Definitive radiotherapy with or without chemotherapy for clinical stage T4N0-1 non-small cell lung cancer

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Purpose: To determine failure patterns and survival outcomes of T4N0-1 non-small cell lung cancer (NSCLC) treated with definitive radiotherapy.

Materials and Methods: Ninety-five patients with T4N0-1 NSCLC who received definitive radiotherapy with or without chemotherapy from May 2003 to October 2014 were retrospectively reviewed. The standard radiotherapy scheme was 66 Gy in 30 fractions. The main concurrent chemotherapy regimen was 50 mg/m2 weekly paclitaxel combined with 20 mg/m2 cisplatin or AUC 2 carboplatin. The primary outcome was overall survival (OS). Secondary outcomes were failure patterns and toxicities.

Results: The median age was 64 years (range, 34 to 90 years). Eighty-eight percent of patients (n = 84) had an Eastern Cooperative Oncology Group performance status of 0–1, and 42% (n = 40) experienced pretreatment weight loss. Sixty percent of patients (n = 57) had no metastatic regional lymph nodes. The median radiation dose was EQD2 67.1 Gy (range, 56.9 to 83.3 Gy). Seventy-one patients (75%) were treated with concurrent chemotherapy; of these, 13 were also administered neoadjuvant chemotherapy. At a median follow-up of 21 months (range, 1 to 102 months), 3-year OS was 44%. The 3-year cumulative incidences of local recurrence and distant recurrence were 48.8% and 36.3%, respectively. Pretreatment weight loss and combined chemotherapy were significant factors for OS. Acute esophagitis over grade 3 occurred in three patients and grade 3 chronic esophagitis occurred in one patient. There was no grade 3–4 radiation pneumonitis.

Conclusion: Definitive radiotherapy for T4N0-1 NSCLC results in favorable survival with acceptable toxicity rates. Local recurrence is the major recurrence pattern. Intensity modulated radiotherapy and radio-sensitizing agents would be needed to improve local tumor control.

Keywords: Carcinoma, Non-small-cell lung, Radiotherapy, Definitive, T4, Local tumor control, Survival

Introduction

Before the introduction of chemotherapy, stage III non-small cell lung cancer (NSCLC) had poor survival outcomes with single local treatments such as surgery or radiotherapy (RT). Addition of induction chemotherapy to RT significantly improved survival [1,2], and subsequent trials reported that concurrent chemo-radiotherapy (CRT) was better than sequential CRT.
Combined with surgery, induction chemotherapy also improved survival for NSCLC patients [5]. Moreover, the addition of neoadjuvant RT to chemotherapy was suggested to improve mediastinal downstaging and the rate of complete resection [6]. As a result, management for stage III NSCLC is currently based on resectability. For resectable patients, surgery is performed with neoadjuvant CRT or adjuvant chemotherapy and RT. On the other hand, concurrent CRT is considered the standard therapy for unresectable stage III NSCLC. However, determination of resectability is difficult in "potentially resectable" subgroups such as stage IIIA, which has led to controversy in the management of stage IIIA NSCLC patients.

Stage IIIA subgroups have markedly different clinical characteristics despite having a similar prognosis, and include patients with N2 disease (T1-3N2M0) and patients with either extensive local invasion or ipsilateral additional tumor nodules (T4N0-1M0). It is appropriate to assume that the stage III-N2 and T4N0-1 groups would need different treatment approaches.

A previous prospective trial compared the survival outcomes of trimodality treatment (concurrent CRT induction with an RT dose of 45 Gy followed by surgical resection) and definitive CRT with a total RT dose of 61 Gy in stage IIIA-N2 NSCLC [7]. There was no significant 5-year overall survival (OS) advantage for surgery after CRT, despite improved progression free survival (PFS). In exploratory analysis, trimodality approaches showed significantly better OS than bimodality approaches if lobectomies were performed. However, these results were only for stage IIIA-N2 disease and not stage T4N0-1 disease.

To the best of our knowledge, there has only been one study to date of the treatment outcomes of stage T4N0-1 NSCLC patients treated with definitive CRT [8]. Although the results of that report showed comparable survival with surgery, some physicians still advocate surgery combined with chemotherapy as the only treatment capable of achieving long-term survival in this subgroup [9].

In the absence of phase III studies comparing surgery to RT as a local treatment, additional information about this particular subgroup with extensive local invasion and minimal nodal disease is crucial. In contrast to the single study on definitive CRT, there have been many reports on the surgical outcomes for stage T4N0-1 patients [10-12]. We here evaluated the survival outcomes of definitive RT with or without chemotherapy in stage T4N0-1 NSCLC patients. Furthermore, by evaluating failure patterns, especially local failure and toxicities, we identified additional strategies for improving patient survival.

### Materials and Methods

1. Patients

Between May 2003 and October 2014, 102 patients with T4N0-1 disease received definitive CRT at Asan Medical Center. After excluding seven patients who ended treatment with an RT dose of 50 Gy, we retrospectively reviewed 95 patients. The first author (Kim YJ) reviewed all patients and tumor stage was determined according to the American Joint Committee on Cancer (AJCC) 7th edition TNM stage classification. Known prognostic factors such as Eastern Cooperative Oncology Group performance status (ECOG-PS) and weight loss were also analyzed. Pretreatment weight loss was defined as 5% or more weight loss over the previous 6 months.

Clinical staging work-ups included pathologic confirmation, chest computed tomography (CT), 18-fluoro-deoxyglucose positron emission tomography computed tomography (18\(^{\text{FDG-PET-CT}}\)), brain magnetic resonance imaging, and bronchoscopy. Diagnostic thoracotomy was not mandatory and the T4 status was mainly evaluated on imaging. Lymph nodes with a diameter ≥1 cm and definite FDG uptake were classified as clinically malignant. Otherwise, pathologic confirmation was performed whenever possible.

2. Radiotherapy

Planning CT was obtained in a supine, arms-raised position using a 2.5 mm-slice thickness. Four-dimensional CT (4DCT) images reflecting respiratory motion were collected from January 2005 onwards.

Gross tumor volume (GTV) of the primary lung lesion delineation was performed using the lung setting (W = 1,700, L = −300), while GTV of the hilar and mediastinal lymph nodes was delineated in a mediastinal setting (W = 600, L = 40). For patients with 4DCT, the internal target volume was also contoured. Clinical target volume (CTV) was delineated for patients with a relatively small GTV by a 5-mm expansion from the GTV. For patients with primary lung lesion invading mediastinum, we also included the involved mediastinal lymph node areas to the CTV. For bulky tumors, there was no CTV.

The planning target volume (PTV) margin was 7 mm radially and 10 mm longitudinally. In patients treated with induction chemotherapy, the postchemotherapy volume was defined as the GTV. Organs at risk include healthy lung tissues, the esophagus, the spinal cord, and the heart.

The standard RT scheme was 66 Gy with 2.2 Gy per fraction. After irradiating with 44 Gy or 52.8 Gy, repeated CT simulation was routinely performed. We performed chest X-rays (CXRs)
every week. Patients who showed dramatic volume changes were requested for earlier repeated CT simulation. Treatment verification was performed by weekly kV imaging guidance using set-up correction based on carina and bony anatomy.

Three-dimensional conventional RT was used as a routine policy, and intensity modulated radiotherapy (IMRT) was considered when the following normal organ constrains were difficult to meet; maximal dose ($D_{\text{max}}$) for spinal cord <50 Gy, mean lung dose (MLD) <20 Gy, volume of lung receiving at least 20 Gy (V20) <30%, mean esophagus dose <35 Gy.

3. Chemotherapy
The main concurrent chemotherapy regimen was 50 mg/m$^2$ weekly paclitaxel combined with 20 mg/m$^2$ cisplatin or AUC 2 carboplatin. Although concurrent CRT is a standard format for definitive CRT, patients with very large tumors and a high risk of radiation-related toxicities were initially treated with induction chemotherapy for volume reduction. The standard chemotherapy regimen was 2 cycles of 1,000 mg/m$^2$ gemcitabine (days 1 and 8) with 70 mg/m$^2$ cisplatin (day 1) in sequential schedules every 3 weeks. Patients older than 75 years old, with a poor performance status or with poor lung function were recommended for definitive RT alone.

4. Follow-up and toxicity scoring
During definitive CRT, all patients were interviewed weekly with a complete blood count test and CXR. Patients were routinely followed up by a radiation oncologist and/or medical oncologist with chest CT and CXR 1 month after treatment, every 3 months during the first 2 years, and every 6 months until 5 years after treatment.

Locoregional recurrence (LR) was defined as primary tumor and regional lymph node progression. We also investigated if the recurrence occurred in radiation field or not. Distant metastasis (DM) was defined as intrapulmonary recurrence outside the radiation field or in ipsilateral different lobes, the contralateral lung, malignant pleural effusion, pleural metastasis, or other distant organs. Although we used AJCC 7th edition TNM stage classification, we considered ipsilateral different lobe metastasis as DM. We accepted that the patients with ipsilateral separate lung nodule had better prognosis than those with the other M1 factors in AJCC 6th edition. However, we assumed that there must be a difference between the assessment of initial stage and recurrence. One of the aims of the present study was to evaluate the local tumor control of definitive RT. If intrapulmonary recurrence outside the radiation field had been considered as LR, we would fail to show the true effect of definitive RT.

Adverse events that occurred during treatment and within 3 months after CRT were defined as acute toxicities. The severity of toxicity was assessed according to the National Cancer Institute's Common Terminology Criteria for Adverse Events (CTCAE) ver. 4.03, for hematologic complications, radiation esophagitis, and pneumonitis.

5. Statistics
Follow-up durations were calculated from the date that treatment started. Survival was calculated using Kaplan-Meier method from the date treatment started until death. The p-values were determined by the log-rank test for univariate analysis and with a Cox proportional hazard model for multivariate analysis. Significant factors revealed by univariate analysis ($p < 0.05$) were included for multivariate analysis. Cumulative incidence of LR and DM were also calculated using Kaplan-Meier method (one minus survival). A Student t-test was used to assess the relationship between chemotherapy and acute toxicities. All statistical tests were performed using SPSS ver. 21.0 (IBM, Armonk, NY, USA).

Results
1. Patient characteristics
The characteristics of the 95 patients are summarized in Table 1. The median follow-up time was 21 months (range, 1 to 102 months). Fifteen patients (16%) had invasive confirmation of T4 status, such as thoracotomy. The most common reason for T4 status was pulmonary artery or vein invasion in 30 patients (32%) (Table 2).

2. Treatment characteristics
The majority of patients ($n = 68, 72\%$) received a total RT dose of 66 Gy in 30 fractions; the EQD2 (equivalent dose in 2 Gy fractions) was 67.1 Gy ($\alpha/\beta$ 10 Gy for tumor). The median radiation dose was also EQD2 67.1 Gy (range, 56.9 to 83.3 Gy). Five patients were irradiated with 66 Gy, and four patients received 60 Gy with a fraction size of 2 Gy. The remaining patients received variable doses of 50–77 Gy in fraction sizes of 1.8–10 Gy. One patient was treated with hypofractionated RT using the Arc technique with a dose regimen of 50 Gy/5 fractions, EQD2 83.3 Gy. This patient had only spine invasion without regional node metastasis.

IMRT was applied to nine patients (9\%) (Table 3). Of these, eight patients had vertebral invasion, and IMRT was used to save their spinal cord. The remaining patient did not have...
vertebral invasion but had a primary lung lesion widely abutted to the thoracic spines, so he also received IMRT to adjust the spinal cord dose. A 64-year-old man diagnosed with cT4N0 was treated with hypo-fractionated RT with a fraction size of 10 Gy up to 50 Gy, and another patient was treated with

Table 1. Patients’ characteristics

| Characteristic                  | No. (%) |
|--------------------------------|---------|
| Age (yr), median (range)        | 64 (34–90) |
| Gender                         |         |
| Male                           | 87 (92) |
| Female                         | 8 (8)   |
| ECOG performance status        |         |
| 0                              | 11 (12) |
| 1                              | 73 (77) |
| 2                              | 9 (9)   |
| 3                              | 2 (2)   |
| Weight loss                    |         |
| High (≥5%)                     | 40 (42) |
| Low (<5%)                      | 55 (58) |
| TNM stage                      |         |
| cT4N0M0                        | 57 (60) |
| cT4N1M0                        | 38 (40) |
| Histology                      |         |
| Squamous cell carcinoma        | 67 (71) |
| Adenocarcinoma                 | 17 (18) |
| Large cell carcinoma           | 4 (4)   |
| Unspecified                    | 7 (7)   |
| Chemotherapy                   |         |
| None                           | 24 (25) |
| Sequential                     | 3 (3)   |
| Induction chemotherapy + concurrent | 13 (14) |
| Concurrent                     | 55 (58) |

ECOG, Eastern Cooperative Oncology Group.

Table 2. Reasons for T4

| Involvement                    | No. (%) |
|--------------------------------|---------|
| Great vessels                  | 51 (54) |
| Pulmonary artery/vein          | 30 (32) |
| Aorta                          | 16 (17) |
| Subclavian artery/brachiocephalic vein | 5 (5)   |
| Superior vena cava             | 3 (3)   |
| Mediastinum                    | 27 (28) |
| Carina/trachea                 | 13 (14) |
| Spine                          | 9 (9)   |
| Heart                          | 7 (7)   |
| Nerves                         |         |
| Recurrent laryngeal nerve      | 5 (5)   |
| Vagus nerve                    | 1 (1)   |
| Esophagus                      | 1 (1)   |

* Nineteen patients had two or more reasons for T4.

Table 3. Characteristics of patients who received intensity-modulated radiotherapy

| Case no. | Age | Reason for T4 | N stage | EQD2 (Gy) | Fraction size | Chemotherapy | LR | LRFS | DM | DMFS | Alive | OS | G3 acute toxicities |
|----------|-----|---------------|---------|-----------|--------------|--------------|----|------|----|------|--------|----|-------------------|
| 1        | 64  | Spine         | 0       | 50        | 10           | 66           | N  | Y    | N  | N    | 83.3   | N  | Y                 |
| 2        | 44  | Mediastinum   | 0       | 50        | 20           | 66           | N  | Y    | N  | N    | 86.6   | N  | N                 |
| 3        | 42  | Spine         | 0       | 66        | 2.0          | 66           | Y  | Y    | N  | N    | 67.1   | Y  | Y                 |
| 4        | 73  | Spine         | 0       | 66        | 2.2          | 66           | Y  | N    | Y  | N    | 67.1   | Y  | N                 |
| 5        | 52  | Spine         | 0       | 66        | 2.2          | 66           | Y  | N    | Y  | N    | 67.1   | Y  | N                 |
| 6        | 58  | Spine         | 0       | 66        | 1.8–2.2      | 66           | Y  | N    | Y  | N    | 67.5   | Y  | N                 |
| 7        | 35  | Spine         | 0       | 66        | 2.0          | 66           | Y  | Y    | N  | N    | 66.6   | Y  | N                 |
| 8        | 71  | Spine         | 0       | 66        | 2.2          | 66           | Y  | Y    | N  | N    | 66.2   | Y  | N                 |
| 9        | 70  | Spine         | 0       | 66        | 2.2          | 66           | N  | Y    | N  | N    | 66.2   | Y  | N                 |

RT, radiotherapy; EQD2, equivalent dose in 2 Gy fractions; LR, locoregional recurrence; LRFS, locoregional recurrence-free survival; DM, distant metastasis; DMFS, distant metastasis-free survival; OS, overall survival; SIB, simultaneous integrated boost.
simultaneous integrated boost. The PTV<sub>CTV</sub> received 56 Gy in 28 fractions, while the PTV<sub>GTV</sub> received 61.6 Gy. LR occurred in three patients (crude rate, 30%), and four patients experienced DM (crude rate, 44.4%). OS ranged from 10 to 99 months, and G3 acute toxicities occurred in two patients with neutropenia, esophagitis and anemia. Hematologic toxicities were more likely with concurrent chemotherapy, and the patient with esophagitis initially had a lesion mostly closed to esophagus.

The median RT duration was 44 days (range, 14 to 103 days). Three patients had over 90 days of RT. Initially, the aim of CRT in these patients was induction for surgical resection. However, they remained inoperable despite preoperative CRT and decided to receive additional RT to a definitive dose. There were some extended time intervals in three patients to assess operability.

A total of 71 patients received chemotherapy; 55 patients (78%, 55/71 patients) were treated with concurrent CRT alone while 13 patients (18%, 13/71 patients) also received induction chemotherapy. The remaining patients received sequential chemotherapy (4%, 3/71 patients). The most common CRT regimen was 50 mg/m<sup>2</sup> weekly paclitaxel combined with 20 mg/m<sup>2</sup> cisplatin (n = 48, 71%). Carboplatin (AUC 2) replaced cisplatin in 14 patients (15%). The remaining patients received cisplatin combined with gemcitabine (n = 3) and etoposide (n = 3).

The most common regimen in induction/sequential chemotherapy was 1,000 mg/m<sup>2</sup> gemcitabine (days 1 and 8) with 70 mg/m<sup>2</sup> cisplatin (day 1) (n = 13, 81%) although the number of cycles varied (range, 2 to 4 cycles).

![Fig. 1. Overall survival (OS) of 95 T4N0-1M0 patients (Kaplan-Meier).](image-url)

### Table 4. Univariate and multivariate analysis for OS

| Variable                        | No. | Univariate |                |          | Multivariate |                |          |
|---------------------------------|-----|------------|----------------|----------|--------------|----------------|----------|
|                                |     | Median OS (mo) | SD | 95% CI | p-value | HR | 95% CI | p-value |
| Age (yr)                        |     |             |     |        |         |                 |          |
| <60                             | 30  | 27         | 5.91 | 15.42–38.58 | 0.868 | - | - |
| ≥60                             | 65  | 31         | 10.83 | 9.78–52.22 | - | - |
| ECOG performance status         |     |             |     |        |         |                 |          |
| 0                               | 11  | 83         | 17.79 | 148.12–117.88 | 0.059 | - | - |
| 1                               | 73  | 27         | 5.52 | 16.19–37.81 | - | - |
| 2                               | 9   | 29         | 15.17 | 0.00–58.73 | - | - |
| 3                               | 2   | 3          | - | - | - | - |
| Weight loss                     |     |             |     |        |         |                 |          |
| ≥5%                             | 40  | 17         | 2.64 | 11.84–22.17 | 0.000 | 0.38 | 0.23–0.64 |
| <5%                             | 55  | 73         | 9.82 | 53.75–92.25 | - | - |
| Histology                       |     |             |     |        |         |                 |          |
| Squamous cell carcinoma         | 67  | 29         | 5.47 | 18.29–39.71 | 0.788 | - | - |
| Adenocarcinoma                  | 17  | 47         | 13.05 | 21.43–72.57 | - | - |
| Large cell carcinoma            | 4   | 21         | 12.00 | 0.00–44.52 | - | - |
| Unspecified                     | 7   | 27         | 14.40 | 0.00–55.23 | - | - |
| N stage                         |     |             |     |        |         |                 |          |
| N0                              | 57  | 34         | 10.90 | 12.63–55.37 | 0.156 | - | - |
| N1                              | 38  | 23         | 5.19 | 12.83–33.18 | - | - |
| Chemotherapy                    |     |             |     |        |         |                 |          |
| Any                             | 71  | 32         | 13.78 | 5.00–59.00 | 0.042 | 1.87 | 1.05–3.35 |
| None                            | 24  | 23         | 5.58 | 12.06–33.95 | 0.045 | - | - |

OS, overall survival; SD, standard deviation; CI, confidence interval; HR, hazard ratio; ECOG, Eastern Cooperative Oncology Group.
percent of patients (n = 24) received definitive RT alone due to old age, poor ECOG-PS, and/or poor lung function.

3. Overall survival and failure patterns

With a median follow-up time of 21 months (range, 1 to 102 months), the median OS was 23 months (range, 1 to 102 months) and the 3-year OS was 44% (Fig. 1). Pretreatment weight loss and a combination of chemotherapy modalities were significant factors in OS (Table 4). Other factors, such as age (<60 years vs. ≥60 years), ECOG-PS, histologic type, and nodal stage (N0 vs. N1), did not show significance.

Excluding two patients who were followed up at another hospital, a total of 93 patients were included in the failure pattern analysis. The cumulative incidence of LR is shown in Fig. 2. The 3-year cumulative incidence of LR was 48.8%, and LR incidence reached a plateau after 3 years. The 3-year cumulative incidence of DM was 36.3% (Fig. 3). The incidence of DM did not increase after 2 years.

The major failure pattern was LR (n = 39, 42%), especially isolated in-field recurrence (n = 23, 25%) (Table 5). One patient experienced isolated out-field recurrence in the contralateral hilar lymph node after 25 months. This patient received salvage RT and remained in disease-free status. DM occurred in 28 patients (29%), and over half of these patients were also diagnosed with LR (n = 15, 16%).

4. Toxicities

Thirteen patients (14%) had acute toxicities above grade 3 (Table 6). There were 16 total events due to one patient who experienced neutropenia, anemia and thrombocytopenia and another patient with neutropenia and esophagitis. All acute toxicities resolved and did not proceed to chronic complications. Among patients with acute toxicities, 10 received concurrent CRT (with or without induction chemotherapy) and the rest were treated with RT alone. However, there was no statistically significant association between concurrent chemotherapy and acute toxicities.

Grade 3 chronic thrombocytopenia was present in two

![Fig. 2. Cumulative incidence of local recurrence (LR) of 93 T4N0-1M0 patients (Kaplan-Meier).](image1)

![Fig. 3. Cumulative incidence of distant metastasis (DM) of 93 T4N0-1M0 patients (Kaplan-Meier).](image2)

| Table 5. Patterns of failure (n = 93) | No. of patients (%) |
|--------------------------------------|---------------------|
| Recurrence                           |                     |
| No                                   | 42 (45)             |
| Yes                                  | 51 (55)             |
| Locoregional failure                 | 39 (42)             |
| Isolated in-field failure            | 23 (25)             |
| Isolated out-field nodal failure     | 1 (1)               |
| Locoregional failure + distant failure| 15 (16)            |
| Distant failure                      | 28 (29)             |
| Isolated distant failure             | 13 (13)             |
| Locoregional distant failure + distant failure | 15 (16) |

| Table 6. ≥Grade 3 toxicities         |                      |
|--------------------------------------|---------------------|
| Toxicity                             | Acute p-value       |
|                                      | CRT RT alone        |
| Hematologic                          |                      |
| Neutropenia                          | 4 1 0               |
| Anemia                               | 5 2 0               |
| Thrombocytopenia                     | 1 0 2               |
| Esophagitis                          | 3 0 1               |
| Pneumonitis                          | 0 0 0               |
| Total events                          | 13 3 0.697 3        |
| Total patients                       | 10 3 3              |

CRT, chemo-radiotherapy; RT, radiotherapy.
patients, and one patient had grade 3 esophagitis. Two patients received concurrent CRT and one patient was treated with induction chemotherapy followed by concurrent CRT. There were no grade 4 toxicities.

**Discussion and Conclusion**

Although the standard treatment for most stage IIIA NSCLC patients is concurrent CRT, there is still controversy over whether surgery has a role in treatment. A prospective study that compared induction concurrent CRT followed by surgery and definitive CRT only included IIIA-N2 disease patients [7]. Although there were no significant difference in 5-year OS in either arms (trimodality 27.2% vs. bimodality 20.3%, p = 0.24), 5-year PFS was significantly improved by trimodality treatment (22.4% vs. 11.1%, p = 0.017). After additional matching analysis, the OS was significantly improved in the surgical arm if a lobectomy was performed (36.1% vs. 17.8%, p = 0.002). However, for pneumonectomy, there was a trend towards poorer OS in the surgical arm (21.9% vs. 23.6%, p = not significant).

Although stage IIIA-N2 and T4N0-1 NSCLC patients have similar prognoses, they have different clinical characteristics. T4N0-1 patients have extensive local tumor burden, while N2 disease has less primary tumor burden, with invasion of mediastinal lymph nodes. We expect that T4N0-1 disease would be more difficult to resect than stage IIIA-N2 disease, although there is no definite evidence for this. According to Garrido et al. [11], 22.7% of T4N0-1 patients and 10.9% of IIIA-N2 patients were unresectable, even with induction chemotherapy. These patients therefore had a delay in making other treatment choices. The three cases of stage T4N0-1 in the present study had treatment periods above 90 days. Prolonged RT periods lead to concerns about reduced survival.

Achievement of complete resection is an important factor for survival. When patients were treated with surgery alone, the complete resection rates were 37.5%–46.2% and could be improved to 65.9% with induction chemotherapy [11,12]. However, approximately 30% of patients still needed adjuvant treatment. Despite the effectiveness of adjuvant RT in incompletely resected NSCLC patients, it can be a significant burden [13]. Moreover, surgical resection can cause considerable treatment-related mortality. Albain et al. [7] reported that 7.9% of NSCLC patients (n = 16) died of surgery-related causes, while 2.1% (n = 4) in definitive CRT arm died of treatment related causes. Some surgical series have a reported mortality of up to 19.7% [14]. For this reason, we assumed that definitive CRT would be a better choice for T4N0-1 patients.

Only one study, Reymen et al. [8] has evaluated the treatment outcomes of definitive CRT in T4N0-1 patients, although there are substantial surgical data for this disease [10,11,14] (Table 7). Reymen et al. [8] reported on the long-term survival of stage T4N0-1 and single station IIIA-N2 NSCLC patients treated with definitive CRT. In 42 T4N0-1 patients in that study followed up at a median of 48 months, the median OS was 34 months, with a 3-year OS of 44% and a 5-year OS of 25%, similar to our present study (median OS 23 months, 3-year OS 44%, 5-year OS 36.3%). Compared to previous surgical reports, definitive CRT showed comparable results with complete resection. Completely resected T4N0-1 patients showed a 3-year OS of 41.9%–71.4% and a 5-year OS of 27.9%–53.2%. Although the survival outcomes were similar, definitive CRT was more feasible when considering treatment-related mortality. Postoperative mortality was reported as 0%–19.1%, while definitive CRT rarely led to treatment-related death (Table 7).

LR (42%) was the major failure pattern in our current stage T4N0-1 patients. Isolated in-field recurrence was especially dominant (25%). This may due to the large volume of primary T4 tumors. There were few reports on failure patterns in T4N0-1 patients treated surgically. One previous study reported that the main cause of late death was LR in 58.62% of patients (17/29 patients). Recurrent tumors were located in the hilar and mediastinal lymph nodes in that study, and in two patients the recurrent tumor was in the left atrium. It seemed that surgery had a similar rate of LR rate [10]. Despite the omission of elective nodal irradiation, out-field nodal recurrence was rare. Distant recurrence occurred in 29% of patients, and over half of these cases also had LR.

Toxicity was acceptable with 13 events of grade 3 hematologic toxicities in 10 patients (11%) and three cases of grade 3 acute esophagitis (3%). The incidence of grade 3 esophagitis was lower than in a previous study (17%) [8]. This discrepancy can be attributed to our inclusion of patients receiving definitive RT alone. Also, T4N0-1 patients have a reduced possibility of having their esophagus in the radiation field compared with N2 patients unless the primary mass largely abuts the esophagus. Only one grade 3 chronic esophagitis was reported, and there was no grade 3 radiation pneumonitis. No grade 4 toxicities were reported.

From our present result, we concluded that improvement of in-field local tumor control is the main issue in stage T4N0-1 disease. Radiation dose escalation should be considered first. A Radiation Therapy Oncology Group (RTOG) 0617 study...
Definitive radiotherapy for T4N0-1 NSCLC compared standard-dose (60 Gy/30 fractions) and dose-escalated (74 Gy/37 fractions) radiation in stage III NSCLC. The outcome was poorer in the dose-escalated arm (median survival 28.7 months vs. 19.5 months, \( p = 0.0007 \)) without significant differences in toxicities over grade 3 [15]. This result was not clearly explained, and uncertainty remains regarding the intermediate dose escalation between 60 Gy and 74 Gy. Although many radiation oncologists prescribe intermediate dose levels, intermediate dose escalation was not effective in a previous multi-institutional propensity score matched analysis of stage III NSCLC patients [16]. The RTOG 0617 study enrolled all stage III NSCLC patients and not a specific subgroup. In that study, DM was slightly dominant in both treatment arms (standard dose arm 1.5-year DM, 35.3% vs. 44%; high dose arm, 42.4% vs. 47.8%). Mediastinal lymph nodal and distant recurrence might have increased and offset the effect of dose escalation due to the inclusion of patients who had high tumor burden in the nodal area. Our present study focused on T4N0-1 patients who were expected to have less DM. Our center used an intermediate dose escalation regimen of 66 Gy/30 fractions, and the 3-year cumulative LR was 48.4%. We suggest that further dose escalation would improve local tumor control for T4N0-1 patients but not for other stage III subgroups. The next dose escalation phase 3 trials should focus on the T4N0-1 subgroup.

Adoption of IMRT could have safety benefits [17]. In the current study, we did not use IMRT for dose escalation but to satisfy dose constraints. With IMRT, nine patients who could not meet the dose constraints were able to receive the same radiation dose as the other patients. These patients achieved a low rate of toxicities (Table 3). We suggest that further dose escalation would improve local tumor control for T4N0-1 patients but not for other stage III subgroups. The next dose escalation phase 3 trials should focus on the T4N0-1 subgroup.

Application of radio-sensitizing agents could also be beneficial. Traditional platinum-based chemotherapy agents are known to have radiation-sensitizing effect [21]. Gemcitabine and paclitaxel were introduced more recently and are thought to be superior radio-sensitizers [18]. Still, there are continuing efforts to find better radio-sensitizers. Choy et al. [20] reported the feasibility of pemetrexed. For T4N0-1 patients who were expected to have less DM, we suggest that further dose escalation would improve local control. The next dose escalation phase 3 trials should focus on the T4N0-1 subgroup.

Table 7. Results of T4N0-1M0 non-small cell lung cancer

| Author/characteristic            | Stage | Treatment                  | All patients with stage IIIA-N2 (n = 136, T4N0-1 [67], IIIA-N2 [69]) | Result                                                                 | Postoperative mortality (%) |
|----------------------------------|-------|----------------------------|------------------------------------------------------------------------|-----------------------------------------------------------------------|-----------------------------|
| Garrido et al. [11]/ prospective | IIIA  | Induction chemotherapy + surgery | Median survival 15.9 mo<sup>a</sup>                                    | 3-yr OS 36.8%, 5-yr OS 21.2%                                         | 7.8                         |
| Wu et al. [10]/ retrospective    | T4 (left atrium, pulmonary vein) | Surgery                  | Median survival 60.6 mo<sup>a</sup>                                    | 3-yr OS 71.4%, 5-yr OS 53.2%                                         | 0                           |
| Pitz et al. [14]/ retrospective  | T4    | Surgery                  | Only completely resected patients without postoperative mortality      | 5-yr OS 33.3%                                                        | 19.1                        |
| Reymen et al. [8]/ prospective   | T4N0-1 | Definitive CRT            | Median survival 34 mo (median follow-up 48 mo)                           | 3-yr OS 44%, 5-yr OS 25%                                             | 0                           |
| Current study/ retrospective     | T4N0-1 | Definitive (C)RT          | Median survival 23 mo (median follow-up 21 mo)                           | 3-yr OS 44%, 5-yr OS 36.3%                                           | 0                           |

OS, overall survival; CRT, chemo-radiotherapy; RT, radiotherapy.

<sup>a</sup>No significant difference in overall median survival time between stage T4N0-1 and IIIA-N2.
Another novel class of radio-sensitizers is nanoparticles (NP). NPs can encapsulate radiation sensitizing agents, protecting them from degradation, or sensitize cancer cells to radiation via physicochemical properties such as high Z number [21].

Combination thereby with surgery should be selectively considered for patients with minimal invasion to adjacent organs expected to obtain complete resection. A multidisciplinary team approach would be appropriate for the decision-making process.

Our present study had some limitations of note. First, staging was mainly based on imaging. Only 16% of our patients (n = 15) had invasive confirmations such as thoracotomy for T4 staging. PET-CT was routinely checked, so we expected >90% accuracy in the diagnosis of a T4 status; however, some patients might have had T3 tumors [22]. Discriminating N1 and N2 lymph nodes conglomerated with the primary mass was also difficult. The sensitivity and specificity of CT for detecting mediastinal nodal metastases have been reported as 52%–64% and 62%–69%, respectively [23]. Therefore, there is a significant risk of understaging nodal status.

Second, patients did not receive the same treatment regimen. Although the majority received definitive CRT, 25% were treated with RT alone, 15% received induction chemotherapy, and 2% received sequential treatment. However, all patients had an appropriate reason for choosing their treatment regimen. The RT-alone group contained patients with old age, poor performance, or poor pulmonary function. Induction chemotherapy was recommended for patients with very large tumors to avoid radiation toxicity. We feel that the present study reflects the real situation in the clinic.

Lastly, this was a retrospective study with inherent biases that could have affected our results.

In conclusion, definitive radiotherapy for T4N0-1 NSCLC results in favorable survival with acceptable toxicity rates. LR, especially in-field recurrence, is a major recurrence pattern. For improving local tumor control, the application of IMRT and additional radio-sensitizing agents is needed.

Conflict of Interest

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

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