Dose-escalated radiotherapy improved survival for esophageal cancer patients with a clinical complete response after standard-dose radiotherapy with concurrent chemotherapy

Wei Zhang1,.*
Yijun Luo2,*
Xiaoli Wang2
Gaohua Han1
Peng Wang1
Wei Yuan3
Sheng-Bin Dai1

1Department of Oncology, Taizhou People’s Hospital, Taizhou, Jiangsu, People’s Republic of China;
2Department of Oncology, The People’s Hospital of Jiangxi Province, Nanchang, Jiangxi, People’s Republic of China;
3Department of Cardiology, Taizhou People’s Hospital, Taizhou, Jiangsu, People’s Republic of China

Purpose: For esophageal cancer patients with a clinical complete response (cCR) after standard-dose radiotherapy with concurrent chemotherapy, data on the survival outcomes and recurrence patterns remain scarce. To evaluate the impact of dose escalation on overall survival for this subset of patients, we carried out the current investigation.

Materials and methods: Medical records of 80 esophageal cancer patients with a cCR after standard-dose radiotherapy with concurrent chemotherapy at our center from 2010 to 2014 were allocated into the standard-dose group (50.4 Gy, observation group) or the radiation dose-escalation group (59.4 Gy, control group). In this cohort study, we compared the outcomes between the 2 groups.

Results: There were no differences in patient characteristics between the 2 groups. The median recurrence-free survival and overall survival times for all patients were 38 and 54 months, respectively. Patients in the control group had significantly better 5-year recurrence-free survival rate (12% vs 0%, p=0.019) and 5-year overall survival rate (42.8% vs 21.0%, p=0.028) than the observation group. Additionally, local control rate was significantly higher in the control group (p=0.04), and ~60% of treatment failures were local failures even for patients achieving cCR after chemoradiotherapy. There were no significant differences in treatment-related toxicities between the groups.

Conclusion: The results of the current study suggest that for esophageal cancer patients with a cCR after standard-dose radiotherapy with concurrent chemotherapy, those with dose-escalated radiotherapy showed significantly better local control, recurrence-free survival, and overall survival than patients receiving 50.4 Gy radiotherapy.

Keywords: esophageal squamous cell carcinoma, radiotherapy, survival outcome, recurrence patterns

Introduction

Esophageal cancer (EC) is one of the most common cancers worldwide and has a poor prognosis due to high rates of local recurrence (LR) and distant metastasis.1,2 Squamous cell carcinoma is the most common histology of EC in Asia.3 About one-half of patients presented with locally advanced stage at the time of diagnosis.4

Definitive chemoradiotherapy (CRT) has been recommended as the optimal treatment for patients who are medically inoperable or have locally advanced stage tumor based on the results of the Radiation Therapy Oncology Group (RTOG) 85-01 trial.5 Despite curative treatments, the 5-year overall survival (OS) rate for esophageal
squaamous cell carcinoma (ESCC) patients is ~20%. The importance of optimizing radiation dose for definitive management is particularly high for EC since treatment outcomes remain poor. Attempts to improve OS by escalating the dose of radiotherapy with concurrent chemotherapy has been assessed in the INT 0123 trial. OS after 64.8 Gy dose was not superior or even lower to the 50.4 Gy. Based on those results, NCCN esophageal cancer guidelines recommend radiation doses of 50.4 Gy for definitive concurrent CRT (CCRT) in EC.

However, because of high local failure and distant failure rates after treatment, new approaches such as intensification of CRT and escalation of the radiation dose have been attempted. Some studies have revealed that high radiation doses can bring survival benefits. However, some studies showed that dose-escalated radiotherapy do not confer a benefit to EC patients and can even lead to more treatment-related toxicities. Of note, patients with ESCC receiving CRT seem to have a disparity in treatment response and outcomes. For radiosensitive EC, the standard radiation dose of 50.4 Gy is sufficient, while for the resistant patient, it is difficult to achieve a complete response (CR) even via radiation dose escalation, which may further increase treatment-related toxicities. Ma and colleagues declared that for patients with ESCC receiving CRT, the standard radiation dose of 50.4 Gy is sufficient, while for the resistant patient, it is difficult to achieve a complete response (CR) even via radiation dose escalation, which may further increase treatment-related toxicities.

Patient selection
We retrospectively reviewed the records of patients with EC who underwent CRT in 2 hospitals between January 2009 and March 2014. The criteria for enrollment were as follows: 1) All the patients had histologically proven primary ESCC and were treated with definitive CRT, and 2) only patients who achieved a CR after planned radiation (50.4–56 Gy) were included in the study. In addition, we excluded patients with thoracic esophageal squamous cell carcinoma who did not receive sufficient CCRT doses (<50.4 Gy). Ultimately, 80 patients who achieved a CR were recruited for this study, and we categorized them into 2 groups based on the radiation doses to compare their treatment outcomes: group 1 (patients receiving CCRT <56 Gy) and group 2 (patients receiving CCRT ≥59.4 Gy). The median total dose and fraction size of RT in groups 1 and 2 were 52 Gy (50.4–56 Gy, fraction size: 1.80/2.0 Gy) and 62.0 Gy (59.4–64.8 Gy, fraction size: 1.80/2.0 Gy), respectively. The study selection process is summarized in Figure 1.

CR for the primary tumor was defined as the absence of evidence of a residual tumor on esophageal endoscopy; on computed tomography (CT) of the neck, chest, and abdomen; and on endosonography of the esophagus. CR for lymph nodes was defined according to the Response Evaluation Criteria in Solid Tumors guidelines. Pretreatment evaluations included barium tests; esophageal endoscopy; CT of the neck, chest, and abdomen; and endosonography of the esophagus. Patients were staged according to the 2010 (7th) American Joint Committee on Cancer staging criteria. Tumor length was measured by esophagography before treatment. This study was approved by the Institutional Review Board of Taizhou People’s Hospital (IRB 20170401) and the People’s Hospital of Jiangxi Province (IRB160815), and all patients had signed written informed consent prior to the study.

CRT
Radiotherapy was administered as 3-dimensional conformal radiotherapy or intensity-modulated radiation therapy. A conventional fractionation schedule (daily 1.8–2.0 Gy per fraction, 5 days per week) was used in all patients. The delineation of clinical target volumes was based on CT, barium esophagogram, endoscopic examination, and positron emission tomography (PET) imaging. The clinical target volumes (CTVs) were a 3-cm proximal and distal margin, and a 0.5–0.8-cm radial margin was added to the GTV. The planning target volume (PTV) encompassed a 1-cm proximal...
and distal margin and a 0.5-cm radial margin based on CTVs. The nodal CTV was defined by a 0.5–1-cm expansion around the nodal GTV. The PTV was the CTV plus a uniform 0.5-cm expansion margin. In the dose-escalation group, 37 patients with CR received dose escalation after standard radiotherapy (50.4 Gy), the escalated dose was from 9 to 14.4 Gy, and the total dose was ranged from 59.4 to 64.8 Gy.

All patients received platinum-based chemotherapy concurrently with irradiation. The additional drug was either a 5-fluorouracil or taxane (docetaxel or paclitaxel).

**Definition of event**

Treatment-induced toxicities were assessed using the National Cancer Institute Common Terminology Criteria for Adverse Events (version 4.0). Severe adverse events were defined as toxicity grade ≥3. Survival analyses were performed for recurrence-free survival (RFS) and OS. OS was measured from the first day of treatment to the date of death from any cause or the last follow-up, whichever occurred first. RFS was defined as the first evidence of any treatment failure. The patterns of treatment failure were defined according to the initial site of recurrence. Locoregional recurrence was defined as a recurrence occurring in the radiation field. Recurrences at any other sites were considered as distant recurrences.

**Follow-up**

All patients were examined weekly during RT to monitor treatment toxicities and their general condition. Routine evaluations included physical exam, hematologic and biochemical profiles, and esophagography. Follow-up evaluations were performed after the completion of all treatment, every 3 months for 2 years, and then every 6 months thereafter. The follow-up evaluations included physical examination, esophagography, esophagogastroduodenoscopy, chest and abdominal CT, and

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**Figure 1** Study flow diagram.

**Abbreviations:** DCRT, definitive chemoradiotherapy; cCR, clinical complete response; CRT, chemoradiotherapy.
PET when available. Other necessary examinations were conducted according to the clinical situation.

**Statistical analysis**

All statistical analyses were carried out using SPSS (version 20.0; IBM Corporation, Armonk, NY, USA). $\chi^2$ test was performed to compare the difference of patients’ characteristics. The rates of OS and RFS were calculated by the Kaplan–Meier method, with comparison using a log-rank test. All statistical tests were 2-tailed, and $p<0.05$ was considered statistically significant.

**Results**

**Patient characteristics**

The patient characteristics are shown in Table 1. In total, 58.8% of patients were male, with a median age of 60, while 41.2% were female, with a median age of 67.5, and 43 patients were in the standard-dose group and 37 patients were in the dose-escalation group. Notably, more patients with stage III disease were included in the high-dose group, although the difference was not significant (62.2% vs 46.5%, $p=0.162$). The median follow-up exceeded 54 months, with a maximum follow-up of 91 months. Patients in the 2 groups were similar with respect to age, sex, Karnofsky performance status, tumor location, tumor length, and clinical stage ($p>0.05$ in all cases).

**Survival**

The median RFS of the patients in the conventional group and dose-escalation group were 35 and 44 months, respectively. Also, the 3- and 5-year RFS rates were 70.3% and 12% in the radiation dose-escalation group, which were significantly

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Abbreviations: KPS, Karnofsky performance score; 3D-CRT, 3-dimensional conformal radiotherapy; IMRT, intensity-modulated radiotherapy; RT, radiotherapy.
Dose-escalated radiotherapy improved survival for EC patients

higher than those in the standard-dose group (42% and 0%, \( p = 0.019 \)).

The median OS was 48 months (conventional group) and 59.8 months (dose-escalation group) for the 2 groups, respectively. The 3- and 5-year OS rates were 75.3% and 21% for patients in the conventional group and 87.7% and 42.8% for those in the dose-escalation group, respectively. The 5-year OS rates seemed higher in the high-dose group (42.8%) than in the conventional-dose group (21%), which showed a significant difference (\( p = 0.028 \)). The estimated Kaplan–Meier curves for OS and RFS are shown in Figures 2 and 3, respectively.

Patterns of failure

Table 2 depicts the rates and sites of the first recurrence. A total of 55 patients (68.8%) experienced treatment failures, including locoregional failure (LRF) alone in 29 patients (36.3%), distant metastasis alone in 19 (23.8%), and combined LRF and distant metastasis in 7 (8.8%); 25 (31.3%) had no evidence of disease. The incidence of LRF was significantly lower in the control vs the observation group (64.5% in the observation group vs 37.5% in the control group, respectively; \( p = 0.04 \)). The distant failure rate (25.8% vs 45.8%) between 2 groups did not reach statistical significance.

Treatment-related toxicity

The toxicities induced by CCRT were mainly skin reaction, esophagitis, hematologic toxicity, and radiation pneumonia. Treatment-induced toxicities for all patients are detailed in Table 3. No treatment-related deaths were found. As for hematological adverse events, events graded \( \geq 3 \) during CRT were seen in 18/43 patients (41.8%) and 21/37 patients (56.7%), respectively. On the other hand, nonhematological side effects, radiation esophagitis, were seen at a high frequency. The incidences of grade \( \geq 3 \) esophagitis was higher in the control group than in the observation group (8/37, 21.6% vs. 4/43, 9.3%). The rate of pneumonitis was 4 (10.8%) in the standard-dose group, whereas it was 10 (25.0%) in the dose-escalation group. No significant differences were observed between the 2 groups in the rates of other toxicities.

Discussion

The question of optimal radiation dose in the treatment of locally advanced EC has long been a topic of investigation. Since the establishment of 50.4 Gy as the standard dose in the management of locally advanced EC (RTOG 94-05), there have been multiple attempts at optimization of radiation dose.

A great number of studies were done to investigate the potential benefits of higher RT dose. The study of Zhang

Figure 2 OS in the standard-dose group (group 1: patients receiving CCRT <56 Gy) and dose-escalation group (group 2: patients receiving CCRT \( \geq 59.4 \) Gy).

Note: \( p = 0.006 \).

Abbreviations: OS, overall survival; CCRT, concurrent chemoradiotherapy.
et al. found that patients in the higher dose group (>51 Gy) had a better 3-year local control rate (36% vs 19%) and a disease-free survival rate (25% vs 10%) than those in the low-dose group (≤51 Gy). No significant differences were found between high- and low-dose groups in OS (p=0.054). A study was conducted by Suh et al. to compare the prognosis between the 2 groups (the patients received a dose of ≤60 or >60 Gy). The high-dose group experienced a significantly lower LRF rate (32% vs 69%, p<0.01). However, no significant difference was noted between the 2 groups regarding OS (p=0.26). In line with these findings, the results from a study by He et al. showed that the high-dose group had a significantly lower local failure rate (17.9% vs 34.3%, p=0.024) than the low-dose group, but the OS was not significantly different (p=0.617). However, some studies claimed that high-dose radiotherapy may provide a survival benefit. In the retrospective analysis by Kim et al., the correlation between radiation dose and OS for EC patients treated with definitive CRT was investigated. The results suggested that patients who underwent high-dose radiotherapy had significantly better survival than patients receiving <60 Gy RT when treated with concurrent chemotherapy. Similarly, a study conducted by Chen et al. found that a dose higher than that used for standard radiotherapy may lead to better survival for ESCC patients undergoing CCRT. These conflicting results revealed that the optimal definitive radiation dose should be managed on an individual basis.

Patients with ESCC receiving CRT seem to show a disparity in treatment response. For the radiosensitive EC,
Dose-escalated radiotherapy may provide survival benefit. Conversely, it is very difficult to confer a benefit to a radiation-resistant patient, and dose escalation even may lead to increase the radiation-related toxicities. At present, data on the survival outcomes and recurrence patterns of patients with EC who achieved a CR after standard-dose radiotherapy remain scarce. In this study, we investigated the survival outcomes and recurrence patterns in patients who achieve cCR after standard radiation dose radiotherapy. Furthermore, we also explored whether the higher radiation dose therapy could confer benefits in local control or OS in those patients.

To the best of our knowledge, this is the first retrospective analysis on whether the higher radiation dose could confer benefits for EC patients who achieved a CR after standard-dose radiotherapy (50.4 Gy). There are 2 principal findings in this study. First, we demonstrated that at least 60% of ESCC patients who achieve cCR after CRT developed disease recurrences, with LR being the predominant failure pattern. However, the LR rate in the control group was significantly lower than that in the standard-dose group. Second, as shown in Figure 2, the survival time and the RFS in the control group were longer than that in the observation group, which were statistically different (p<0.05). The results of the current study suggest that the subgroup of patients who underwent dose escalation had significantly better LC and OS than patients receiving 50.4 Gy when treated with concurrent chemotherapy. Furthermore, the CRT-induced toxicities were also acceptable in the dose-escalation group.

In this scenario, it is our institutional policy to perform a dose-escalation radiotherapy aimed at achieving a better tumor control. Unfortunately, LR rates remained suboptimal (37.5%) even after dose-escalation radiotherapy. Such rates are significantly higher than those identified (<10%) in patients who underwent resection.18–20 McLoughlin et al21 suggested that after chemoradiation, cCR was not substantially predictive of pathological CR. Similarly, a study conducted by Stiles et al22 also revealed that a CR on posttreatment PET scan predicts but should not be assumed to be synonymous with complete pathologic response in EC patients. Accordingly, only 23%–40% of cCR patients actually have no residual cancer on pathological examination.23–25 In summary, although the lesion has reached clinical remission after CRT, there may still be residual lesions in the primary site. Therefore, dose-escalation radiotherapy may eliminate the residual lesion and improve the local control and OS.

This study has a few limitations that need to be considered when interpreting the results. The study is limited by its retrospective nature, and we cannot account for potential selection bias, which may limit the generalizability of our results. Second, our sample size is small, and further independent studies in larger populations are needed to confirm and validate our results. Finally, it is possible that treatment-related toxicities were underestimated due to the study’s retrospective setting.

**Conclusion**

Taken together, our results indicate that dose-escalated radiotherapy yields more favorable survival outcomes in EC patients who achieved a cCR after standard-dose radiotherapy. Further independent studies in larger populations are needed to confirm and validate our results.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


