Effects of Surgery Combined with Chemoradiotherapy on Short- and Long-Term Outcomes of Early-Stage Nasopharyngeal Carcinoma

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Objective: The efficacy of surgery as the primary treatment modality for nasopharyngeal carcinoma (NPC) is yet to be clarified. Therefore, we aimed to explore the short- and long-term efficacy of surgery for early-stage NPC.

Methods: We retrospectively evaluated 341 patients diagnosed with early-stage NPC between September 2010 and December 2015. Among them, 58 patients underwent endoscopic nasopharyngectomy combined with chemoradiotherapy, whereas 283 patients underwent conventional chemoradiotherapy. The patients who underwent concurrent chemoradiotherapy or radiotherapy alone were matched to patients who underwent surgery in a 1:2 ratio using propensity score matching to analyze the clinical efficacy of each therapeutic modality. The primary endpoint was survival, and the secondary endpoints were tumor regression rate and reduction in Epstein-Barr virus (EBV)-DNA levels.

Results: After matching, 156 patients were enrolled (58 patients in the surgery group; 98 patients in the non-surgery group). The baseline data of the matched patients had good inter-group comparability (All $P > 0.05$). The surgery group had significantly higher 5-year overall survival (98.30% vs. 91.70%), disease-free survival (98.30% vs. 81.40%), and recurrence-free survival (100.00% vs. 90.10%) rates than did the non-surgery group (All $P < 0.05$). In total, 0 and 14 patients in the surgery and non-surgery groups, respectively, had residual cancer at the end of treatment ($P = 0.001$). All patients in the surgery group tested negative for EBV-DNA, whereas two patients in the non-surgery group tested positive. The incidence of hematologic toxicity during treatment was similar between the two groups (All $P > 0.05$). Still, the incidence of severe oral mucositis was lower in the surgery group than in the non-surgery group (37.9% vs. 54.08%, $P = 0.051$).

Conclusion: Surgery can improve the clearance rate of EB virus and reduce tumor residue. Surgery may be a safe and effective treatment for early NPC.

Keywords: nasopharyngeal carcinoma, endoscopic surgery, radiotherapy, prognosis

Introduction

Radical endoscopic surgery is the primary treatment modality for early laryngeal cancer¹ and thyroid cancer.² Endoscopic surgery not only enables complete tumor resection but also preserves organ functions. Nasopharyngeal carcinoma (NPC) is the most prevalent type of head and neck cancer in Southern China,³ with the most common pathological type being WHO type III, which is sensitive to radiotherapy. Most patients with NPC are diagnosed at the middle to the advanced stage of the disease due to the inaccessibility of the nasopharynx and the nonpalpable nature of early symptoms. Middle- to late-stage NPC is generally responsive to

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chemoradiotherapy. However, some patients experience treatment failure primarily due to residual disease, recurrence, distant metastasis, and nasopharyngeal hemorrhage.^{4,5} To ensure better local control rates, clinicians tend to increase the local radiation dose, which results in more radiotherapy-related complications.

The development of NPC screening modalities and advancements in imaging examinations have, in turn, increased the early detection rate of early-stage NPC in recent years.⁶ We previously found that endoscopic nasopharyngectomy yields better overall survival (OS) and lower incidence of xerostomia than conventional chemoradiotherapy.⁷ One study also showed that endoscopic nasopharyngectomy is not inferior to chemoradiotherapy with respect to recurrence and metastasis.⁸ Liu et al even reported residual tumors in a proportion of NPC patients who underwent chemoradiotherapy.⁹ Although surgery has been established as an effective modality for post-treatment residual or recurrent tumors,¹⁰ its efficacy as the initial treatment modality for NPC remains unclear. Further, the usefulness of neck dissection in NPC has rarely been studied. Thus, this study aimed to investigate the short- and long-term efficacy of endoscopic nasopharyngectomy combined with chemoradiotherapy with respect to the tumor regression rate and prognosis in patients with early NPC.

Data and Methods

Study Design and Patients

Data for the patients with NPC were retrieved from the database of The People's Hospital of Guangxi Zhuang Autonomous Region (Nanning, China). This study was approved by our institutional ethical committee (NO.20080301). We evaluated 341 patients diagnosed with early-stage NPC between September 2010 and December 2015. The inclusion criteria were: (1) treatment-naïve pathologically diagnosed early NPC (stage I and II); (2) Karnofsky score >80 points; (3) no distant metastasis. Exclusion criteria: (1) T1 patients with oropharynx invasion; (2) T2 patients with internal carotid artery involvement or posterior parapharyngeal space invasion, or patients with parapharyngeal lymph node enlargement. In total, 341 NPC patients were included. Among them, 58 underwent endoscopic nasopharyngectomy with or without neck dissection, whereas 283 underwent chemoradiotherapy. All patients were staged according to the

Union for International Cancer Control NPC staging system (7th Edition, 2010).

Treatment

Surgery

The detailed procedures for administering endoscopic nasopharyngectomy were performed as previously reported.⁷ All patients were administered general anesthesia before the surgery. Nasopharyngeal tumors were resected using a nasal electromotor, a plasma scalpel guided by a nasal endoscope. The tumor margins after primary resection were evaluated using magnetic resonance imaging (MRI) of the nasopharynx and skull base as well as intraoperative observations. For N1 patients, functional neck dissection was performed concurrently.

Radiotherapy

All patients received radiotherapy. For conventional radiotherapy, total radiation dose was 70~76Gy for nasopharyngeal lesions and 60~70Gy for cervical lesions in the non-surgery group, while total radiation dose was 60~64 Gy for nasopharyngeal lesions and 60~64 Gy for cervical lesions in the surgery group. For intensity-modulated radiation therapy (IMRT), the prescribed dose was 69~72Gy for nasopharyngeal lesions and cervical lesions in the non-surgery group, while the prescribed dose was 59.4~63.8Gy for nasopharyngeal lesions and cervical lesions in the surgery group.

Chemotherapy

In general, the patients with stage II disease (T2N0, T1N1 and T2N1) were recommended to receive concurrent chemotherapy. The chemotherapy programs were: (1) Nedaplatin 80mg/m² was administered via intravenous drip infusion on day 1, followed by a 120-h continuous infusion of 2000mg/m² 5-FU every four weeks for 2~3 cycles; or (2) Cisplatin 30mg/m² was administered via intravenous drip infusion on day 1~3, followed by a 120-h continuous infusion of 2000 mg/m² 5-FU every four weeks for 2~3 cycles.

Detection of Blood Cells

The blood tests for hemoglobin level, neutrophil counts, platelet counts, and white blood cell (WBC) counts were obtained before, during, and after treatment was carried out. Myelosuppression was assessed per the criteria for evaluating toxicity and side effects of drugs in the

Common Terminology Criteria for Adverse Events (Version 4.03) issued by the US National Cancer Institute.

Evaluation of Tumor Regression Rate

All patients underwent contrast-enhanced MRI of the nasopharynx and skull base before and at the end of treatment. The tumor regression rate was evaluated based on the Response Evaluation Criteria in Solid Tumors (version 1.1).¹¹

Evaluation of Oral Mucositis

Acute reactions in the oral and pharyngeal mucosa were evaluated according to the Common Terminology Criteria for Adverse Events (Version 4.03). Grade ≥ 3 mucosal reactions were classified as severe mucositis, whereas grade 1 or 2 reactions were mild mucositis.

Follow-Up

After completing treatment, patients were subsequently followed up every 3 months during the first years, every

6 months during the second year, and annually from the third year onwards.

The OS was defined as the period from the diagnosis of NPC to the date of death from any cause or the censoring of patients at the last follow-up. Disease-free survival (DFS) was defined as the period from the diagnosis to the date of disease recurrence, metastasis, or death. Relapse-free survival (RFS) was defined as the interval from the diagnosis to first recurrence or final follow-up. Distant metastasis-free survival (DMFS) was defined as the time from the diagnosis to first distant metastasis or final follow-up.

Statistical Analysis

The primary endpoint was survival, and the secondary endpoints were tumor regression rate and the reduction in Epstein-Barr virus (EBV-DNA) levels. All statistical analyses were performed using the R statistical software (ver.3.6.1). Propensity score matching of 1:2

Table 1 Baseline Characteristics of 341 Patients

Characteristics	All Patients (n=341)	Non-Surgery (n=283)	Surgery (n=58)	χ^2/t	P value
Age	47.26 (10.68)	47.63 (10.77)	45.45 (10.13)	1.42	0.158
Sex					
Male	230 (67.45%)	183 (64.66%)	47 (81.03%)	5.88	0.025
Female	111 (32.55%)	100 (35.34%)	11 (18.97%)		
T stage					
I	90 (26.39%)	56 (19.79%)	34 (58.62%)	37.36	<0.001
2	251 (73.61%)	227 (80.21%)	24 (41.38%)		
N stage					
0	174 (51.03%)	131 (46.29%)	43 (74.14%)	14.94	<0.001
I	167 (48.97%)	152 (53.71%)	15 (25.86%)		
Clinical stage					
I	58 (17.01%)	33 (11.66%)	25 (43.10%)	33.71	<0.001
II	283 (82.99%)	250 (88.34%)	33 (56.90%)		
Radiotherapy					
IMRT ^a	232 (68.04%)	194 (68.55%)	38 (65.52%)	0.20	0.774
2DRT ^b	109 (31.96%)	89 (31.45%)	20 (34.48%)		
Concurrent Chemotherapy					
Without	45 (13.20%)	39 (13.78%)	6 (10.34%)	0.50	0.546
With	296 (86.80%)	244 (86.22%)	52 (89.66%)		
Residual_position					
No	289 (84.75%)	231 (81.63%)	58 (100.00%)	12.57	0.007
NP ^c	29 (8.50%)	29 (10.25%)	0 (0.00%)		
LN ^d	22 (6.45%)	22 (7.77%)	0 (0.00%)		
NP&LN	1 (0.29%)	1 (0.35%)	0 (0.00%)		

Notes: ^aIntensity modulated radiotherapy; ^b2-dimensional radiotherapy; ^cNasopharynx; ^dCervical lymph nodes. Bold formatting indicates statistically significant.

Figure 1 Comparison of survival curves between residual and non-residual group in patients with early-stage nasopharyngeal carcinoma (n=341) ((A) Overall survival; (B) Disease-free survival; (C) Relapse-free survival; (D) Distant metastasis-free survival).

scheme was applied to develop comparable cohorts of patients with surgery and without surgery. Count data were compared using the χ^2 test, whereas measurement data were analyzed using the *t*-test or analysis of variance. Non-normally distributed measurement data were compared using the rank-sum test. Survival curves were plotted using the Kaplan-Meier method for univariate survival analysis and then analyzed using the Log rank test. Multivariate analysis was performed using the Cox proportional-hazards model. $P < 0.05$ was considered statistically significant.

Results

Patient Characteristics

Sex, T stage, N stage, clinical stage, and position of residual tumors were significantly different between the surgery and the non-surgery groups (All $P < 0.05$), as shown in Table 1. In total, 51 of the 341 patients had

residual tumors at the end of treatment (28 patients with nasopharyngeal residue; 22 patients, cervical residue; and one patient, nasopharyngeal plus cervical residues). There were 7 and 44 patients with T1 and T2 disease, respectively, who had residual tumors. At the end of treatment, the residual group had lower disease-free survival (DFS), recurrence-free survival (RFS), and distant metastasis-free survival (DMFS) rates than did the non-residual group (All $P < 0.05$), but there was no significant difference in their OS rates ($P > 0.05$) (Figure 1).

Matching results

After matching, 156 patients were included in the analysis (58 patients in the surgery group; 98 patients in the non-surgery group). There were no significant differences in age, sex, T stage, N stage, clinical stage, and radiotherapy modality (All $P > 0.05$) between the two groups (Table 2).

Table 2 Baseline Characteristics of Patients After Propensity Score Matching

Characteristics	All Patients (n=156)	Non-Surgery (n=98)	Surgery (n=58)	χ^2/t	P value
Age	47.06 (10.69)	48.01 (10.95)	45.45 (10.13)	1.45	0.149
Sex					
Male	123 (78.85%)	76 (77.55%)	47 (81.03%)	0.27	0.709
Female	33 (21.15%)	22 (22.45%)	11 (18.97%)		
T stage					
I	84 (53.85%)	50 (51.02%)	34 (58.62%)	0.85	0.426
2	72 (46.15%)	48 (48.98%)	24 (41.38%)		
N stage					
0	111 (71.15%)	68 (69.39%)	43 (74.14%)	0.40	0.601
I	45 (28.85%)	30 (30.61%)	15 (25.86%)		
Clinical stage					
I	57 (36.54%)	32 (32.65%)	25 (43.10%)	1.72	0.253
II	99 (63.46%)	66 (67.35%)	33 (56.90%)		
Radiotherapy					
IMRT ^a	113 (72.44%)	75 (76.53%)	38 (65.52%)	2.21	0.163
2DRT ^b	43 (27.56%)	23 (23.47%)	20 (34.48%)		
Concurrent Chemotherapy					
No	27 (17.31)	21 (21.43)	6 (10.34)	3.13	0.093
Yes	129 (82.69)	77 (78.57)	52 (89.66)		
Residual position					
No	141 (90.38%)	84 (85.71%)	58 (100.00%)	9.10	0.014
NP ^c	11 (7.05%)	10 (10.20%)	0 (0.00%)		
LN ^d	3 (1.92%)	3 (3.06%)	0 (0.00%)		
NP&LN	1 (0.64%)	1 (1.02%)	0 (0.00%)		

Note: Bold formatting indicates statistically significant.

Abbreviations: ^aIntensity modulated radiotherapy; ^b2-dimensional radiotherapy; ^cNasopharynx; ^dCervical lymph nodes.

Table 3 Univariate Survival Analysis of 156 Patients with Nasopharyngeal Carcinoma

Variable	OS		DFS		RFS		DMFS	
	χ^2	P value	χ^2	P value	χ^2	P value	χ^2	P value
Age group	4.37	0.037	6.25	0.012	6.49	0.011	1.45	0.229
Sex	2.91	0.088	2.52	0.112	2.50	0.114	0.61	0.436
T stage	0.08	0.783	0.30	0.586	0.02	0.894	0.31	0.581
N stage	5.30	0.021	3.68	0.055	0.16	0.689	7.18	0.007
Clinical stage	2.27	0.132	1.47	0.225	0.06	0.807	2.65	0.104
RT technique	0.93	0.334	0.28	0.595	0.10	0.750	1.19	0.276
Endoscopic surgery	5.43	0.020	8.31	0.004	5.76	0.016	2.87	0.090

Note: Bold formatting indicates statistically significant.

Abbreviations: OS, overall survival; DFS, disease-free survival; RFS, relapse-free survival; DMFS, distant metastasis-free survival; RT, radiotherapy.

Survival Analysis

The median follow-up was 63 months (29–104 months) with the following stats: 12 patients died, 8 patients experienced recurrence, and 8 patients developed distant metastases, 1 patient developed recurrence and distant metastases. In total,

0 and 14 patients in the surgery and the non-surgery groups, respectively, had a residual tumor at the end of the treatment ($P=0.001$). The surgery group had significantly higher 5-year OS (98.30% vs. 91.70%), DFS (98.30% vs. 81.40%), RFS (100.00% vs. 90.10%), and DMFS (98.30% vs. 91.30%) rates

Figure 2 Comparison of survival curves between surgery and non-surgery group in patients with early-stage nasopharyngeal carcinoma (n=156) ((A) Overall survival; (B) Disease-free survival; (C) Relapse-free survival; (D) Distant metastasis-free survival).

than did the non-surgery group ($P=0.020$, 0.004 , 0.016 , and 0.090 , respectively, Table 3 and Figure 2). Multivariate analysis showed that surgery is an independent variable influencing DFS (Figure 3).

Subgroup Analysis

In the survival analysis of 57 patients with stage I NPC, the results showed no significant difference in OS, DFS, RFS, and DMFS between the surgery and non-surgery groups (All $P>0.05$). In the survival analysis of 99 patients with stage II NPC, the surgery group had higher OS, DFS, and RFS rates than did the non-surgery group ($P=0.079$, 0.037 , and 0.077 , respectively, Figure 4). Among stage II patients with lymph node metastasis, the neck dissection group ($n=14$) had higher 5-year OS (92.90% vs. 87.10%), DFS (92.90% vs. 75.90%), RFS (100.00% vs. 90.30%), and DMFS (92.90% vs. 82.00%) than did the non-dissection group ($n=31$), but the differences were not significant (All $P>0.05$, Figure 5). In 4 of the 14 patients who

underwent endoscopic nasopharyngectomy combined with neck dissection, the postoperative pathological analysis indicated by the absence of metastatic carcinoma in the cervical lymph nodes. The representative cases are presented in Figure 6.

Changes in EBV-DNA

The average pretreatment EBV-DNA level was $0.57 \log_{10}$ copies/mL for the surgery group and $0.58 \log_{10}$ copies/mL for the non-surgery group ($P=0.967$). At the end of the treatment, all patients in the surgery group tested negative for EBV-DNA, whereas two patients in the non-surgery group tested positive for EBV-DNA.

Inter-Group Comparison of Radiation Dose and Toxic Effects

The median nasopharyngeal radiation dose was 60 Gy for the surgery group and 70 Gy for the non-surgery group, with significant difference ($P<0.001$). The incidence of

Figure 3 Forest plots depicting the multivariate association of clinicopathological characteristics with overall survival (**A**) and disease-free survival (**B**). *Indicates statistically significant.

hematologic toxicities during treatment was similar between the two groups (Figure 7). The incidence of severe oral mucositis throughout treatment was significantly different between the surgery group and the non-surgery group (37.9% vs. 54.08%, $P=0.051$). The mean weight loss at the end of treatment was 6.09 ± 3.72 kg in the surgery group and 5.22 ± 2.80 kg in the non-surgery

group, with no significant difference ($P=0.098$), as shown in Figure 8.

Discussion

The efficacy of surgery as the primary treatment modality for NPC is yet to be clarified. This study found that surgery combined with chemoradiotherapy can

Figure 4 Comparison of survival curves between surgery and non-surgery group in patients with stage II nasopharyngeal carcinoma (n=99) ((A) Overall survival; (B) Disease-free survival; (C) Relapse-free survival; (D) Distant metastasis-free survival).

improve survival in NPC patients, particularly those with stage II disease. The OS and DFS rates of the surgery group were higher than those of the non-surgery group even after propensity score matching. Multivariate Cox regression analysis also showed that surgery is an independent factor influencing the DFS of NPC patients.

Residual tumors after treatment are found in approximately 7% to 13% of NPC patients;¹² similar finding was obtained in the current study (14.9%). However, unlike previous studies, we evaluated the tumor regression rate at the end of treatment. The common types of residual tumors include local (nasopharyngeal) and regional (cervical) residual disease, with the latter occurring more frequently than the former at the end of treatment.¹³ In this study, nasopharyngeal residues (n=28) were slightly more common than cervical residues (n=22), but both of these occurred only in the chemoradiotherapy group. No

tumor residue was found in the surgery group. Although biopsy is widely recognized as the gold standard for diagnosing residual NPC, tumor tissues are difficult to extract through biopsy because 69.4% of lesions occur outside the nasopharynx.¹⁴ Thus, currently, residual tumors are primarily diagnosed using contrast-enhanced MRI.^{13,15} However, the nasopharyngeal biopsy is an ideal diagnostic modality for residual lesions protruding into the nasal cavity. Biopsied tissues can also be subjected to immunohistochemistry for several markers, such as the Ki-67 labeling index, to understand tumor proliferation.¹⁶ Positron emission tomography-computed tomography (PET-CT) is more accurate than MRI for evaluating local lesions, but also more expensive.¹⁷ Existing literature on the prognostic impact of residual NPC is relatively limited and inconclusive. One study found no significant difference in survival rates between patients with and without residual disease after NPC treatment, and therefore,

Figure 5 Comparison of survival curves between neck dissection and non-neck dissection in patients with T1-2N1 nasopharyngeal carcinoma (n=45) ((A) Overall survival; (B) Disease-free survival; (C) Relapse-free survival; (D) Distant metastasis-free survival).

recommended follow-up observation for these patients.¹⁸ In contrast, another study reported that the group with the residual disease should be actively treated as this group has lower survival rates than the group with no residual disease.¹⁹ Similarly, the group with residual disease in this study also had lower DFS, RFS, and DMFS than the group with no residual disease. T staging has been previously found to be the main influencing factor of local failure.²⁰ We also found a higher rate of residual disease of 17.5% in T2 patients, whereas it was only 7.8% in T1 patients. This implied that the risk of residual tumor is positively correlated with the tumor size.

The EBV-DNA level at three months after treatment has been reported to be associated with tumor residue,¹³ and this association could be possible

because NPC cells are a possible source of plasma EBV-DNA. Tumor residue, local recurrence, and distant metastasis are likely to occur in patients with detectable EBV-DNA even after treatment.²¹ In this study, all patients in the surgery group tested negative for EBV-DNA at the end of treatment, whereas two patients in the chemoradiotherapy group remained positive for EBV-DNA. This was probably because the surgical resection removed the nasopharyngeal and cervical lesions, and consequently, reduced the main sources of EBV.²² Wang et al also reported that the EBV clearance rate is an independent risk factor influencing NPC prognosis.²³

We previously found a higher 3-year OS rate in early NPC patients undergoing surgery than in patients undergoing chemoradiotherapy (98.44% vs. 91.98%), whereas

Figure 6 Pre- and post-operative MRI and high-definition endoscopic images. (A) Pre-operative MRI shows that the tumor is located in the right pharyngeal recess (red arrow). (B) Pre-operative MRI shows that lymph node (LN) metastases is located in the right side of the neck (red arrow). (C, D) The post-operative MRI did not show tumor residual. (E) Pre-operative endoscopic examination shows the tumor is located in the left nasopharynx. (F) Pre-operative endoscopic examination shows the tumor is located in the right nasopharynx. (G, H) The post-operative endoscopic examination images show no visible tumor residual.

there was no significant difference in DFS.⁷ Based on these findings, we extended the follow-up period in the current study and similarly found better OS and DFS rates in the surgery group than in the non-surgery group. However, in subgroup analysis, the current study found that the efficacy of surgery is similar to that of radiotherapy among stage I NPC patients, in line with another study, except that the previous study evaluated ten newly diagnosed localized stage I NPC patients, and the surgery group did not undergo postoperative chemoradiotherapy.⁸ On contrast, among the stage II patients, surgery was superior to chemoradiotherapy with respect to the DFS. Further analysis of the N1 patients showed that 14 of the 15 patients who underwent endoscopic surgery also underwent functional neck dissection. The results of the survival analysis showed better survival in those who underwent dissection, but the difference was not statistically significant. This could be because of the limited number of cases.

A previous study showed that cells from lymph node metastases could form new colonies of metastatic cancer cells in the lungs or other organs.²⁴ Hence, preventing or reducing lymph node metastases can lower the risk of tumor metastasis. Wong et al suggested that surgical treatment is more appropriate for locally advanced NPC without distant metastasis,²⁵ because the ratio of hypoxic cells increases with the size of the tumor, which significantly diminishes the radiation sensitivity of the tumor cells. These indicated that the promotion in DFS for NPC patients treated with surgery may be associated with removing radioresistant disease as compared with radiotherapy.

The incidence of severe oral mucositis in the current study was lower in the surgery group than in the non-surgery group, possibly because the former was exposed to a lower radiation dose. As the mucosa is an early responding tissue, the mucosa is associated with the overall radiation dose. The oral and

Figure 7 The hematologic toxicity between surgery and non-surgery group in patients with early-stage nasopharyngeal carcinoma (n=156, all $P>0.05$) ((A) Hemoglobin; (B) Neutrophils; (C) Platelets; (D) Leukocytes).

pharyngeal mucosa will begin to change when the dose of radiotherapy ranges from 1500 cGy to 2000 cGy, and oral and pharyngeal ulcers may occur when the dose reaches up to 3000 cGy.²⁶ Compared with conventional radiotherapy, IMRT improves the local control rate and overall survival of NPC but is also associated with a higher incidence of severe radiation-induced oral mucositis (ROM).²⁷ Furthermore, both the surgery and non-surgery groups showed similar rates of hematologic toxicity, which indicated that surgery did not increase the side effects of treatment. In addition to achieving complete resection of nasopharyngeal tumors, endoscopic nasopharyngectomy has the benefit of being minimally invasive and allowing faster postoperative recovery. Patients with cervical lymph node metastasis underwent functional neck dissection, which preserved the sternocleidomastoid

muscle, internal and external jugular veins, and accessory nerve and had no effect on postoperative blood return.

This study has some limitations. First, the possibility of bias could not be eliminated because of the retrospective study design, but propensity score matching was performed to compensate. Second, some patients were not tested for EBV-DNA. Third, due to the limited number of N1 patients who underwent endoscopic nasopharyngectomy without neck dissection, the exact prognostic impact of neck dissection for NPC remains unclear and needs to be further clarified in future studies with a larger sample size.

In conclusion, surgery combined with chemoradiotherapy may be a safe and effective treatment modality for early-stage NPC. Further, it can reduce tumor residue and the occurrence of severe ROM, and improve the clearance rate of EB virus.

Figure 8 The grade of oral mucositis and weight change between surgery and non-surgery group in patients with early-stage nasopharyngeal carcinoma (n=156, both $P>0.05$) ((A) Oral mucositis; (B) Weight change).

Ethics Statement

The Ethics Committee of The People's Hospital of Guangxi Zhuang Autonomous Region approved the study protocol, in accordance with the guidelines of the Helsinki Declaration. All patients provided written informed consent.

Author Contributions

All authors made substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; took part in drafting the article or revising it critically for important intellectual content; reviewed the manuscript and agreed on the journal to which the article will be submitted; agreed on all versions of the article before submission, during revision, the final version accepted for publication, and any significant changes introduced at the proofing stage; and agree to take responsibility and be accountable for the contents of the article.

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Disclosure

The authors report no conflicts of interest in this work.

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