was no significant difference between the two groups in terms of the one-year infection recurrence rate after surgery (p=0.281). Three patients (1.6%) in the TT group and 12 (5.4%) in the TE group complained of bleeding within the first 24 hours. The rate of bleeding was significantly lower in the TT group than the TE group (p=0.001). There were no fatalities in either group.

Conclusion: In both groups, the rate of reinfection accorded with the requirement for postoperative antibiotics. For recurrent tonsillitis, TT was as effective as TE. However, TT was superior in terms of the risk of bleeding.

Keywords: Recurrent tonsillitis, tonsillotomy, tonsillectomy, bipolar tonsillotomy

Introduction

Tonsil surgery is one of the most common surgical interventions performed by ear, nose, and throat (ENT) specialists. However, differences between therapeutic techniques, including in terms of indications, are still under investigation. Although the number of cases indicated for surgery due to infection has decreased, a large proportion of cases still have an infectious origin (1, 2). Many cases of tonsillitis can be treated with antibiotics, but surgery is the first line treatment in patients with recurrent or chronic tonsillitis who do not respond to medical treatment. Various techniques have been applied in tonsil surgery. The disadvantage of the tonsillotomy (TT) technique is that the tonsils can regrow to serve as a source of infection. Therefore, many previous studies concluded that TT was not suitable for patients with chronic tonsillitis (3). The main advantage of the TT technique is the preservation of the tonsil capsule, which serves as a biological dressing, covering the pharyngeal muscles such that they are not directly exposed to trauma, or to inflammation arising from contact with saliva. Thus, both postoperative pain and healing time are reduced (4, 5). In general, ENT physicians prefer total tonsillectomy (TE) to TT for patients with frequent episodes of tonsillitis. In recent years, however, TT has been performed in patients with a history of frequent infections. But still, studies on the application of TT, which is a safe method for treating cases with frequent episodes of tonsillitis, are insufficient. The present study was performed...
to compare the frequency of postoperative upper respiratory tract infection in patients undergoing TE versus TT.

**Methods**

The study was carried out with approval from the Ethics Committee of Van Training and Research Hospital of the University of Health Sciences (approval number 2018/07, granted on December 4, 2018). This retrospective cohort study reviewed the consecutive medical records of all children undergoing tonsil surgery at our tertiary center. In this study, 1,254 patients who underwent tonsillar surgery at our center from June 2014 through July 2018 were evaluated retrospectively in terms of surgical indications and approach. Only patients with chronic and/or recurrent tonsillar infection, and at least one year follow-up, were evaluated. A total of 393 patients were included in this study. The indications were determined according to Key Action Statement 2-3 of the current guidelines for TE in children (6). Patients with obstructive sleep apnea syndrome, continuous snoring, nasal congestion, abnormal dentofacial/orofacial development, recurrent/chronic adenoiditis, recurrent or chronic otitis media with effusion, recurrent or chronic sinusitis with adenoid hypertrophy, or suspicion of malignancy, were excluded from the study. Patients undergoing paracentesis and tube insertion were also excluded, along with those with immunodeficiency and hematological disorders or syndromes. Only patients who underwent surgery because of complaints of frequent tonsillitis were included in the study.

The patients were divided into two groups. The first group underwent general anesthesia for bipolar electrocautery (15 W), TT and bleeding control. The tonsillar tissue was cauterized using bayonet nonbonding bipolar forceps. The ForceTriad Energy Platform (Medtronic-Covidien, Inc., Minneapolis, MN, USA) was used as a generator (240 V, 50 Hz). The cauterization power was fixed at 15 W. TT was performed on the anterior plica line of each tonsil by cauterizing the surface several times. The full thickness of the tonsil was cauterized from the superior to the inferior direction, perpendicular to the level of the plica. The medial portion of the cauterized area was extracted by cutting with scissors (Figure 1).

The second group underwent cold dissection TE under general anesthesia with using sutures for the control of bleeding. We used vertical and horizontal sutures (4-0 braided absorbable polyglactin 910 sutures; Vicryl®, Ethicon, Somerville, NJ, USA). During the postoperative period, patients with a diagnosis of recurrent throat infections were identified from among the electronic National Health Service hospital records. Patients who received antibiotic therapy at our center, or treatment for symptoms of upper respiratory tract infection from other physicians (especially pediatricians and family physicians), were identified from among the National Health Service data. After surgery, an analysis was made of the treatment outcomes of patients with upper respiratory tract infection. The patients were followed up for 12–48 months. During the same period, their parents were also contacted by telephone to determine the frequency of reinfection and their satisfaction with the surgery (Table 1). Patient’s parents also completed a questionnaire pertaining to postoperative satisfaction (scale of 1–10).

**Statistical Analysis**

Descriptive statistics were used to analyze continuous (quantitative) variables. Data are expressed as the mean ± standard deviation (SD). Categorical variables are expressed as numbers and percentages. The sample size required for each group was calculated with the Collins sampling calculation formula. For an estimated type 1 error rate of 5% (z=1.96) and power of 80%, it was calculated that each group should consist of at least 123 patients. However, to increase the power, it was planned that each group would contain at least 174 patients. The independent t-test was used to compare the mean scores of the groups on parametric continuous variables. The chi-square test was used to analyze categorical variables. The z-ratio test was used to compare proportions. Minitab (version 17.0; Minitab Inc., State College, PA, USA) and the IBM Statistical Package for Social Sciences software for Windows version 24.0 (IBM SPSS Corp.;
Armonk, NY, USA) were used for the analyses, in all of which p<0.05 was taken to indicate statistical significance.

Results
A total of 393 patients [212 males (53.9%) and 181 females (46.1%)] were included in this retrospective study. All patients were treated for chronic or recurrent tonsillitis. TT and TE were performed in 174 and 219 patients, respectively. The mean age of the patients undergoing TT for recurrent tonsillitis was 8.03 years (range: 4-15 years), while that of the patients undergoing TE was 7.4 years (range: 4-15 years). The mean follow-up period was 24.84 months (range: 12-48 months) in the TT group and 21.49 months (range: 12-48 months) in the TE group.

In the TT group, the mean number of times antibiotic therapy was prescribed for recurrent tonsillitis per year was 5.89 pre-operatively and 1.02 after surgery. Furthermore, 14 (8.1%) TT group patients had frequent tonsillitis postoperatively, of whom four underwent reoperation (TE) because of frequent tonsillitis. The remaining 10 patients did not accept repeat surgery despite frequent tonsillitis; they had fewer tonsillitis episodes than pre-operatively and preferred antibiotic therapy.

In the TE group, the mean number of times antibiotic therapy was prescribed for recurrent tonsillitis per year was 5.74 pre-operatively, which decreased to 0.64 postoperatively. In the TE group, 12 (5.4%) patients required antibiotic therapy due to recurrent pharyngitis after surgery. The antibiotics prescribed for patients diagnosed with tonsillitis or pharyngitis are shown in Figure 2.

Three patients (1.6%) in the TT group exhibited bleeding during the first 24 hours after surgery. In patients who presented with bleeding in the postoperative tonsillar region, the clots therein were cleared and treatment was provided in the form of a cold-water gargle and cotton impregnated with pano-tocaine–adrenaline. In one patient, bleeding continued despite treatment, so TE was performed under general anesthesia. In the remaining two patients, bleeding was self-limited by medical method. In the TE group, 12 patients (5.4%) presented with bleeding. Eight of these patients were taken to the operating room and sutured again under general anesthesia. In the remaining four patients, bleeding was self-limited by medical methods. There were no fatal complications in either group. The rate of bleeding was significantly lower in the TT group than the TE group (p=0.001), while there was no significant difference in mean postsurgical satisfaction evaluation score between the groups (Table 2).

Discussion
The indications, efficacy, necessity, risks, and benefits of tonsil surgery have been discussed in previous reports. Can TT replace TE in cases requiring tonsil surgery? The Paradise et al. (5) criteria for TE can be applied for assessment of clinical status and interpretation of examination results, and to guide the follow-up of patients scheduled for tonsil surgery. Patients with frequent tonsil infections should be evaluated clinically for up to one week for fever, painful cervical lymphadenopathy, and bacterial lymphadenopathy, which may indicate bacterial infec-

Table 2. Patient characteristics: comparison between the tonsillotomy and tonsillectomy groups

|                        | Tonsillotomy group (TT), n=174 | Tonsillectomy group (TE), n=219 |
|------------------------|-------------------------------|---------------------------------|
| Gender (male/female)   | 100/74                        | 112/107                         |
| Age (years)            | 8.03±(2.53)                   | 7.4±(2.73)                      |
| Average follow-up period (months) | 24.84±(10)               | 21.49±(9,9)                     |
| Number of patients with postoperative bleeding (%) | 3 (1.6)                  | 12 (5.4)                        |
| Number of patients requiring surgery after postoperative bleeding | 1                          | 8                               |
| Mean number of antibiotic prescriptions due to tonsillitis in the year before surgery | 5.89±(1.14)             | 5.74±(1.15)                     |
| Number of recurrent pharyngitis or tonsillitis episodes in the year after surgery (%) | 14 (8.1)                  | 12 (5.4)                        |
| Mean number of antibiotics prescriptions for tonsillitis or pharyngitis in the year after surgery | 1.02±(1.58)             | 0.64±(0.97)                     |
| Number of patients requiring revision surgery during follow-up | 4                          | –                               |
| Average family satisfaction score after surgery (scale of 1–10) | 7.66±(0.96)             | 7.77±(0.99)                     |

Data are presented as n (%) for categorical variables. Data are presented as mean ± SD or median (min, max) for continuous variables. p<0.05 was taken to indicate statistical significance.

Figure 2. Antibiotics used in the treatment of tonsillitis. "Others" included ampicillin, amoxicillin, cefaclor, ampicillin + sulbactam, and azithromycin.
Bleeding has been reported at a rate of 0.1–20% after TE, with a mortality ratio due to bleeding of 1/1,000–1/170,000. Some studies reported that postoperative bleeding was the most common cause of mortality following TE. Chronic or recurrent acute infections result in more bleeding due to destruction of the tonsil capsule and adhesion to the surrounding tissue (13, 14). Vicini et al. (15) compared TE and TT groups in terms of bleeding (a total of 450 cases). The rate of postoperative bleeding was 6.8% in the TE group and 0.7% in the TT group, and it was concluded that TT was safer and more effective than TE. In the present study, the rate of postoperative bleeding was 1.6% in the TT group and 5.4% in the TE group; again, indicating that TT was much safer than TE in terms of bleeding. Howev-

ergic long-term risks should be taken into consideration in the decision-making process regarding whether to perform TE or adenoidectomy.

This retrospective cohort study included children treated with one of two different surgical techniques for tonsil infections. A major limitation of this study was the absence of histological and bacteriological data.

Conclusion
All organs and tissues in the human body are crucial. We found that TT was safer than, and equally effective as, TE, and can therefore be recommended for the management of recurrent tonsillitis during childhood.

Ethics Committee Approval: Ethics committee approval was received for this study from the Ethics Committee of Van Training and Research Hospital of the University of Health Sciences (approval number 2018/07, granted on December 4, 2018).

Informed Consent: Informed consent was obtained from the parents of all participants included in the study.

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