

Original Article

Survival Benefits From Surgery for Stage IVa Head and Neck Squamous Cell Carcinoma: A Multi-institutional Analysis of 1,033 Cases

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Objectives. Head and neck squamous cell carcinomas (HNSCs) are frequently diagnosed at the locoregional advanced stage (stage IVa), but controversy remains regarding whether stage IVa HSNCs should be treated with upfront surgery or definitive chemoradiation therapy (CRT). The purpose of this study was to compare overall survival (OS) and disease-free survival (DFS) in patients with stage IVa HNSC treated primarily by surgery with curative intent with/without (neo)adjuvant treatment (surgery group) versus those treated primarily with CRT (CRT group).

Methods. We reviewed data of 1,033 patients with stage IVa HNSC treated with curative intent at 17 cancer centers between 2010 and 2016.

Results. Among 1,033 patients, 765 (74.1%) received upfront surgery and 268 (25.9%) received CRT. The 5-year OS and DFS rates were 64.4% and 62.0% in the surgery group and 49.5% and 45.4% in the CRT group, respectively. In multivariate analyses, OS and DFS were better in the surgery group than in the CRT group (odds ratio [OR] for death, 0.762; 95% confidence interval [CI], 0.592–0.981; OR for recurrence, 0.628; 95% CI, 0.492–0.802). In subgroup analyses, the OS and DFS of patients with oropharyngeal cancer were better in the surgery group (OR for death, 0.548; 95% CI, 0.341–0.879; OR for recurrence, 0.598; 95% CI, 0.377–0.948). In the surgery group, patients with laryngeal cancer showed better OS (OR for death, 0.432; 95% CI, 0.211-0.882), while those with hypopharyngeal cancer DFS was improved (OR for recurrence, 0.506; 95% CI, 0.328–0.780).

Conclusion. A survival benefit from surgery may be achieved even in patients with stage IVa HNSC, particularly those with oropharyngeal and laryngeal cancer. Surgery led to a reduction in the recurrence rate in patients with hypopharyngeal cancer.

Keywords. Squamous Cell Carcinoma of Head And Neck; Oral Neoplasms; Oropharyngeal Neoplasm; Laryngeal Neoplasms; Hypopharyngeal Neoplasms; Survival, Disease-Free Survival; Survival Rate; Prognosis; Surgery

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INTRODUCTION

Head and neck squamous cell carcinoma (HNSC) is the sixth most common type of cancer [1], accounting for more than 700,000 new cases and 350,000 cancer deaths worldwide [2]. HNSCs may arise from the oral cavity (354,864 cases/yr [2.1% of all cancers]), larynx (177,422 [1.0%]), oropharynx (92,887 [0.5%]), or hypopharynx (80,608 [0.4%]) [2] and are frequently diagnosed at an advanced stage (stage IV). The prognosis of these patients is poor. Furthermore, between 2004 and 2015, the incidence rate for stage IV HNSC increased significantly, by 26.1% [3,4].

Stage IV disease is subdivided into stages IVa and IVb. In the seventh edition of the American Joint Committee on Cancer (AJCC) staging system, the term "advanced resectable" defining stage IVa was replaced with the term "moderately advanced," while the previous definition of stage IVb as "advanced unresectable" was replaced with "very advanced." A resectable tumor is one in which the gross tumor can be removed without any residual tumor, such that local control can be achieved. However, in locally advanced HNSC, identifying resectable tumors is often difficult. In addition, a substantial proportion of stage IVa disease, although resectable, is currently treated nonsurgically, either due to institutional preference or to the patient's refusal of surgery. Clear guidelines for the treatment of stage IVa disease are thus far lacking, due to the ethical concerns associated with conducting a randomized controlled trial (RCT) comparing surgery versus nonsurgical treatment in patients with advanced HNSC. Recently, several clinical trials have shown that nonsurgical treatment in some patients with advanced HNSC may lead to satisfactory oncologic outcomes [5-8]. Nonetheless, for patients with stage IVa HNSC, surgical therapy remains the preferred approach as better oncologic outcomes are expected.

To address this issue, we conducted a large-scale, multi-institutional clinical study of patients with stage IVa HNSC. Specifically, overall survival (OS) and disease-free survival (DFS) were compared in patients with stage IVa HNSC treated primarily

HIGHLIGHTS

- It remains unclear whether upfront surgery improves overall survival (OS) and disease-free survival (DFS) in patients with stage IVa head and neck squamous cell carcinoma (HNSC) compared to definitive chemoradiation therapy.
- A survival benefit from the upfront surgical treatment may be achieved even in patients with stage IVa HNSC.
- Upfront surgical treatment improved OS and DFS in patients with stage IVa HNSC, particularly in those with oropharyngeal and laryngeal cancer.
- In patients with hypopharyngeal cancer, surgery improved DFS, but not OS.

with surgery or nonsurgical modalities.

MATERIALS AND METHODS

Study design

This study was designed by the members of the Research Committee of the Korean Society of Head and Neck Surgery. The study protocol was approved by the Institutional Review Board of the Catholic University of Korea (IRB No. XC 19RCD10096) and the IRBs of all participating institutions. Data were collected retrospectively from medical reports of patients with stage IVa HNSC initially treated in 17 hospitals between January 2010 and December 2016. The hospitals belonged to the following academic medical institutions: Ajou University Hospital, Catholic Medical Center (six St. Mary's hospitals), Dankook University Hospital, Inha University Hospital, Jeonbuk National University Hospital, Korea Institute of Radiological and Medical Sciences (Korea Cancer Center Hospital), Korea University Medicine (three Korea University Hospitals), Seoul National University Bundang Hospital, and Yonsei University Health System (two Severance Hospitals). All of these institutions are high-volume centers with extensive experience in the treatment of HN-SCs by well-trained multidisciplinary teams.

Patients who met the following criteria were included in the study: diagnosed with HNSC involving the oral cavity, oropharynx, larynx, or hypopharynx; with clinically staged Iva disease according to the 8th edition of the AJCC staging system (for human papilloma virus [HPV]-positive oropharynx cancer, stage IV disease is reserved only for distant metastases [M1 disease]. Therefore, among oropharyngeal cancer patients, only HPVnegative or not-tested patients were included in this study); no previous treatment for the same diagnosis (cancer of the head and neck) before hospitalization; no other malignant disease within 5 years at the time of diagnosis and follow up of >12 months. The patients were divided into two groups based on the treatment modality. Patients in the surgery group were those treated primarily by surgery with curative intent with/without (neo)adjuvant treatment, and patients in the CRT group those treated primarily by chemotherapy and radiotherapy. Clinical staging was conducted in the outpatient clinic and/or the operating room and consisted of a physical examination using a flexible or rigid endoscope and radiological evaluation such as computed tomography, magnetic resonance imaging, ultrasonography, and positron emission tomography. Therapeutic strategies differed slightly between institutions as they were decided by institutional policy, although they largely met the guideline of the National Comprehensive Cancer Network.

Statistical analysis

All analyses were performed using SPSS software ver. 16.0 (SPSS Inc., Chicago, IL, USA). A *P*-value < 0.05 was considered to in-

dicate statistical significance. The significance of the relationships between the two groups and clinical factors were analyzed using a chi-square test, Fisher's exact test, or Student *t*-test, as appropriate. The Kaplan Meier method was used to calculate 5-year OS and DFS rates; differences in survival rates between the two groups were assessed using the log-rank test. Multivariate analyses using a Cox proportional-hazards model were performed to compare the factors with prognostic potential indicated by univariate analyses. The primary endpoint was OS, calculated as the length of time from diagnosis until death or last follow-up. The secondary endpoint was DFS, calculated as the length of time from diagnosis until first documented recurrence or death.

RESULTS

Patients

Data were collected from 1,405 patients. After the exclusion of 372 patients who did not meet the inclusion criteria (p16(+) oropharyngeal cancer [n=177], under- or over-staging [n=175], and other reasons [n=20]), the final study population consisted of 1,033 patients divided among the participating institutions as follows: Ajou University Hospital (n=67), Catholic Medical Center (n=307), Dankook University Hospital (n=29), Inha University Hospital (n=51), Jeonbuk National University Hospital (n=89), Korea Cancer Center Hospital (n=68), Korea University Medicine

(n=111), Seoul National University Bundang Hospital (n=79) and Yonsei University Health System (n=232). The schema of the cohort is described in Fig. 1. The median follow-up among surviving patients was 31 months (range, 12–132 months). The median 859 male and 174 female patients had a median age of 61 years (range, 14–101 years). Regarding clinical T stage, 150 (14.5%), 280 (27.1%), 152 (14.7%), and 451 (43.7%) patients had stage T1 to T4a cancers, respectively. The staging of the cervical lymph nodes was N0 in 96 patients (9.3%), N1 in 105 patients (10.2%), and N2 in 832 (80.5%) patients. The surgery group consisted of 765 patients (74.1%) and the CRT group contained 268 patients (25.9%). The anatomical subsites of the HNSCs in patients of both groups are shown in Fig. 2. The baseline characteristics of the patients in the two groups are summarized in Table 1.

OS of patients with HNSC

There were 232 deaths in the surgery group and 98 in the CRT group. The 5-year OS rates were 64.4% (95% confidence interval [CI], 62.4%–66.4%) and 49.5% (95% CI, 46.5%–52.5%), respectively. There were 249 and 107 overall recurrences, with 5-year DFS rates of 62.0% (95% CI, 59.9%–64.1%) and 45.4% (95% CI, 41.5%–49.3%), respectively. The Kaplan-Meier survival curves are shown in Fig. 3. The results of the univariate analyses are presented in Supplementary Table 1. In multivariate analyses, OS and DFS were significantly higher in the surgery group (Table 2). The recurrence patterns in both groups are summarized

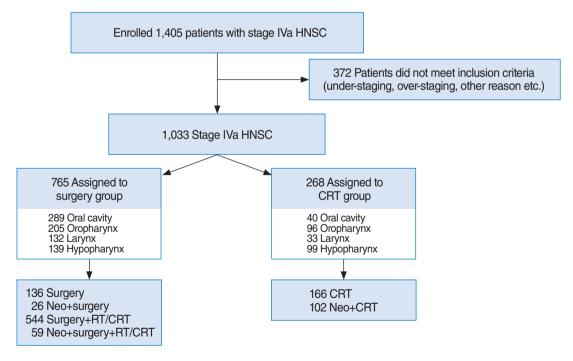


Fig. 1. Schema of the cohort. The study design, including the allocation of patients to the surgery group and CRT group according to the primary treatment modality, is shown. HNSC, head and neck squamous cell carcinoma; Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment; CRT group, patients treated primarily with chemotherapy and radiotherapy; Neo, neoadjuvant chemotherapy; RT, radiotherapy; CRT, chemoradiation therapy.

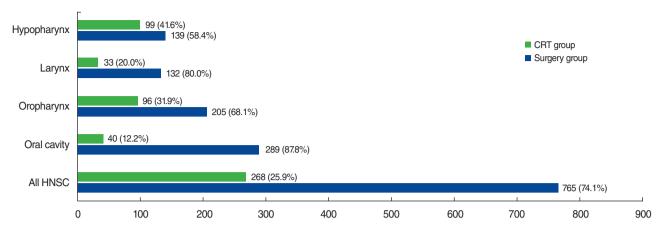


Fig. 2. The proportion of surgery-based treatments in patients (n=1,033) with stage IVa head neck cancer, shown by anatomical location of the primary tumor. CRT group, patients treated primarily with chemotherapy and radiotherapy; Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment; HNSC, head and neck squamous cell carcinoma.

Table 1. Baseline characteristics of all patients

Variable	Total (n=1,033)	Surgery group (n=765)	CRT group (n=268)	P-value
Age (yr)				0.979
≥60	431 (41.7/100)	319 (41.7/74.0)	112 (41.8/26.0)	
<60	602 (58.3/100)	446 (58.3/74.1)	156 (58.2/25.9)	
Sex				0.007*
Male	859 (83.2/100)	622 (81.3/72.4)	237 (88.4/27.6)	
Female	174 (16.8/100)	143 (18.7/82.2)	31 (11.6/17.8)	
T stage				0.116
Non T4a	582 (56.3/100)	442 (57.8/75.9)	140 (52.2/24.1)	
T4a	451 (43.7/100)	323 (42.2/71.6)	128 (47.8/28.4)	
N stage				0.056
N0-1	201 (19.5/100)	160 (20.9/79.6)	41 (15.3/20.4)	
N2	832 (80.5/100)	605 (79.1/72.7)	227 (84.7/27.3)	
Follow-up period (mo)	31 (12–132)	33 (12–122)	23 (12–132)	0.004*

Values are presented as number of patients (%within group/%within variable) or median (range).

Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment; CRT group, patients treated primarily with chemotherapy and radiotherapy.

^{*}Statistically significant (P<0.05).

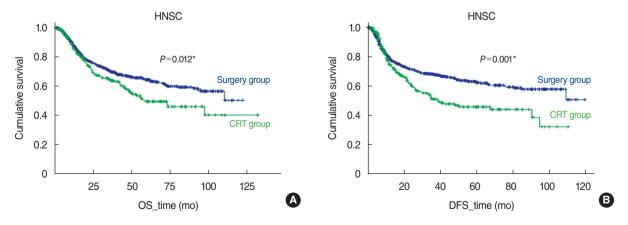


Fig. 3. Kaplan-Meier survival curves for patients with stage IVa HNSC (n=1,033). (A) Overall survival (OS), (B) disease-free survival (DFS) treated surgically (surgery group) or nonsurgically (CRT group). HNSC, head and neck squamous cell carcinoma; Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment; CRT group, patients treated primarily with chemotherapy and radiotherapy. *Statistically significant (*P*<0.05).

in Supplementary Table 2. The majority of first recurrences were locoregional in both groups. There were 163 recurrences in 765 patients (21.2%) in the surgery group, and 94 recurrences in 268 patients (34.9%) in the CRT group.

Survival according to HNSC subgroups

The benefits of surgery as determined by OS and DFS were evaluated according to the prognostic indicators identified in the univariate analyses.

Primary tumor site

The Kaplan-Meier survival curves based on the primary tumor site are shown in Supplementary Fig. 1 and Fig. 4. In the multivariate analyses, OS and DFS rates were significantly higher in patients with oropharyngeal and laryngeal cancer than in those with oral cancer. The results are summarized in Table 2.

1) Oral cavity

In the surgery group, the 5-year OS and DFS rates were 59.8% (95% CI, 56.4%–63.2%) and 52.5% (95% CI, 49.0%–56.0%), respectively. In the CRT group the corresponding rates were

53.1% (95% CI, 42.4%–63.8%) and 39.3% (95% CI, 26.9%–51.7%), respectively. The differences between the two groups were not significant.

2) Oropharynx

The 5-year OS and DFS rates were 76.9% (95% CI, 73.5%–80.3%) and 71.8% (95% CI, 68.2%–75.4%), respectively, in the surgery group and 53.1% (95% CI, 46.4%–59.8%) and 54.5% (95% CI, 48.2%–60.8%), respectively, in the CRT group. In the multivariate analyses, after adjustment for covariates, OS and DFS were significantly higher in the surgery group (Supplementary Table 3, Fig. 5).

3) Larynx

The OS and DFS rates were 69.0% (95% CI, 64.0%–74.0%) and 63.9% (95% CI, 58.1%–69.7%), and 42.9% (95% CI, 29.6%–56.2%) and 53.4% (95% CI, 41.8%–65.0%), respectively, in the two groups. In multivariate analyses, after adjustment for covariates, OS was significantly higher in the surgery group (Supplementary Table 4, Fig. 5). Patients in the surgery group included those who underwent total larvngectomy (n=84,

Table 2. Multivariate analyses (Cox proportional hazard model) of the clinical parameters predicting the prognosis of patients with stage IVa HNSC (n=1,033)

Variable	Overall survival		Disease-free survival	
	OR (95% CI)	P-value	OR (95% CI)	P-value
Age ≥60 yr	1.550 (1.231–1.952)	0.001*	1.386 (1.112–1.728)	0.004*
Primary site		<0.001*		<0.001*
Oral cavity	1.000	-	1.000	-
Oropharynx	0.558 (0.410-0.757)	<0.001*	0.522 (0.390-0.698)	<0.001*
Larynx	0.615 (0.429-0.881)	0.008*	0.631 (0.454-0.878)	0.006*
Hypopharynx	1.031 (0.771–1.379)	0.835	0.747 (0.558–1.000)	0.050
T4a stage	1.157 (0.924–1.450)	0.204	1.362 (1.094–1.697)	0.006*
Surgery group	0.762 (0.592-0.981)	0.035*	0.628 (0.492-0.802)	<0.001*

HNSC, head and neck squamous cell carcinoma; OR, odds ratio; CI, confidence interval; Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment.

^{*}Statistically significant (P<0.05).

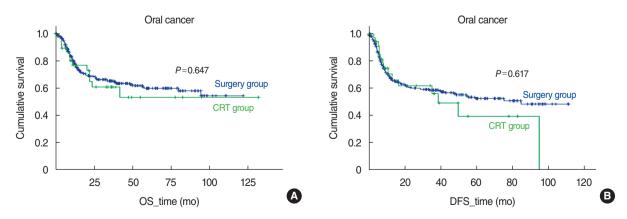


Fig. 4. Kaplan-Meier survival curves according to treatment modality and primary tumor site or T4a stage. Oral cancer (n=329): (A) overall survival (OS) and (B) disease-free survival (DFS).

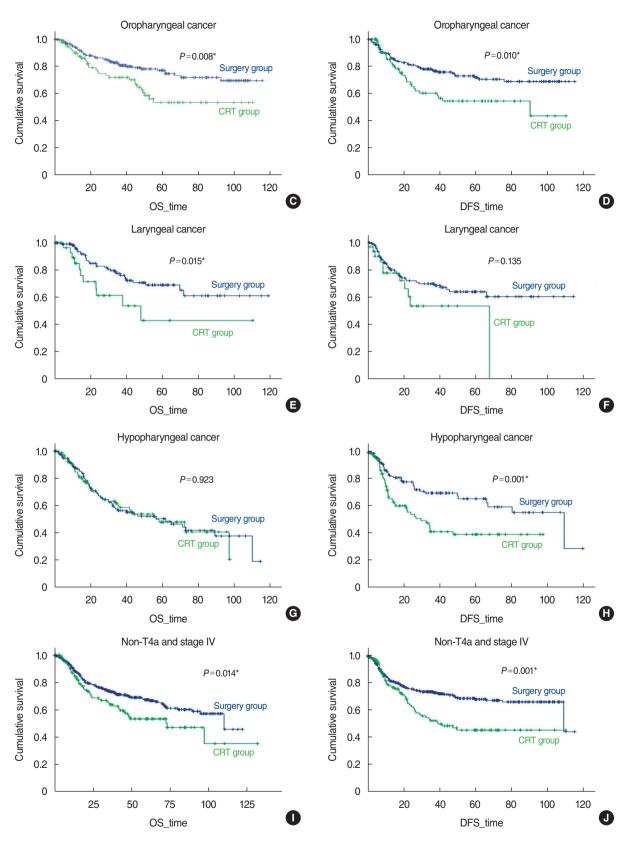


Fig. 4. (Continued) Oropharynx (n=301): (C) OS and (D) DFS. Larynx (n=165): (E) OS and (F) DFS. Hypopharynx (n=238): (G) OS and (H) DFS. Non-T4a (n=582): (I) OS and (J) DFS. Surgery group, patients treated primarily by surgery with curative intent with/without (neo)adjuvant treatment; CRT group, patients treated primarily with chemotherapy and radiotherapy. *Statistically significant (*P*<0.05).

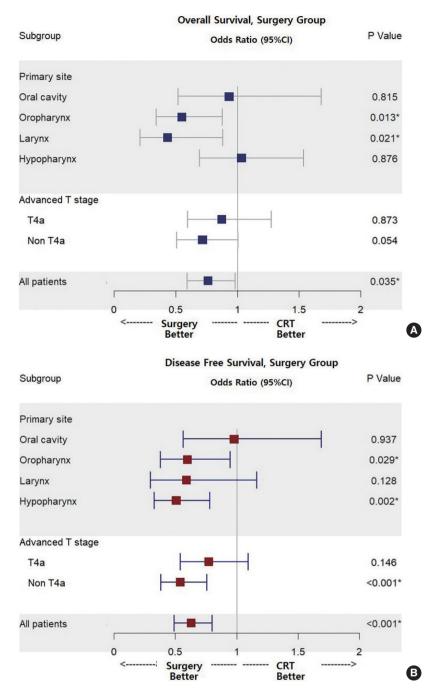


Fig. 5. Survival benefits from surgery by primary tumor site or T4a stage. (A) Overall survival, (B) disease-free survival. CI, confidence interval; CRT group, patients treated primarily with chemotherapy and radiotherapy. *Statistically significant (*P*<0.05).

63.6%) and those who underwent partial laryngectomy (n=48, 36.4%).

4) Hypopharynx

The OS and DFS rates were 50.2% (95% CI, 45.1%–52.3%) and 64.5% (95% CI, 59.2%–69.8%), and 47.7% (95% CI, 41.6%–53.8%) and 37.7% (95% CI, 31.8%–43.6%), respectively. In multivariate analyses, after adjustment for covariates,

DFS was significantly higher in the surgery group (Supplementary Table 5, Fig. 5).

T4a vs. non-T4a disease

1) T4a disease

For patients with stage T4a disease, the 5-year OS and DFS rates were 59.9% (95% CI, 56.6%–63.2%) and 53.5% (95% CI, 50.1%–56.9%), respectively, in the surgery group and 42.2%

(95% CI, 35.3%–49.1%) and 46.6% (95% CI, 40.8%–52.4%), respectively, in the CRT group. The differences between the two groups were not significant (P=0.340 and P=0.234).

2) Non-T4a disease (T1-3N2)

For patients with non-T4a stage disease, the OS and DFS rates were 67.4% (95% CI, 64.8%–70.0%) and 67.8% (95% CI, 65.2%–70.4%), and 53.4% (95% CI, 48.4%–58.4%) and 45.1% (95% CI, 39.8%–50.4%), respectively. The differences between the two groups were significant (P=0.014 and P=0.001). In multivariate analyses, OS was higher in the surgery group (OR for death, 0.716; 95% CI, 0.509–1.006) but did not significantly differ from that of the CRT group (P=0.054). By contrast, DFS was significantly higher in the surgery group (OR for recurrence, 0.538; 95% CI, 0.382–0.759; P<0.001) (Supplementary Table 6, Fig. 5).

DISCUSSION

To the best of our knowledge, this is the largest study of a stage IVa HNSC cohort reported in the English-language medical literature. Ethical issues have hindered RCTs comparing the oncologic outcomes of advanced HNSC patients treated surgically or nonsurgically. In major hospitals of South Korea, 74.1% of patients with advanced HNSC have been treated surgically during the past 6 years (2010–2016), a therapeutic strategy supported by our results showing improved survival in these patients. After adjustment for covariates expected to affect prognosis, surgery reduced the odds of death and disease recurrence in patients with stage IVa HNSC by 26% and 38%, respectively. In their RCT, Iver et al. [9] found either no differences in outcomes or a slight advantage favoring surgery-based treatment in stage III/IV HNSC. However, that study included a broad spectrum of cancer stages, from T1N2M0 to T4N2M0, and did not specifically address the prognosis of patients with stage IVa disease. An additional benefit of surgery is that it generates pathologic information, including surgical margins, perivascular invasion, perineural invasion, and pathologic stage, all of which are potent prognostic predictors and may lead to patient-tailored treatment, including the need for chemotherapy or radiation therapy. In our patient cohort, 80% of patients in the surgery group underwent adjuvant treatment due to positive surgical margins (30%) or other reasons (50%).

Surgery is the generally accepted treatment of choice for oral cancer. Thus, analyses of survival in patients treated surgically or nonsurgically were not possible, as the majority of patients were treated by surgery and a control group of chemoradiation therapy (CRT) patients was not available. Previous reports also support surgery as the preferred treatment in oral cancer [9-11]. In the RCT of Iyer et al. [9], among 32 patients with locally advanced oral cancer, those who underwent surgery followed by radiotherapy (SRT) had better survival than those treated with

CRT (68% vs. 12%). Spiotto et al. [10] used the National Cancer Database to compare the survival outcomes of 6,900 patients with stage III/IVa oral cancer. SRT was associated with better survival than CRT alone (3-year OS, 53.9% vs. 37.8%).

Among oropharyngeal cancers, the incidence of HPV-positive tumors is steadily increasing but their characteristics differ from those of HPV-negative tumors [12,13]. In our study, only HPVnegative or non-tested patients were included. In these patients, surgery reduced the odds of death by 46% and disease recurrence by 41%. Our findings suggest that surgery-based treatments can achieve better survival than CRT in patients with HPV-negative oropharyngeal cancer. However, the results reported in the literature vary. The multicenter retrospective study by Song et al. [14] compared the prognosis of 586 patients treated by SRT versus CRT, including 419 patients with stage IVa oropharyngeal cancer, and reported similar survival outcomes. Zenga et al. [15] reported that among patients with T4 oropharyngeal cancer, particularly in the era of HPV-positive disease, nonsurgical treatment had a significant negative association with OS (OR, 2.79), disease-specific survival (OR, 3.38), and DFS (OR, 2.59). Kelly et al. [16] compared survival outcomes among patients with cT1-2 N1-2b HPV-negative oropharyngeal cancer and found that upfront surgery was not associated with a better OS than achieved with CRT (OR, 1.01). Roden et al. [17] analyzed the National Cancer Database and found that although CRT was the most common treatment modality for stage III-IVb oropharyngeal cancer (48.1%) in the United States, patients who underwent surgery and CRT had a higher 3-year OS (88.5%) than did patients treated with CRT alone (74.2%).

Surgery reduced the odds of overall death by 57% in our patients with laryngeal cancer. However, two randomized trials supporting nonsurgical treatment have had a significant impact on actual practice [18,19]. Trends in the treatment of advanced laryngeal cancer in the United States [20] changed between 1985 and 2007, with CRT increasing from 7% to 45% and surgery decreasing from 42% to 32%. However, these two trials covered stage III/IV disease (possibly analyzing T1N2M0 and T4N2M0 together), which may have resulted in biased results. A study based on the data of 7,019 patients from a national hospital-based cancer registry [21] reported that treatment outcomes for patients with stage III and IV disease differed, thus complicating drawing a unique conclusion. A study by Patel et al. [22] using data from 8,703 patients entered in the National Cancer Database showed that for those with T4 disease, surgery improved OS (OR, 0.80; 95% CI, 0.62-0.92), but in patients with non-T4 disease and a high nodal burden (T2-T3N2-N3) it worsened survival (OR, 1.25; 95% CI, 1.04-1.51) compared to CRT. Other studies [23,24] have also reported better survival in patients with T4 disease treated surgically than nonsurgically and recommended total laryngectomy for those with T4 laryngeal cancer.

Whether patients with advanced hypopharyngeal cancer without distant metastasis should be treated by SRT or CRT remains

controversial. Retrospective studies have included patients with stage III and IV disease and the results have differed. To the best of our knowledge, ours is the first study limited to stage IVa hypopharyngeal cancer in a large patient population. Although surgery did not confer a survival benefit, it reduced the odds of recurrence by 49%. Chung et al. [25] reported very similar results for 266 patients with stage III/IV disease. They showed that survival following nonsurgical treatment (44.6% for induction chemotherapy followed by [chemo]radiotherapy [ICT], 39.6% for CRT) was comparable to that of SRT (45.3%) and that the salvage rate after nonsurgical treatment was higher (12.5% for ICT, 15.6% for CRT, and 3.8% for SRT). Some authors have reported that survival and recurrence rates are not significantly different between treatment modalities [26,27], whereas others have reported that surgery improves OS and DFS in patients with stage III/IV disease [28].

The main limitation of our study was its retrospective design. In addition, as this was a multi-center study, patients were inevitably treated using slightly different modalities. In addition, the number of patients in the CRT group was relatively small, particularly in the subgroup with oral cancer, which is almost always treated surgically. Despite these limitations, our study provides the basis for a much-needed guideline for the treatment of stage IVa HNSC.

In summary, surgery can be expected to confer survival benefits in patients with stage IVa HNSC and OS was improved in patients with stage IVa oropharyngeal and laryngeal cancer. Furthermore, surgery reduced the recurrence rate in patients with hypopharyngeal cancer, although it did not improve OS. However, overall, surgery does not confer a survival benefit in patients with advanced T stage (T4a) HNSCs.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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SUPPLEMENTARY MATERIALS

Supplementary materials can be found via https://doi.org/10. 21053/ceo.2020.01732.

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