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ORIGINAL RESEARCH

Cancer Medicine WILEY

Transoral surgery for superficial head and neck cancer: National Multi-Center Survey in Japan

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Abstract

Head and neck cancers, especially in hypopharynx and oropharynx, are often detected at advanced stage with poor prognosis. Narrow band imaging enables detection of superficial cancers and transoral surgery is performed with curative intent. However, pathological evaluation and real-world safety and clinical outcomes have not been clearly understood. The aim of this nationwide multicenter study was to investigate the safety and efficacy of transoral surgery for superficial head and neck cancer. We collected the patients with superficial head and neck squamous cell carcinoma who were treated by transoral surgery from 27 hospitals in Japan. Central pathology review was undertaken on all of the resected specimens. The primary objective was effectiveness of transoral surgery, and the secondary objective was safety including incidence and severity of adverse events. Among the 568 patients, a total of 662 lesions were primarily treated by 575 sessions of transoral surgery. The median tumor diameter was 12 mm (range 1-75) endoscopically. Among the lesions, 57.4% were diagnosed as squamous cell carcinoma in situ. The median procedure time was 48 minutes (range 2-357). Adverse events occurred in 12.7%. Life-threatening complications occurred in 0.5%, but there were no treatment-related deaths. During a median follow-up period of 46.1 months (range 1-113), the 3-year overall survival rate, relapse-free survival rate, cause-specific survival rate, and larynx-preservation survival rate were 88.1%, 84.4%, 99.6%, and 87.5%, respectively. Transoral surgery for superficial head and neck cancer offers effective minimally invasive treatment.

Clinical trials registry number: UMIN000008276.

KEYWORDS

head and neck cancer, larynx preservation, pharyngeal cancer, superficial cancer, transoral surgery

1 | INTRODUCTION

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Worldwide incidence and mortality of oropharyngeal cancer are reported as 92,887 and 51,005 and those of hypopharyngeal cancer are 80,608 and 34,984 in 2018.¹ The prognosis is still poor even with multimodal treatment because most patients have locally advanced disease with lymph node involvement at the time of diagnosis and have a propensity for developing distant metastasis.^{2,3}

The standard treatment for resectable oro- and hypopharyngeal cancer is laryngopharyngectomy with pharyngeal reconstruction, leading to a loss of natural speech and a difficulty of swallowing.⁴ An alternative treatment is chemoradiotherapy, which can preserve organ and function. However, it often caused serious adverse effects, such as dysphagia, due to severe mucositis and xerostomia, negatively affecting patients' quality of life.⁵

The ideal approach to improve the patients' survival and to preserve organ and function is early detection of cancer and applying minimally invasive treatment.⁶ Tumor located within the epithelium and subepithelial layer was categorized as superficial cancer.⁷ Muto et al. reported that narrow band imaging (NBI: Olympus Co., Ltd.) enabled virtual chromoendoscopy and early detection of superficial head and neck cancer.⁸ Then, NBI is now widely used in clinical practice in many countries.⁹⁻¹⁵

For superficial lesions, endoscopic mucosal resection (EMR) and endoscopic submucosal dissection (ESD) have been indicated and showed effectiveness.¹⁶⁻²⁴ Recently, transoral video-assisted surgery (TOVS) and endoscopic laryngopharyngeal surgery (ELPS) have also been indicated.²⁵⁻²⁹ Together, EMR, ESD, TOVS, and ELPS are classified as transoral surgery (TOS). While TOS has been widely indicated for superficial head and neck cancer, their pathological evaluation is not standardized. Then, the clinical management after TOS is not also standardized. In addition, the real-world effectiveness and safety of TOS for superficial head and neck cancer have not been well defined. We, therefore, conducted a national multi-center survey of TOS in Japan.

2 | MATERIALS AND METHODS

2.1 | Participants

Patients who were primarily treated by TOS from April 2001 through July 2012 were retrospectively collected from 27 hospitals in Japan. The inclusion criteria were as follows: (a) tumors pathologically diagnosed as squamous cell carcinoma (SCC), (b) tumors invasion was pathologically limited within subepithelial layer, (c) no exposure of tumor cells to the vertical margin (negative vertical margin), (d) macroscopic tumor location in the oropharynx, hypopharynx, or supraglottis, (e) no regional lymph node metastasis on computed tomography, and (f) no other active advanced cancer in the head or neck region.

Written informed consent was obtained from all patients for the procedures in this study. Patients with concomitant primary cancer in any other organ were excluded. If cancers in other organs have been curatively treated when initial TOS was indicated, the patients were included. This study was approved by ethics committees in all participating hospitals and was registered in UMIN Clinical Trials Registry (UMIN000008276).

2.2 | Transoral surgery (TOS)

TOS is defined as a procedure or an operation to perform mucosectomy for which a surgical device and visual guidance are inserted from mouth. Ablation procedure is not included in TOS. EMR and ESD were mainly performed by gastroenterologists. Others were mainly performed by head and neck surgeons.

2.3 | Outcomes/Survey variables

The primary objective was effectiveness of TOS, and the secondary objective was safety including incidence and severity of adverse events. The survey variables were as follows: (a) the clinicopathological characteristics of patients with superficial SCC of head and neck, (b) adverse events associated with TOS, (c) incidences of local recurrence, regional lymph node recurrence, and distant recurrence after TOS and subsequent treatments, (d) incidence of and treatment regimen for metachronous cancer, and (e) the survival data on follow-up duration (overall survival, relapse-free survival, cause-specific survival, and larynx-preservation survival after TOS).

2.4 | Histopathological analysis

One certified pathologist (S.F.) performed centralized pathology review of registered patients and excluded the patients without SCC, those with SCC with muscularis propria invasion, and those with histologic cancer types other than SCC. As a second step, 10 certified pathologists developed a new set of diagnostic criteria to distinguish subepithelial invasive SCC from SCC *in situ* for this study. The criteria were as follows; at least one solitary nest of epithelial neoplastic cells is present in the stroma clearly separated from intraepithelial carcinoma or intraepithelial carcinoma with a thickness of 500 µm or greater. As a third step, we conducted a central pathological review board by three certified pathologists (M.F., T.N., and M.I.). This board defined the presence or absence of invasion blinded to the clinical findings according to the diagnostic criteria. Consensus decision making was used to make final pathological diagnosis.

2.5 | Local, regional lymph node and distant recurrence

Local recurrence was defined as a development of tumor at the treatment site of TOS. Regional lymph node and distant recurrence were defined as an abnormal enlargement of lymph node and a new lesion in distant location detected on the computed tomography, respectively.

2.6 | Metachronous cancer

Metachronous cancer was defined as cancer detected in the region after initial TOS that was clearly separate from the resection scar. Metachronous cancer in other organs was defined as cancer arising in organs other than the head and neck after initial TOS.

2.7 | Statistical analysis

p-values for categorical data were calculated by using Kruskal-Wallis test for trends in the median procedure times and using Fisher's exact test for other variables related to the safety of transoral surgery, respectively. Overall survival rates, relapsefree survival rates, and cause-specific survival rates were estimated using the Kaplan–Meier method and tested by log-rank tests. Cumulative incidence of metachronous head and neck cancers, metachronous cancers arising in other organs, and larynx-preservation survival that events were laryngectomy and all death were estimated using the Kaplan–Meier method. We defined the time to the development of a metachronous cancer as the period from the day of TOS to the day of diagnosis of a metachronous cancer. All data were analyzed with SAS (version 9). All authors had access to the study data and have reviewed and approved the final manuscript.

3 | RESULTS

3.1 | Participants

A total of 599 patients with superficial head and neck cancer (700 lesions) were registered. We excluded 12 patients (14 lesions) with no available pathological specimens and 10 patients (11 lesions) with inadequate follow-ups. The specimens

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of the remaining 577 patients (675 lesions) were carefully screened by one certified pathologist per protocol. This screening excluded 4 patients (4 lesions) with non-cancerous lesions, 2 patients (2 lesions) with SCC invasion to muscular layer, 2 patients (5 lesions) with insufficient specimen to evaluate pathological findings, and 1 patient (2 lesions) with a tumor of a histologic type other than SCC (spindle cell carcinoma). Finally, a total of 568 patients (662 lesions) were included in the analysis. The total number of TOS sessions for 568 patients was 575 because 7 other sessions for synchronous lesions were performed on another day (Figure 1).

Table 1 shows the demographic characteristics of the study patients. The median age was 66 years (range 33 to 89), and 534 (94.0%) of the subjects were men. The performance status was 0 in 539 patients (94.9%). The most common reason for the detection of superficial head and neck cancer was endoscopic examination before or after the treatment of esophageal cancer (366 patients, 64.4%). The total number of patients with previous head and neck cancer and history of other cancer was 141 and 531 (includes overlapping patients), respectively. Among the 531 patients with previous cancer, 416 (78.3%) had history of esophageal cancer. Treatment for these cancers was summarized in Table 2.

Table 3 shows the demographic characteristics of the treated lesions. Among the 662 lesions, 519 (78.4%) were located in the hypopharynx and 132 lesions (19.9%) were located in the oropharynx. The most common macroscopic types was flat (528 lesions, 79.8%). The procedures for TOS were EMR (307 lesions, 46.2%), ESD (264 lesions, 39.7%), ELPS (31 lesions, 4.7%), and TOVS (31 lesions, 4.7%). A total of 490 lesions (74.0%) underwent en bloc resection. The median tumor diameter was 12 mm (range 1-75) endoscopically and 14 mm (range 1-60) pathologically. The median diameters of the resected tumor specimens in EMR, ESD, ELPS, TOVS, and other procedures were 12, 15, 20, 16, and 13 mm, respectively (ranges: 1-45, 1-60, 2-58, 5-42, and 3-50 mm, respectively). Three hundred and eighty lesions (57.4%) were revealed to be intraepithelial SCC based on the central pathological review on the depth of invasion. The T categories were found to be Tis (380 lesions, 57.4%), T1 (181 lesions, 27.3%), T2 (89 lesions, 13.4%), T3 (11 lesions, 1.7%), and unknown (1 lesion, 0.2%). Subsequent treatment was performed immediately after initial TOS for 20 lesions (3.0%).

3.2 | Adverse events

Among the 575 treatment sessions, most of the procedure was underwent under general anesthesia (545 sessions, 94.8%). The median procedure time was 48 minutes (range 2–357). EMR was performed in a short time (32 minutes, p < 0.0001). Adverse events occurred in 12.7% (73/575). The main adverse events were laryngeal edema (33 sessions, 5.7%),

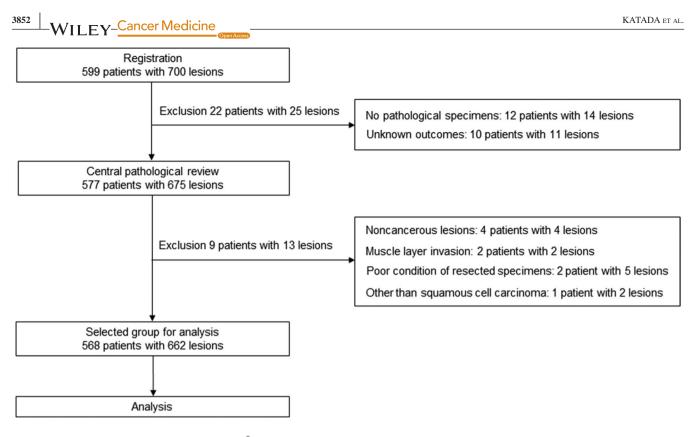


FIGURE 1 Flow chart of patients and lesions. *Seven other sessions for synchronous lesions were performed on another day

subcutaneous emphysema (20 sessions, 3.5%), aspiration pneumonia (14 sessions, 2.4%), and bleeding (11 sessions, 1.9%). Subcutaneous emphysema frequently occurred in ESD (6.3%, p < 0.0178). Temporary tracheotomy was performed in 49 treatment sessions (8.5%). The main reasons for tracheotomy were development of laryngeal edema (22 sessions, 3.8%) and perioperative planned management (21 sessions, 3.7%). There were no treatment-related deaths; however, 3 patients (0.5%) developed life-threatening severe adverse events. Those were as follows: 1) one patient underwent an emergency tracheotomy because of suffocation caused by laryngeal edema after surgery, 2) one patient underwent an emergency tracheotomy because of arterial bleeding and hemostasis was achieved by ligation of the blood vessels, and 3) one patient had transient cardiopulmonary arrest caused by aspiration of food during a meal on the following day and recovered after removal of the foreign object through a tracheostoma (Table 4).

3.3 | Local, regional lymph node and distant recurrence

Median follow-up period was 46.1 months (range 1–113). Recurrence data and their treatment were summarized in Table 5. Among 662 lesions treated by TOS, 53 lesions (8.0%) developed local recurrence. Local recurrence rates of EMR, ESD, and other procedures were 11.7% (35/298), 2.7% (7/258), and 10.4% (11/106), respectively (p < 0.0001). The median diameters of the resected tumor specimens which

developed local recurrence and specimens that did not develop local recurrence were 16 (range: 3-45 mm) and 14 mm (range: 1-60 mm), respectively. There was no relation between tumor size and local recurrence (p = 0.13). Thirtynine lesions (73.6%) were treated by re-TOS. Traditional open surgery with and without laryngectomy were performed in 3 lesions (5.7%) and 2 lesions (3.8%), respectively. Remaining 9 lesions (17.0%) were treated with non-surgical treatment. Regional lymph node recurrence developed in 26 patients (4.6%). Among them, 20 patients (76.9%) developed on the same side of the neck. Radical neck dissection was performed in 15 patients (57.7%), and 8 patients (30.8%) received neck dissection plus postoperative chemotherapy and/or radiotherapy. Three patients (11.5%) received definitive chemoradiotherapy. Three patients (0.5%) had distant recurrence; two had lung metastasis and remaining one had lung and liver metastasis. Two patients (66.7%) were followed up without any treatment, and 1 patient (33.3%) received chemotherapy.

3.4 | Metachronous cancer

A total of 234 metachronous head and neck cancers were diagnosed in 132 patients (23.2%) during the follow-up period. The 3-year cumulative incidence rate of metachronous head and neck cancers after TOS was 16.7% (95% confidence interval, 13.7% to 20.2%) (Figure 2A). Among 234 lesions, 207 (88.5%) were again treated by TOS. Traditional open surgery

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TABLE 1 Patient characteristics	
Total number of patients	568
Main factor leading to detection	
Before/after treatment of esophageal cancer	366 (64.4%)
Before/after treatment of head and neck cancer	83 (14.6%)
Medical checkups	55 (9.7%)
Pharyngolaryngeal paresthesia	49 (8.6%)
Before/after treatment of gastric cancer	15 (2.6%)
Age, median (range)	66 (33–89)
Sex (male)	534 (94.0%)
Performance status (0/1/2/3/4)	539 (94.9%) / 22 (3.9%) / 6 (1.1%) / 1 (0.2%) / 0 (0.0%)
History of head and neck cancer	
Total number of previous head and neck cancers ^a	141
Hypopharynx	43 (30.5%)
Oral cavity	38 (27.0%)
Larynx	31 (22.0%)
Oropharynx	25 (17.7%)
Primary unknown	2 (1.4%)
Maxilla	2 (1.4%)
History of cancer in other organ	
Total number of previous cancers ^a	531
Esophageal cancer	416 (78.3%)
Gastric cancer	74 (13.9%)
Colorectal cancer	12 (2.3%)
Prostate cancer	9 (1.7%)
Lung cancer	5 (0.9%)
Liver cancer	3 (0.6%)
Breast cancer	2 (0.4%)
Skin cancer	2 (0.4%)
Bladder cancer	2 (0.4%)
Malignant lymphoma	2 (0.4%)
Bile-duct cancer	1 (0.2%)
Thyroid cancer	1 (0.2%)
Duodenal cancer	1 (0.2%)
Anal canal cancer	1 (0.2%)

^aIncluding overlapping patients.

with and without laryngectomy were performed in 1 lesion (0.4%) and 4 lesions (1.7%), respectively. Other 20 lesions (8.5%) were treated with non-surgical treatment and the treatment details of 2 lesions (0.9%) were not available (Table 5).

A total of 131 metachronous cancers arising in other organ were diagnosed in 96 patients (16.9%) during the follow-up period. The 3-year cumulative incidence rate of metachronous cancers arising in other organs after TOS was 14.7% (95% CI, 11.9% to 18.0%) (Figure 2B). And treatment for these cancers were summarized in Table 6. Esophagus was the main sites of metachronous cancer arising in other organ. Among 90 metachronous esophageal cancers, endoscopic resection was performed in 74 lesions (82.2%), surgery-based treatment in 11 lesions (12.2%), and chemoradiation-based treatment in 5 lesions (5.6%).

3.5 | Survival

During a median follow-up period of 46.1 months (range 1–113), 3 patients died of superficial head and neck cancer because of 2 distant metastasis and 1 local lymph node

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	Total number of previous cancers ^a	Surgery-based therapy	Endoscopy-based therapy	Chemoradiation -based therapy	Chemotherapy or hormonal therapy	Observation	TACE/RFA	Unknown
Head and neck cancer	141	75	0	60	5	1	0	0
Esophageal cancer	416	120	213	80	0	2	0	1
Gastric cancer	74	37	34	1	2	0	0	0
Colorectal cancer	12	6	2	0	1	0	0	0
Prostate cancer	6	4	0	1	4	0	0	0
Lung cancer	5	3	1	1	0	0	0	0
Liver cancer	c	1	0	0	0	0	2	0
Bladder cancer	2	1	0	1	0	0	0	0
Breast cancer	2	2	0	0	0	0	0	0
Skin cancer	2	1	1	0	0	0	0	0
Malignant lymphoma	2	1	0	1	0	0	0	0
Bile-duct cancer	1	1	0	0	0	0	0	0
Duodenal cancer	1	1	0	0	0	0	0	0
Anal canal cancer	1	0	0	1	0	0	0	0
Thyroid cancer	1	1	0	0	0	0	0	0
Total	672	257	251	146	12	3	2	1
Abbreviations: RFA, rad	Abbreviations: RFA, radiofrequency ablation; TACE, transcatheter arterial chemoembolization.	anscatheter arterial chemo	embolization.					

TABLE 2 Treatment history of head and neck cancer and cancer in other organ

^aIncluding overlapping patients.

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TABLE 3 Lesion characteristics	
Total number of lesions	662
Tumor location	
Oropharynx	132 (19.9%)
Anterior wall / Posterior wall / Lateral wall / Superior wall	9/79/23/21
Hypopharynx	519 (78.4%)
Postcricoid / Pyriform sinus / Posterior wall	33/404/82
Larynx	7 (1.1%)
Laryngeal epiglottis / Laryngeal arytenoid / Aryepiglottic folds	4/2/1
Oral cavity	4 (0.6%)
Oral floor / Hard palate / Buccal mucosa	1/1/2
Macroscopic type	
Flat / Elevated / Unknown	528 (79.8%) / 127 (19.2%) / 7 (1.1%)
Treatment methods	
Endoscopic mucosal resection (EMR)	307 (46.2%)
Endoscopic submucosal dissection (ESD)	264 (39.7%)
Endoscopic laryngopharyngeal surgery (ELPS)	31 (4.7%)
Transoral videolaryngoscopic surgery (TOVS)	31 (4.7%)
Laser microlaryngeal surgery	17 (2.6%)
Direct mucosectomy	12 (1.8%)
Number of resected specimens	
En bloc	490 (74.0%)
Piecemeal	172 (26.0%)
Number of segments obtained by piecemeal resection	
2/3/4/5/6/7/8/9/10/11	85/39/13/11/10/7/3/1/2/1
Tumor diameter on endoscopic images, median (range) ^a	12 (1–75)
Tumor diameter of resected specimens, median (range) ^b	14 (1–60)
EMR / ESD / ELPS / TOVS / Other procedures	12 (1-45) / 15 (1-60) / 20 (2-58) / 16 (5-42) / 13 (3-50)
Endoscopic depth of invasion for resected lesions	
Intraepithelial / Subepithelial / Difficult to evaluate	472 (71.0%) / 158 (23.8%) / 32 (4.8%)
Histopathological depth of invasion (central diagnosis)	
Intraepithelial / Subepithelial	380 (57.4%) / 282 (42.6%)
T category	
Tis / T1 / T2 / T3 / Unknown	380 (57.4%) / 181 (27.3%) / 89 (13.4%) / 11 (1.7%) / 1 (0.2%)
Lymphatic invasion	19 (2.9%)
Venous invasion	16 (2.4%)
Horizontal margin positive for cancer in the resected specimen	309 (46.7%)
Subsequent treatment immediately after initial transoral surgery	20 (3.0%)
^a Missing data for 20 nations	

^aMissing data for 29 patients.

^bMissing data for 1 patient.

metastasis and 25 patients died of metachronous cancer arising in other organ. Specific sites of cancer among those 25 patients were esophageal cancer in 8 patients, lung cancer in 6 patients, colorectal cancer in 3 patients, gastric cancer in 2 patients, liver cancer in 2 patients, bile duct cancer in 2 patients, duodenal cancer in 1 patient, and ureteral cancer in 1 patient. The 3-year overall survival rate (Figure 2C) was 88.1% (95% CI, 85.0% to 90.6%), the 3-year relapse-free survival rate (Figure 2D) was 84.4% (95% CI, 81.0% to 87.3%), the 3-year cause-specific survival rate (Figure 2E) was 99.6% (95% CI, 98.5% to 99.9%), and the 3-year larynx-preservation survival rate (Figure 2F) was 87.5% (95% CI, 84.3% to 90.1%).

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TABLE 4 Variables related to the safety of transoral surgery

Total number of treatment57526322290
sessions
Methods for anesthesia 0.0353
General anesthesia545 (94.8%)242 (92.0%)214 (96.4%)89 (98.9%)
Intravenous anesthesia 29 (5.0%) 20 (7.6%) 8 (3.6%) 1 (1.1%)
None 1 (0.2%) 1 (0.4%) 0 (0.0%) 0 (0.0%)
Procedure time, median (range), 48 (2–357) 32 (2–240) 60 (15–357) 71 (6–300) <0.0001 min ^a
Adverse events 73 (12.7%) 28 (10.6%) 36 (16.2%) 9 (10.0%) 0.1399
Laryngeal edema 33 (5.7%) 18 (6.8%) 13 (5.9%) 2 (2.2%) 0.2593
Subcutaneous emphysema 20 (3.5%) 5 (1.9%) 14 (6.3%) 1 (1.1%) 0.0178
Aspiration pneumonia 14 (2.4%) 4 (1.5%) 9 (4.1%) 1 (1.1%) 0.1704
Bleeding 11 (1.9%) 5 (1.9%) 5 (2.3%) 1 (1.1%) 0.923
Stenosis 3 (0.5%) 1 (0.4%) 2 (0.9%) 0 (0.0%) 0.7573
Cerebral infarction 2 (0.3%) 1 (0.4%) 1 (0.5%) 0 (0.0%) 1
Dermatitis caused by iodine 1 (0.2%) 1 (0.4%) 0 (0.0%) 0 (0.0%) 1
Tooth injury 1 (0.2%) 0 (0.0%) 0 (0.0%) 1 (1.1%) 0.1565
Mediastinitis 1 (0.2%) 0 (0.0%) 1 (0.5%) 0 (0.0%) 0.5426
Temporary tracheotomy 49 (8.5%) 22 (8.4%) 22 (9.9%) 5 (5.6%) 0.4964
Reason for tracheotomy
$\begin{array}{ccc} \text{Development of laryngeal} & 22 (3.8\%) & 17 (6.5\%)^{\text{b}} & 4 (1.8\%) & 1 (1.1\%) & 0.0113 \\ \text{edema}^{\text{b}} & \end{array}$
$ \begin{array}{ccc} \text{Perioperative planned} & 21 (3.7\%) & 6 (2.3\%)^{\text{b}} & 12 (5.4\%) & 3 (3.3\%) & 0.1869 \\ & \text{management}^{\text{b}} \end{array} $
Difficulty for intraoperative 2 (0.3%) 0 (0.0%) 1 (0.5%) 1 (1.1%) 0.1453 bleeding management 2 (0.3%) 0 (0.0%) 1 (0.5%) 1 (1.1%) 0.1453
Unknown 5 (0.9%) 0 (0.0%) 5 (2.3%) 0 (0.0%) 0.0243
Life-threatening severe adverse 3 (0.5%) 1 (0.4%) 2 (0.9%) 0 (0.0%) 0.7573 event
Treatment-related death 0 (0%) 0 (0%) 0 (0%) —

^aMissing data for 21 patients.

^bOne overlapping patient.

Survival was analyzed based on the depth of invasion (carcinoma in situ vs. cancer with subepithelial invasion) and results were compared. The 3-year overall survival rates (Figure 2G), the 3-year relapse-free survival rates (Figure 2H), and the 3-year cause-specific survival rates (Figure 2I) were 88.2% (95% CI, 83.7% to 91.4%) vs. 88.4% (95% CI, 83.6% to 91.8%) (p = 0.47), 88.1% (95% CI, 83.7% to 91.4%) vs. 80.1% (95% CI, 74.6% to 84.6%) (p = 0.002), and 100% vs. 99.6% (95% CI, 97.1% to 99.9%) (p = 0.055), respectively.

Survival based on the T category was analyzed. The 3year overall survival rates of Tis, T1, T2, and T3 tumors (Figure 2J) were 88.2% (95% CI, 83.7% to 91.4%), 92.2% (95% CI, 86.7% to 95.5%), 79.3% (95% CI, 68.4% to 86.8%), and 100% (p = 0.037), respectively. The 3-year relapse-free survival rates of Tis, T1, T2, and T3 tumors (Figure 2K) were 88.1% (95% CI, 83.7% to 91.4%), 84.7% (95% CI, 78.0% to 89.5%), 71.2% (95% CI, 59.8% to 79.9%), and 81.8% (95% CI, 44.7% to 95.1%) (p < 0.0001), respectively. The 3-year cause-specific survival rates of Tis, T1, T2, and T3 tumors were 100%, 98.7% (95% CI, 94.9% to 99.7%), 100%, and 100% (p = 0.068), respectively.

4 | DISCUSSION

This is the first report of national multi-center survey of TOS for superficial head and neck cancer based on the standardized pathological evaluation. During a median follow-up period
 TABLE 5
 Recurrence, metachronous head and neck cancer, and their treatment after transoral surgery

then treatment after transorar surgery	
Local recurrence ($n = 662$ lesions)	53 (8.0%)
Treatment for recurrent lesions	
Transoral surgery	39 (73.6%)
Traditional open surgery	5 (9.4%)
With laryngectomy	3 (5.7%)
Without laryngectomy	2 (3.8%)
Observation	3 (5.7%)
Definitive chemoradiotherapy	2 (3.8%)
Radiotherapy	2 (3.8%)
Argon plasma coagulation	1 (1.9%)
Laser ablation	1 (1.9%)
Regional lymph node recurrence (n = 568 patients)	8 26 (4.6%)
Location of recurrent lesions	
Only same side	20 (76.9%)
Only opposite side	2 (7.7%)
Both sides	2 (7.7%)
Unknown	2 (7.7%)
Treatment for recurrent lesions	
Neck dissection	15 (57.7%)
Neck dissection + postoperative chemotherapy	3 (11.5%)
Neck dissection + postoperative radiotherapy	3 (11.5%)
Neck dissection + postoperative chemoradiotherapy	2 (7.7%)
Definitive chemoradiotherapy	3 (11.5%)
Distant recurrence ($n = 568$ patients)	3 (0.5%)
Location of recurrent lesions	
Lung	2 (66.7%)
Lung + Liver	1 (33.3%)
Treatment for recurrent lesions	
Chemotherapy	1 (33.3%)
Observation	2 (66.7%)
Metachronous head and neck cancer $(n = 568 \text{ patients})$	132 (23.2%) with 234 lesions
Treatment for metachronous lesions	
Transoral surgery	207 (88.5%)
Traditional open surgery	5 (2.1%)
With laryngectomy	1 (0.4%)
Without laryngectomy	4 (1.7%)
Argon plasma coagulation	9 (3.8%)
Radiotherapy	6 (2.6%)
Observation	3 (1.3%)
Definitive chemoradiotherapy	1 (0.4%)
Chemotherapy	1 (0.4%)
Unknown	2 (0.9%)

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of 46.1 months (range 1–113), the 3-year overall survival rate and the 3-year cause-specific survival rate was 88.1% (95% CI. 85.0% to 90.6%) and 99.6% (95% CI, 98.5% to 99.9%), respectively. There was no treatment-related death.

The most important clinical benefit of TOS was that it could preserve organ and function sparing patients from potentially devastating adverse events of radial surgery or chemoradiation. In this study, a total of 53 local recurrence (8.0%) developed after completion of TOS. However, 39 recurrent lesions (73.6%) were treated by re-TOS. As for regional lymph node recurrence, most of the patients were treated by radical neck dissection or radical neck dissection plus chemotherapy and/or radiotherapy. Only 4 patients (0.7%) underwent laryngectomy (3 for local recurrence and 1 for metachronous head and neck cancer). Therefore, 99.3% (564/568) of the patients overall enjoyed preservation of organ and function. Calculated 3-year larynx-preservation survival rate was very high at 87.5%.

The local recurrence rate was significantly lower in the *en* bloc resection (5.3%) than in the piecemeal resection (15.7%, p < 0.0001). Previous studies reported that the size of tumors that can be resected *en* bloc by EMR is limited and that EMR tends to have a higher rate of local recurrence.^{30,31} In this study, *en* bloc resection rate of EMR was 55.4% for median tumor size of 14 mm. Because the average size of *en* bloc resected specimens by EMR is 10.3 ± 6.1 mm, EMR may be suitable for small lesions if *en* bloc resection can be performed.

The most frequent adverse event was laryngeal edema. Temporary tracheostomy was indicated in 49 (8.5%) of 575 treatment sessions. Among them, 22 procedures (44.9%, 22/49) directly attributed to laryngeal edema and 2 procedures (0.3%) were due to difficulty for intraoperative bleeding management. In contrast, 21 procedures (42.9%, 21/49) were indicated for the planned tracheostomy to avoid airway obstruction potentially caused by laryngeal edema, bleeding, or aspiration after TOS even in the cases with absence of intraoperative adverse events. However, the indication for planned tracheostomy was not clear because all such adverse events did not cause airway obstruction. Then, we have to clear the definite indication of planned tracheostomy to introduce the TOS as a minimally invasive treatment.

The rate of postoperative stenosis in the present study was only 0.5% (3/575). In the three cases who developed stenosis, the pathological tumor diameters were 15, 16, and 45 mm, respectively. And, all lesion located in the pyriform sinus. The possible reason developed stenosis might be associated with the tumor lesion regardless of the tumor size because the pyriform sinus is directly connected to the cervical esophagus which is physiological stenotic part.

Indication for TOS has not been clearly determined. In this study, pathological criteria for intraepithelial SCC and

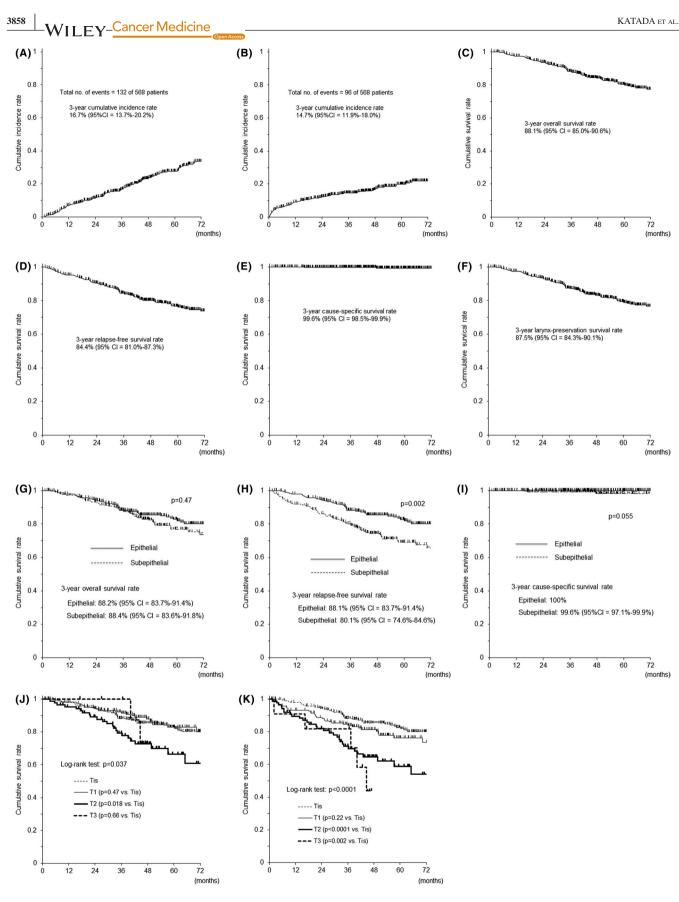


FIGURE 2 Cumulative incidence of metachronous cancers and survival rate. A, Cumulative incidence rate of metachronous head and neck cancers. B, Cumulative incidence rate of metachronous cancers arising in other organs. C, Overall survival rate. D, Relapse-free survival rate. E, Cause-specific survival rate. F, Larynx-preservation survival rate. G, Overall survival rates according to the histopathological depth of invasion. I, Cause-specific survival rates according to the histopathological depth of invasion. I, Cause-specific survival rates according to the histopathological depth of invasion. J, Overall survival rates according to the T category. K, Relapse-free survival rates according to the T category

	Total number of metachronous cancers ^a	Endoscopic resection	Surgery-based therapy	Chemoradiation-based therapy	Chemotherapy or hormonal therapy	Observation	TACE/ RFA ^b
Esophageal cancer	06	74	11	5	0	0	0
Gastric cancer	16	12	4	0	0	0	0
Lung cancer	10	1	6	2	1	0	0
Colorectal cancer	5	1	3	0	0	1	0
Bile-duct cancer	4	0	3	0	1	0	0
Liver cancer	2	0	1	0	0	0	1
Duodenal cancer	1	0	0	0	0	1	0
Prostate cancer	1	0	0	0	1	0	0
Urinary tract cancer	1	0	1	0	0	0	0
Thyroid cancer	1	0	1	0	0	0	0
Total	131	88	30	7	3	2	1

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subepithelial SCC have been clearly defined. Using this criteria, relapse-free survival rates were significantly different between two groups, while overall survival was similar. Cause-specific survival rate was not statistically different between the two groups because both groups had nearly 100% cause-specific survival. These results indicated that our pathological criteria for subepithelial invasion is clinically useful to stratify the risk for recurrence but not survival after TOS.

Early detection of head and neck cancer continues to be difficult worldwide. Screening of cancer in the head and neck is not a common practice. However, early detection is important because advanced head and neck cancer has poor prognosis and conventional treatments adversely affect the patients' quality of life. Image enhanced endoscopy such as NBI is revealed to be useful for early detection of head and neck cancer.¹² However, it is not routinely used in Western countries, while they were high incidence area for head and neck cancer. We would like to emphasize the benefit of image-enhanced endoscopy and hope it will be used in routine clinical practice especially in countries with known high incidence of head and neck cancer.

Our study has several limitations. This is a retrospective study and the duration of follow-up was relatively short. Although this national multi-center survey showed realworld outcomes and benefit of TOS for superficial head and neck cancer and we have shown clinically meaningful pathological criteria of subepithelial invasion, a prospective study would provide a better assessment of individual management of TOS.

In conclusion, TOS for superficial head and neck cancer appears to be an excellent organ preserving minimally invasive treatment that results in excellent cause-specific survival.

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CONFLICT OF INTEREST

^bTACE: transcatheter arterial chemoembolization, RFA: radiofrequency ablation.

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AUTHOR CONTRIBUTIONS

CK, MM, SF, TEY, YAS, HW, TS, AO, TAK, and RH contributed to conception and design. CK, KOK, YAS, and MI

Treatment of metachronous cancer arising in other organ

TABLE 6

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contributed to collection and assembly of data. CK, MM, SF, TEY, YAS, and RH contributed to data analysis and interpretation. CK, MM, SF, and TEY wrote the manuscript. RH provided financial support. CK, KOK, YAS, MI, MM, and RH contributed to administrative support. TOY, AW, TI, SY, IT, HM, YUS, AT, TOK, NH, MA, AS, KOK, YA, TM, HU, KO, KG, SH, YO, ST, YN, KT, KEK, MA, NS, and AA provided study materials or patients. All authors provided final approval of manuscript.

ETHICAL CONSIDERATION

This study was approved by the institutional review board at Kitasato University School of Medicine (Approval ID; B10-134).

DATA AVAILABILITY STATEMENT

The data and other items supporting the results of the study will be made available upon reasonable request.

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REFERENCES

- Bray F, Ferlay J, Soerjomataram I, Siegel RL, Torre LA, Jemal A. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin.* 2018;68:394-424.
- 2. Hoffman HT, Karnell LH, Shah JP, et al. Hypopharyngeal cancer patient care evaluation. *Laryngoscope*. 1997;107:1005-1017.
- Seiwert TY, Cohen EE. State-of-the-art management of locally advanced head and neck cancer. Br J Cancer. 2005;92: 1341-1348.
- Bova R, Goh R, Poulson M, Coman WB. Total pharyngolaryngectomy for squamous cell carcinoma of the hypopharynx: a review. *Laryngoscope*. 2005;115:864-869.
- Chao KS, Deasy JO, Markman J, et al. A prospective study of salivary function sparing in patients with head-and-neck cancers receiving intensity-modulated or three-dimensional radiation therapy: initial results. *Int J Radiat Oncol Biol Phys.* 2001;49:907-916.
- Gogarty DS, Shuman A, O'Sullivan EM, et al. Conceiving a national head and neck cancer screening programme. *J Laryngol Otol.* 2016;130:8-14.
- Japan Society for Head and Neck Cancer. *General rules for clinical studies on head and neck cancer*. Tokyo: Kanehara Co., Ltd.; 2018. (in Japanese).
- Muto M, Nakane M, Katada C, et al. Squamous cell carcinoma in situ at oropharyngeal and hypopharyngeal mucosal sites. *Cancer*. 2004;101:1375-1381.
- Katada C, Tanabe S, Koizumi W, et al. Narrow band imaging for detecting superficial squamous cell carcinoma of the head and neck in patients with esophageal squamous cell carcinoma. *Endoscopy*. 2010;42:185-190.

- Katada C, Muto M, Nakayama M, et al. Risk of superficial squamous cell carcinoma developing in the head and neck region in patients with esophageal squamous cell carcinoma. *Laryngoscope*. 2012;122:1291-1296.
- Katada C, Yokoyama T, Yano T, et al. Drinking alcohol, multiple dysplastic lesions and the risk of field cancerization of squamous cell carcinoma in the esophagus and head and neck region. *Gastroenterology*. 2016;151:860-869.
- Muto M, Minashi K, Yano T, et al. Early detection of superficial squamous cell carcinoma in the head and neck region and esophagus by narrow band imaging: a multicenter randomized controlled trial. *J Clin Oncol.* 2010;28:1566-1572.
- Goda K, Dobashi A, Yoshimura N, et al. Dual-focus versus conventional magnification endoscopy for the diagnosis of superficial squamous neoplasms in the pharynx and esophagus: a randomized trial. *Endoscopy*. 2016;48:321-329.
- Tateya I, Morita S, Muto M, et al. Magnifying endoscope with NBI to predict the depth of invasion in laryngo-pharyngeal cancer. *Laryngoscope*. 2015;125:1124-1129.
- Kikuchi D, Iizuka T, Yamada A, et al. Utility of magnifying endoscopy with narrow band imaging in determining the invasion depth of superficial pharyngeal cancer. *Head Neck*. 2015;37:846-850.
- Shimizu Y, Yamamoto J, Kato M, et al. Endoscopic submucosal dissection for treatment of early stage hypopharyngeal carcinoma. *Gastrointest Endosc.* 2006;64:255-259.
- Iizuka T, Kikuchi D, Hoteya S, Yahagi N, Takeda H. Endoscopic submucosal dissection for treatment of mesopharyngeal and hypopharyngeal carcinomas. *Endoscopy*. 2009;41:113-117.
- Muto M, Satake H, Yano T, et al. Long-term outcome of transoral organ-preserving pharyngeal endoscopic resection for superficial pharyngeal cancer. *Gastrointest Endosc.* 2011;74:477-484.
- Okada K, Tsuchida T, Ishiyama A, et al. Endoscopic mucosal resection and endoscopic submucosal dissection for en bloc resection of superficial pharyngeal carcinomas. *Endoscopy*. 2012;44:556-564.
- Hanaoka N, Ishihara R, Takeuchi Y, et al. Clinical outcomes of endoscopic mucosal resection and endoscopic submucosal dissection as a transoral treatment for superficial pharyngeal cancer. *Head Neck.* 2013;35:1248-1254.
- Imai K, Tanaka M, Hasuike N, et al. Feasibility of a "resect and watch" strategy with endoscopic resection for superficial pharyngeal cancer. *Gastrointest Endosc.* 2013;78:22-29.
- Satake H, Yano T, Muto M, et al. Clinical outcome after endoscopic resection for superficial pharyngeal squamous cell carcinoma invading the subepithelial layer. *Endoscopy*. 2015;47:11-18.
- Hanaoka N, Ishihara R, Takeuchi Y, et al. Endoscopic submucosal dissection as minimally invasive treatment for superficial pharyngeal cancer: a phase II study. *Gastrointest Endosc*. 2015;82:1002-1008.
- Yoshio T, Tsuchida T, Ishiyama A, et al. Efficacy of double-scope endoscopic submucosal dissection and long-term outcomes of endoscopic resection for superficial pharyngeal cancer. *Dig Endosc*. 2017;29:152-159.
- Shiotani A, Tomifuji M, Araki K, Yamashita T, Saito K. Videolaryngoscopic transoral en bloc resection of supraglottic and hypopharyngeal cancers using laparoscopic surgical instruments. *Ann Otol Rhinol Laryngol.* 2010;119:225-232.

30.

2013;35:1162-1167.

2016;30:323-329.

cers. Jpn J Clin Oncol. 2013;43:782-787.

cancers. Head Neck. 2017;39:1779-1787.

tion. Gastrointest Endosc. 2008;67:799-804.

-WILE

- 26. Okami K, Ebisumoto K, Sakai A, et al. Transoral en bloc resection of superficial laryngeal and pharyngeal cancers. Head Neck. 27. Nakayama M, Katada C, Mikami T, et al. A clinical study of transoral pharyngectomies to treat superficial hypopharyngeal can-28. Tateya I, Muto M, Morita S, et al. Endoscopic laryngo-pharyngeal surgery for superficial laryngo-pharyngeal cancer. Surg Endosc. 29. Watanabe A, Taniguchi M, Kimura Y, et al. Synopsis of transoral endoscopic laryngopharyngeal surgery for superficial pharyngeal cam4.3927 Ishihara R, Iishi H, Takeuchi Y, et al. Local recurrence of large squamous-cell carcinoma of the esophagus after endoscopic resec-
- 31. Furue Y, Katada C, Tanabe S, et al. Effectiveness and safety of endoscopic aspiration mucosectomy and endoscopic submucosal dissection in patients with superficial esophageal squamous-cell carcinoma. Surg Endosc. 2019;33:1433-1440.

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